European Medicines Agency pre-authorisation procedural advice for users of the centralised procedure

This integrated version has been created for printing purposes only. Please refer to the individual question & answers as published in the pre-submission guidance for access to the hyperlinked information.

Questions and answers are being updated continuously, and will be marked by "NEW" or "Rev." with the relevant date upon publication.

This guidance document addresses a number of questions which users of the centralised procedure may have. It provides an overview of the European Medicines Agency’s position on issues, which are typically addressed during the course of pre-submission meetings.

It will be updated regularly to reflect new developments, to include guidance on further pre-authorisation procedures and to reflect the implementation of the new European legislation. Revised topics will be marked by "New" or "Rev" upon publication.

The EMA emphasises the importance of pre-submission meetings between applicants and the EMA/(Co-) Rapporteur. Pre-submission meetings (which should take place approximately 7 months prior to the anticipated date of submission of the application) are a vital opportunity for applicants to obtain procedural, regulatory and legal advice from the EMA. The product team is available to address any questions MAHs may have regarding their pre-authorisation application.

This guidance information and fruitful pre-submission meetings should enable applicants to submit applications, which are in conformity with the legal and regulatory requirements and which can be validated speedily. Pre-submission meetings will also enable applicants to establish contact with the EMA staff closely involved with the application as it proceeds.

**Note:**

It should be highlighted that this document has been produced for guidance only and should be read in conjunction with "The rules governing medicinal products in the European Union", Volume 2A, Notice to Applicants.
Applicants must in all cases comply with all requirements of Community Legislation. Provisions, which extend to EEA countries (i.e. the EU member states, plus Norway, Iceland and Liechtenstein) by virtue of the EEA agreement, are outlined in the relevant sections of the text.
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1. Is my medicinal product eligible for evaluation under the Centralised Procedure? Rev. Feb 10

Regulation (EC) No 726/2004 of the European Parliament and of the Council lays down a centralised Community procedure for the authorisation of medicinal products, for which there is a single application, a single evaluation and a single authorisation allowing direct access to the single market of the Community.

A marketing authorisation granted under the centralised procedure is valid for the entire Community market, which means the medicinal product, may be put on the market in all member states.

1.1. Article 3 of Regulation (EC) No 726/2004 defines the scope and eligibility of applications for evaluation under the centralised procedure through which medicinal products must (“mandatory scope”) or may (“optional scope” or “Generic/Hybrid”) be authorised by the Community.

1.1.1. Mandatory scope (Article 3(1)):

For medicinal products falling within the mandatory scope of the Annex of Regulation (EC) No 726/2004, applicants are obliged to use the centralised procedure by submitting their marketing authorisation application to the EMA. Medicinal products under the mandatory scope belong to one of the following categories:

1. Medicinal products developed by means of one of the following biotechnological processes:
   - recombinant DNA technology;
   - controlled expression of genes coding for biologically active proteins in prokaryotes and eukaryotes including transformed mammalian cells;
   - hybridoma and monoclonal antibody methods;

Similar biological (“biosimilar”) medicinal products which are developed by one of the above biotechnological processes also fall under the mandatory scope of the centralised procedure.

1.1. Advanced therapy medicinal product as defined in Article 2 of Regulation (EC) No 1394/2007
   - Gene therapy medicinal products
   - Somatic cell therapy medicinal products
   - Tissue engineered products

   “Transitional period” applies (Article 29):

   Advanced therapy medicinal products, other than tissue engineered products, which were legally on the Community market in accordance with national or Community legislation on 30 December 2008, shall comply with this Regulation no later than 30 December 2011.

   Tissue engineered products which were legally on the Community market in accordance with national or Community legislation on 30 December 2008 shall comply with this Regulation no later than 30 December 2012.
2. Medicinal products for human use containing a new active substance which, on the date of entry into force of the Regulation (20 November 2005), was not authorised in the Community and for which the therapeutic indication is the treatment of any of the following diseases:

- Acquired immune deficiency syndrome;
- Cancer;
- Neurodegenerative disorder;
- Diabetes;

And with effect from 20 May 2008

- Auto-immune diseases and other auto-immune dysfunctions;
- Viral diseases;

Clarifications on the working definitions of the diseases listed above are available in the "Scientific Aspects and Working definitions for the mandatory scope of the centralised procedure (EMEA/CHMP/121944/2007)".

3. Medicinal products that are designated as orphan medicinal products pursuant to Regulation (EC) No 141/2000.

1.1.2. Optional Scope (Article 3(2)):

For medicinal products falling under the optional scope, applications for the following categories may, at the request of the applicant, be accepted for assessment under the centralised procedure:

1. A medicinal product containing a new active substance which, on the day of entry into force of the Regulation (20 November 2005) was not authorised in the Community (Article 3(2)a).

A new chemical, biological or radiopharmaceutical active substance, as defined in Annex III to Chapter 1 of the Notice to Applicants, includes:

- a chemical, biological or radiopharmaceutical substance not previously authorised as a medicinal product in the European Union;
- an isomer, mixture of isomers, a complex or derivative or salt of a chemical substance previously authorised as a medicinal product in the European Union but differing in properties with regard to safety and efficacy from that chemical substance previously authorised;
- a biological substance previously authorised as a medicinal product in the European Union, but differing in molecular structure, nature of source material or manufacturing process;
- a radiopharmaceutical substance which is a radionuclide, or a ligand not previously authorised as a medicinal product in the European Union, or the coupling mechanism to link the molecule and the radionuclide has not been authorised previously the European Union;

2. A medicinal product, which constitutes a significant therapeutic, scientific or technical innovation or that the granting of authorisation is in the interest of patients at Community level (Article 3(2)b).

For the purpose of determining whether “a medicinal product constitutes a significant therapeutic, scientific or technical innovation”, the Agency will consider if:

- the medicinal product provides a new alternative to patients in treating, preventing or diagnosing a disease, or,
• the medicinal product development is based on significant new scientific knowledge or on the application of a new scientific knowledge, or,
• a new technology or a new application of technology is used for the development or the manufacture of the medicinal product.

Regarding the criteria of ‘interest of patients’, a medicinal product which does not constitute a significant therapeutic, scientific or technical innovation, can be of patient interest at Community level when it addresses a specific health issue, allows access to medicines, or provides another type of contribution to patient care in the Community.

1.1.3. Generic/Hybrid of centralised medicinal product applications (Article 3(3)):

A generic or hybrid medicinal product of a reference medicinal product authorised via the centralised procedure has ‘automatic’ access to the centralised procedure under Article 3(3).

1.1.4. Duplicate/multiple marketing authorisations

Multiple/duplicate or informed consent applications from the same or different marketing authorisation holder for a specific medicinal product with an active substance(s) already authorised via the centralised procedure, have automatic access to the centralised procedure.

1.2. Applications for certain medicinal products for paediatric use may also be eligible for evaluation through the centralised procedure in accordance with the Paediatric Regulation (Regulation (EC) No 1901/2006)

1.2.1. Marketing Authorisation application including paediatric indication(s) for a medicinal product which is not authorised in the Community (Article 28):

A marketing authorisation application for a medicinal product not authorised in the Community on the date of entry into force of the Paediatric Regulation (26 July 2008) and which includes one or more paediatric indication(s) on the basis of studies conducted in compliance with an agreed paediatric investigation plan (PIP).

1.2.2. Applications for a new paediatric indication, a pharmaceutical form and/or a route of administration for nationally authorised medicinal products (Article 29):

Applications for a new paediatric indication, a pharmaceutical form and/or a route of administration for a nationally authorised medicinal product falling under Article 8 of Regulation (EC) No 1901/2006 and which include results of studies conducted in compliance with an agreed PIP. Article 8 of Regulation (EC) No 1901/2006 applies to authorised medicinal products which are protected either by a supplementary protection certificate under Regulation (EEC) No 1768/92, or by a patent which qualifies for the granting of the supplementary protection certificate.

1.2.3. Paediatric Use Marketing Authorisation (PUMA) application (Article 31):

Applications for a PUMA concerns only a medicinal product for human use which is not protected by a supplementary protection certificate under Regulation (EEC) No 1768/92 or by a patent which qualifies
for the granting of the supplementary protection certificate, and which covers exclusively paediatric therapeutic indications, including the appropriate strength, pharmaceutical form or route of administration for that product.

In all cases listed above, the eligibility of a medicinal product for evaluation via the centralised procedure must be requested by the applicant by submitting a Pre-submission request form (Eligibility) to CPeligibility@ema.europa.eu.

References

- Regulation (EC) No 726/2004
- Regulation (EC) No 1394/2007
- "Scientific Aspects and Working definitions for the mandatory scope of the centralised procedure (EMEA/CHMP/121944/2007)".
- The Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2A, Chapter 1 on “Marketing authorisation”
- The Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2A, Chapter 4 on ”Centralised Procedure”
2. How and when should the eligibility request be sent to the EMA? *Rev. Feb 10*

Regardless of whether the product falls into the mandatory or optional scope, or would have "automatic access" or access in accordance with the Paediatric or Advanced Therapy Regulation, an 'eligibility request' should always be submitted using the specific form and accompanied by a justification of eligibility for evaluation under the centralised procedure. The applicant should clearly address the specific criterion fulfilled by the product to be eligible for the centralised procedure (for eligibility criteria see Q1).

Please note that:

1. In cases where products fall under the "mandatory scope" criterion ([Art. 3(1)](EC) No. 726/2004), the relevant justification should be provided.

For Advanced Therapy Medicinal Products (ATMPs), the relevant justification and documentation (including EMA scientific recommendation on classification of ATMPs by the Committee for Advanced Therapies (CAT) if available) should be provided.

NB: Only one criterion can be chosen

2. In cases where products fall under one of the "optional scope" criteria ([Art. 3(2)](EC) No. 726/2004), the justification should consist of a concise summary document of preferably two pages stating why the product should qualify for evaluation through the centralised procedure. The applicant should clearly state in the request which criterion the appended justification concerns:

- Art. 3(2) a: New active substance; or
- Art. 3(2) b Significant therapeutic innovation, or
- Art. 3(2) b Significant scientific innovation or
- Art. 3(2) b Significant technical innovation; or
- Art. 3(2) b Interest of patient at the community level.

NB: Only one criterion can be chosen and must be adequately justified; e.g. eligibility in accordance with Art 3(2)b of Regulation (EC) No. 726/2004 – Significant therapeutic innovation

3. In the following cases where the medicinal product applied for may have "automatic access" to the centralised procedure, this should be the basis for the justification to be submitted. This is the case when the medicinal product applied for, is either:

- a "generic/hybrid" ([Art. 3(3)](EC) No. 726/2004); or
- a duplicate/multiple; or
- an informed consent

to a centrally authorised medicinal product, adequate and relevant information on the already centrally authorised medicinal product should be provided as background information (such as invented name/INN/ Commission Decision date/ type of application submitted and criteria/ indent under which the medicinal product was eligible to access the centralised procedure at the time (EMA letter to be annexed)).
4. When the medicinal product applied for, is either:

- an application including paediatric indication(s) in compliance with an agreed PIP (Art. 28 of Regulation (EC) No 1901/2006); or

- an application consisting of a new paediatric indication, a new pharmaceutical form and/or a new route of administration in compliance with an agreed PIP for a nationally authorised medicinal product (Art. 29 of Regulation (EC) No 1901/2006); or

- an application for a Paediatric Use Marketing Authorisation (PUMA) (Art. 31 of Regulation (EC) No 1901/2006),

adequate and relevant information should be provided (such as copy of the EMA PIP decision to be provided in annex), details of the paediatric indication/form/route applied for and a listing of the study data collected in accordance with the PIP which will be submitted in the planned application).

When submitting a request, the applicant should use the Pre-submission request form (Eligibility) and send it electronically, to: CPeligibility@ema.europa.eu, together with a separate Annex 1 (draft Summary of Product Characteristics) and Annex 2 (Justification for Eligibility) especially required for medicinal products falling under the optional scope of Article 3(2)b.

EMA recommends providing the eligibility request preferably, at the earliest, 18 months before submission of the marketing authorisation application (MAA) and, at the latest, 7 months before the MAA is filed with the EMA, at which point it could be submitted as part of the “letter of intent to submit”. For Eligibility requests submitted as part of the “letter of intent to submit”, Rapporteurs will be automatically appointed following the confirmation of the eligibility to the centralised procedure provided that the planned submission date is within 6-7 months.

The eligibility request and supporting documentation should be submitted to the EMA 10 calendar days before the CHMP meeting (see enclosed table for submission deadlines), so as to ensure its inclusion in the next CHMP agenda.

Any request received after the deadline will be considered the following month.

The eligibility will be evaluated on a case-by-case basis by the EMA/CHMP. The applicant will, in all cases, be informed of the CHMP opinion, the week following the CHMP meeting where the discussion took place.

NB: Review of eligibility applications made under Article 3(2)b will take place over 2 consequent CHMP meetings because of the need to appoint a sponsor(s) to assess the request.

References

- Regulation (EC) No 726/2004
- Regulation (EC) No 1394/2007
- The Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2A, Chapter 4 on “Centralised Procedure”
- “Scientific Aspects and Working definitions for the mandatory scope of the centralised procedure” (EMEA/CHMP/121944/2007)
3. What will be the legal basis for my application? Rev. Feb 08

The applicant should clearly indicate the legal basis for the submission of their application in the EU Application Form, i.e. select one of the following articles of Directive 2001/83/EC:

- Article 8(3) - Full application
- Article 10 - Generic, hybrid or similar biological application
- Article 10a - Well-established use application
- Article 10b - Fixed combination application
- Article 10c - Informed consent application

At pre-submission meetings, it is strongly recommended to discuss the proposed legal basis in view of the available data, with the EMA in order to prevent difficulties at validation.

3.1. Article 8(3) - Full application:

For full applications according to Article 8(3) of Directive 2001/83/EC, the results of pharmaceutical tests (physico-chemical, biological or microbiological), pre-clinical tests (pharmacological and toxicological), and clinical trials need to be submitted. Detailed data requirements are set-out in Annex I to Directive 2001/83/EC, as amended by Directive 2003/63/EC.

Any deviations from these requirements, in particular, absence of a study/test report, requires a justification as to why the results are not provided and whether the requirements as set out in the Annex I to Commission Directive 2001/83/EC, are considered fulfilled.

Justifications are to be provided in the respective non-clinical and clinical overviews in Module 2. Further guidance on the drafting of such justifications is provided below. There is a possibility to use "umbrella" justifications to cover absence of more than one study report or more than one indent provided that is clear that the justification applies to several study reports. There is no need, however, to create and include a document in Module 4 and 5 which (only) refers to the presence of a justification in Module 2.

3.1.1. ‘Full-mixed’ application:

Where Module 4 and/or 5 consists of a combination of reports of limited non-clinical and/or clinical studies carried out by the applicant and of bibliographical references this kind of application has also to be submitted according to Article 8(3) of Directive 2001/83/EC (So-called ‘full-mixed’ application - see also section on ‘mixed’ marketing authorisation application in Part II of Annex I to the Directive).

A justification for not having performed certain tests/trials and for providing literature references instead, should be provided as to why the references provided by the applicant can replace the study reports, and how the results presented fulfill the requirements as set out in the Annex I to Commission Directive 2001/83/EC. The general principles for ‘justifications’ as outlined above also apply to full-mixed applications.

Such literature references, when replacing required study reports, should be included in the relevant Module 4/5 indents and should be summarised in Module 2 as required for any other study report.
“Supportive-only” literature references (i.e. provided in addition to study reports), should be provided in the CTD sections for “references” and do not need to be summarized in Module 2.

3.1.2. Guidance for the preparation of the Non-clinical and/or clinical Overviews in case of Art 8.3 (Full or “Full-mixed”) marketing authorisation applications:

- For each item of section 4.1 and 5.1 of Part I of the Annex I to Dir 2001/83/EC, the Applicant should indicate whether the Application contains the results of pre-clinical tests or clinical trials in the format of detailed study reports (hereafter referred to as “study reports”), and/or in the format of bibliographical references, or no information at all.

- If study reports are provided and cover all the requirements for a specific section, no further justifications are required.

- If results are submitted in the form of bibliographical references for a specific item, a justification is required as to why the references provided by the applicant can replace the study reports, and how the results presented fulfil the requirements as set out in the Annex I to Commission Directive 2001/83/EC.

- If no results are provided for a certain test or trial, a justification is required as to why the results are not provided and whether the requirements as set out in the Annex I to Commission Directive 2001/83/EC, are considered fulfilled. A simple statement such as “Not Applicable” is not an acceptable justification.

Justifications for absence of study reports in each of the sections can be based, for example, on the following principles:

- Specific derogations foreseen in Directive 2001/83/EC;
- Specific derogations foreseen in CHMP Guidelines;
- Animal welfare¹ and ethical considerations² coupled with expert assessment that further tests or trials are unlikely to extend scientific knowledge of subject area;
- Expert assessment that repetition of certain tests or trials is unlikely to extend scientific knowledge of subject area (e.g., extent of clinical experience with active substance at the time of development to replace certain non-clinical tests);
- Scientific argumentation regarding inapplicability of such tests and trials;
- Inability to provide comprehensive data in accordance with Article 14(8) of Regulation (EC) No 276/2004 and as outlined in general provisions of Section 6 of Part II of the Annex to Commission Directive 2001/83/EC (applications in exceptional circumstances);

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² Declaration of Helsinki
3.2. Article 10 - Generic, hybrid or similar biological applications

3.2.1. Generic applications:

According to Article 10(1) of Directive 2001/83/EC, the applicant is not required to provide the results of pre-clinical tests and clinical trials if he can demonstrate that the medicinal product is a generic medicinal product of a reference medicinal product which is or has been authorised under Article 6 of Directive 2001/83/EC for not less than 8 years in a Member State or in the Community.

A generic medicinal product is defined as a medicinal product that has:

- the same qualitative and quantitative composition in active substances as the reference product,
- the same pharmaceutical form as the reference medicinal product,
- and whose bioequivalence with the reference medicinal product has been demonstrated by appropriate bioavailability studies.

This type of application refers to information that is contained in the dossier of the authorisation of the reference medicinal product, for which a marketing authorisation has been granted in the Community on the basis of a complete dossier in accordance with article 8(3), 10a, 10b or 10c of Directive 2001/83/EC.

It should be noted that the period of 8 years from initial authorisation of the reference medicinal product, providing a period of so-called “data exclusivity”, only applies to those reference medicinal products for which the initial application for authorisation was submitted through the centralised procedure after 20 November 2005.

3.2.2. Hybrid applications:

Hybrid applications under Article 10(3) of Directive 2001/83/EC differ from generic applications in that the results of appropriate pre-clinical tests and clinical trials will be necessary in the following three circumstances:

- where the strict definition of a ‘generic medicinal product’ is not met;
- where the bioavailability studies cannot be used to demonstrate bioequivalence;
- where there are changes in the active substance(s), therapeutic indications, strength, pharmaceutical form or route of administration of the generic product compared to the reference medicinal product

In such cases the results of tests and trials must be consistent with the data content standards required in the Annex to the Directive 2001/83/EC as amended by Directive 2003/63/EC.

These applications will thus rely in part on the results of pre-clinical tests and clinical trials for a reference product and in part on new data. Some guidance on the appropriate additional studies required is indicated in Annex IV of the Chapter 1 of the Notice to Applicants.

3.2.3. Similar biological application:

In Article 10(4) of Directive 2001/83/EC it is stated that where a biological medicinal product which is similar to a reference biological product, does not meet the conditions in the definition of generic medicinal products, owing to, in particular, differences relating to raw materials or differences in manufacturing processes of the similar biological medicinal product and the reference biological
medicinal product, the results of appropriate pre-clinical tests or clinical trials relating to these conditions must be provided. The type and quantity of supplementary data to be provided must comply with the relevant criteria stated in Annex I to Directive 2001/83/EC and the related detailed guidelines. The results of other tests and trials from the reference medicinal product’s dossier shall not be provided.

The chosen reference medicinal product must be a medicinal product authorised in the Community, on the basis of a complete dossier in accordance with the provisions of Article 8 of Directive 2001/83/EC.

3.3. Article 10a - Well-established use application:

According to Article 10a of Directive 2001/83/EC, as amended it is possible to replace results of pre-clinical and clinical trials by detailed references to published scientific literature (information available in the public domain) if it can be demonstrated that the active substances of a medicinal product have been in well-established medicinal use within the Community for at least 10 years, with recognised efficacy and an acceptable level of safety. In this regard, the provisions of Annex I (Part II.1) to Directive 2001/83/EC shall apply.

The following criteria for the demonstration of such well-established use should be taken into account:

- the time over which a substance has been used with regular application in patients; quantitative aspects of the use of the substance, taking into account the extent to which the substance has been used in practice, the extent of use on a geographical basis and the extent to which the use of the substance has been monitored by pharmacovigilance or other methods;
- the degree of scientific interest in the use of the substance (reflected in the published scientific literature) and the coherence of scientific assessments;

For such applications, the provisions of the Annex I to Directive 2001/83/EC apply in like manner. They are considered as full and independent applications. Applicants should submit Modules 1, 2 and 3 as described in Part I of Annex I to Directive 2001/83/EC. For Modules 4 and 5, a detailed scientific bibliography shall address all required pre-clinical and clinical characteristics, and should be summarised in Module 2. As with any other full application, if parts of the dossier are incomplete, particular attention must be paid to justify such absences in the non-clinical/clinical overviews.

It should be noted that, if well-known substances are used for entirely new therapeutic indications, it is not possible to solely refer to a well-established use and additional data on the new therapeutic indication together with appropriate pre-clinical and human safety data should be provided. In such case, Article 8(3) of Directive 2001/83/EC should be used as legal basis.

3.4. Article 10b - Fixed combination application:

According to Article 10b of Directive 2001/83/EC, in the case of medicinal products containing active substances used in the composition of authorised medicinal products but not hitherto used in combination for therapeutic purposes, the results of new pre-clinical tests or new clinical trials relating to that combination shall be provided in accordance with Article 8(3)(i) of the same Directive, but it shall not be necessary to provide scientific references relating to each individual active substance.

The combination of active substances within a single pharmaceutical form of administration according to this provision is a so-called ‘fixed combination’.

Applications for fixed combination medicinal products can be accepted and validated under Article 10b on condition that the individual substances have been authorised as a medicinal product in the EEA via a Community or national procedure.
It follows from the wording of Article 10b as well as from Part II.5 of Annex I to the Directive 2001/83/EC as amended, that a full dossier, comprising all the information of modules 1 to 5, has to be provided in relation to the fixed combination. Any absence of specific fixed combination data should be duly justified in the Non-clinical and/or clinical Overviews (see general guidance above).

Although there is no requirement for the inclusion of data on the individual active substances, it is possible to include information on the individual substances (literature or actual data), especially in order to justify the absence of certain specific data on the combination.

3.5. Article 10c - Informed consent application:

According to Article 10c of Directive 2001/83/EC as amended, following the granting of a marketing authorisation, the authorisation holder may allow use to be made of the pharmaceutical, non clinical and clinical documentation contained in the dossier of the medicinal product for the purpose of examining subsequent applications relating to other medicinal products possessing the same qualitative and quantitative composition in terms of active substances and the same pharmaceutical form.

It is a prerequisite for the use of Article 10c as legal basis that consent has been obtained from the marketing authorisation holder of the reference product for all three modules containing the pharmaceutical, pre-clinical and clinical data (modules 3, 4 and 5), and the applicant of the informed consent application should have permanently access to this documentation or should be in possession of the information.

For such informed consent applications, only a complete module 1 should be submitted, including the Application Form with relevant Annexes (e.g. copy of correspondence with the European Commission for multiple applications, if applicable, see also Q9 – multiple applications, and the letter of consent from the MAH of the authorised medicinal product allowing access to modules 2, 3, 4, 5 of the initial dossier and any subsequent documentation submitted)

If the dossier of the authorised medicinal product includes an ASMF, a new letter of access should be included in module 1 of the informed consent application.

References

- Regulation (EC) No 726/2004
- Directive 2001/83/EC, as amended
- The Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2A, Chapter 1
- EMEA guidance for users of the Centralised Procedure for generic/hybrid applications (CHMP/225411/2006)
- CHMP Guideline on similar biological medicinal products (CHMP/437/04)
4. How will I know if the proposed (invented) name of my medicinal product is acceptable from a public health point of view? Rev. Feb 15

In accordance with Article 6 of Regulation (EC) No 726/2004, “each application for the authorisation of a medicinal product for human use (…) otherwise than in exceptional cases relating to the application of the law on trademarks, shall include the use of a single name for the medicinal product.” The Centralised Procedure therefore requires one single name for the medicinal product to be authorised.

According to Article 1(20) of Directive 2001/83/EC, as amended, the name of the medicinal product “may be either an invented name not liable to confusion with the common name, or a common name or scientific name accompanied by a trademark or the name of the Marketing Authorisation Holder”. It is also understood by legislation that a common name is according to Article 1(21) of Directive 2001/83/EC, as amended, “The international non-proprietary name (INN) recommended by the World Health Organisation, or, if one does not exist, the usual common name”.

Although it is not mandatory under Community legislation, in practice, many companies submitting marketing authorisation applications under the Centralised Procedure wish to use invented names for their medicinal products.

As part of the EMA’s role in evaluating the safety of medicinal products in the centralised procedure, it is obliged to consider whether the (invented) name proposed for a medicinal product could create a public-health concern or potential safety risks.

In particular, the (invented) name of a medicinal product:

- should not convey misleading therapeutic or pharmaceutical connotations;
- should not be misleading with respect to the composition of the product;

In order to identify, at an early stage, potential difficulties presented by the (invented) name(s) proposed by an applicant, the EMA/CHMP set up the Name Review Group (NRG), to perform the review of names. The NRG is also responsible for updating the “Guideline on the acceptability of names for human medicinal products processed through the centralised procedure” (EMA/CHMP/287710/2014).

It should be highlighted that when an applicant/MAH wishes to use instead of an invented name the common name or scientific name, together with a trademark or the name of the Marketing Authorisation Holder, this is also subject to NRG review.

4.1. The Name Review Group (NRG)

The NRG is composed of representatives of EU Member States and is chaired by an EMA representative. Representatives of the European Commission and the EMA Secretariat also participate in the work of the group. Other relevant experts (e.g. WHO experts) are consulted on a case-by-case basis.

The NRG meets 6 times a year (approximately every 2 months). Its conclusions are presented for adoption at the subsequent CHMP plenary meeting.

The criteria applied by the NRG when reviewing the acceptability of proposed invented names are detailed in the “Guideline on the acceptability of names for human medicinal products processed through the centralised procedure” (EMA/CHMP/287710/2014), hereafter referred to as the ‘Guideline’.
4.2. The EMA procedure for checking proposed (invented) names

4.2.1. Submission of the (invented) name request by the Applicant/MAH

Provided that the medicinal product is eligible for evaluation under the Centralised Procedure, the applicant should inform the EMA of the proposed (invented) name(s) for their medicinal product at the earliest 18 months and preferably 4-6 months prior to the planned submission date of the marketing authorisation application. See also Question 4a. What are the dates for submission of invented name requests for the deadlines for submission of Proposed (Invented) Names.

Applicants may submit a name review request after eligibility has been confirmed by the CHMP or in parallel to the eligibility request. Applicants are advised to contact the NRG secretariat prior to submission of the name review request form for advice if eligibility is not yet confirmed at that time.

The ‘Proposed (Invented) Name request form’, along with either a draft Summary of Product Characteristics (SmPC) or a product profile and any other relevant information, should be sent to the EMA at the following e-mail address: NRG@ema.europa.eu. An electronic request form (in pdf format) has been developed and replaces the current form in Word format.

Up to two (invented) names can be accepted per Marketing Authorisation Application from which the applicant should select the final name to be used. Up to two newly proposed (invented) names can be considered at each NRG meeting per Marketing Authorisation Application.

It should be noted that once two (invented) names have been deemed acceptable by the NRG for a Marketing Authorization Application, no further review of newly proposed names is allowed unless agreed with EMA on duly justified grounds (i.e. identification of safety issue/health concern after acceptance of (invented) names, conditional acceptability of previously reviewed (invented) names, constraints achieving a global (invented) name, issues relating to the application of the law on trademarks, etc.).

Applicants should follow the criteria described in the ‘Guideline’ when proposing (invented) names and would be expected to review the proposed (invented) name, applying the criteria before requesting that an invented name be considered. Where the applicant deviates from these criteria, justification should be provided.

Where the applicant submits proposed (invented) names intended to be used in the context of multiple marketing authorisations/applications, it shall specifically request the NRG to consider whether the proposed (invented) names cannot be considered potentially confusing with each other (see also question on Multiple Applications).

4.2.2. Consultation with the Member States and NRG discussion/CHMP adoption

The proposed (invented) name(s) and all the background information provided by the applicant(s)/MAH(s) are sent to every NRG contact point nominated by National Competent Authorities (NCAs) of EU Member States for their review and will subsequently be discussed at the NRG meeting. The detailed procedure is described in the ‘Guideline’.

The NRG conclusions/recommendations are presented for adoption to the subsequent CHMP plenary meeting, after which the applicant will be informed of the outcome of the discussion on the acceptability of the proposed (invented) name(s) for their medicinal product together with the reasons and source for the objections(s) raised, where applicable. See also Question 4a. What are the dates for submission of invented name requests? for the dates of NRG discussion/CHMP adoption.
4.2.3. Rejection by NRG/CHMP of a proposed (invented) name

In case of rejection of a proposed (invented) name by NRG/CHMP, the applicant/MAH has the following possibilities:

- To submit up to two new (invented) names proposals, which are checked through the same procedure as described above. In the case that a name has already been accepted in a previous NRG meeting and two new names are submitted, the applicant is required to indicate in the 'Proposed (Invented) Name Request form' which two names should be finally retained in the NRG database.

- To provide a justification to retain the (invented) name (addressing specifically all the objections raised) using the 'Proposed (Invented) Name Request form' and selecting 'Justification Form' in the 'Form Type' area. Such justification will be reviewed as described in the 'Guideline'. If the proposed (invented) name cannot be accepted prior to submission, the Marketing authorisation application can be submitted under either any of the proposed (invented) names or the common name or scientific name accompanied by a trademark or the name of the MAH.

Applicants may submit justifications for rejected names in addition to the entitlement of 2 (invented) names reviewed per meeting. However, only two accepted names can be retained in the NRG database and therefore the applicant should indicate in advance which two names should be retained in case that they are accepted.

At the latest one month prior to the adoption of the CHMP opinion on the concerned MAA, the applicant will in such case have to inform the EMA and the NRG Secretariat on the acceptable invented name of their choice.

- If no suitable invented name has been identified at that stage, the opinion will be adopted using the common name or scientific name accompanied by the name of the MAH. Applicants are hereby reminded that such name also needs NRG review and acceptance by the CHMP prior to the adoption of the opinion. In this case, as soon as the Commission Decision is granted, the MAH may submit a variation to introduce an invented name, on the condition that such name has been considered acceptable by the NRG.

- Exceptionally, provided all means have been exhausted, the applicant/MAH may request the matter to be presented to the CHMP within the context of the evaluation of the medicinal product (e.g. oral explanation).

4.2.4. Change of the (invented) name after the marketing authorisation is granted

In accordance with Commission Regulation (EC) No 1234/2008, the (invented) name of a medicinal product may be changed after a marketing authorisation is granted through a Type IA\textsubscript{INN} (No A.2) variation procedure.

This can be done either in case of a marketing authorisation being granted under INN or common name together with a trademark or the name of the MAH or in case the MAH wants to change the initial invented name.

Such Type IA\textsubscript{INN} variation is possible provided that the check by the Agency on the acceptability of the new name had been finalised and was positive before implementation of the new name. Immediately upon implementation of the change, the MAH must submit a Type IA\textsubscript{INN} variation notification to the Agency for review (see the EMA Post-Authorisation Procedural Advice on Type IA variations).
Taking into account that the MAH will be required to submit the EMA letter of acceptance of the concerned (invented) name as part of the variation application, it is recommended that the proposed invented name be submitted at least 4-6 months in advance of the foreseen implementation date and submission of the Type IAIN variation notification.

References

- Regulation (EC) No 726/2004
- Directive 2001/83 EC as amended
- “Centralised Procedure”, The Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2A, Chapter 4
- “Guideline on the acceptability of invented names for human medicinal products processed through the centralised procedure” (EMA/CHMP/287710/2014)
- Regulation (EC) No 1234/2008
- “Post-Authorisation Procedural Advice Human Medicinal Products” (EMEA-H-19984/03)
5. How shall I compose the complete name of my medicinal product? *Rev. May 11*

Each medicinal product should be placed on the market under a name and in a package suitable to ensure identification and differentiation. A medicinal product authorised under the Centralised Procedure must have the same name in all EU Member States.

The medicinal product should be identified in the product information according to the following rule: the name of the medicinal product should be followed by the strength and the pharmaceutical form. However, when otherwise referring to the medicinal product throughout the product information text, the strength and the pharmaceutical form do not have to be mentioned in the name.

In the SPC, the INN or the common name of the active substance should be used when referring to properties of the active substance(s) rather than the invented name. The use of pronouns (e.g. "it") is encouraged whenever possible.

Thus, whenever the "name of the medicinal product" is specifically required to be provided in the SPC, labelling (on the outer or immediate packaging or on blisters) or the Package Leaflet, it should be written in the following order as:

{(invented) name strength pharmaceutical form}, whereby

- **invented name**: no ® ™ symbols attached
- **Pharmaceutical form**:

  The pharmaceutical form should be stated according to the full "Standard Terms" published by the Council of Europe, in the singular (except for tablets and capsules). Where the Council of Europe short standard term is used on small immediate packaging materials (blisters, strips, small immediate packaging units) in case of space limitation, the short term should be added in brackets in section 3 of the SPC.

  E.g.  (invented name) X mg hard capsules

  (invented name) Y mg/g cream

- **The different strengths of fixed-combination products should be presented separated by a "/"**.

  E.g.  (invented) name 150 mg/12.5 mg tablets

For mock-ups and specimens, this information may be presented on different lines of text or in different font sizes if necessary, provided that the appearance of the name is as an integrated item.

E.g.  (invented) name Z mg/ml

  Solution for injection

Where the INN or the common name is to be provided in addition to an invented name, this should preferably be given on the line of text directly below the complete name.

References

- Directive 2001/83/EC, title I, II and V, as amended
• “Guideline on the readability of the label and package leaflet of medicinal products for human use”, the Rules governing Medicinal products in the European Community, Volume 2C, Notice to applicants

• “Guideline on Summary of Product Characteristics”, the Rules governing Medicinal products in the European Community, Volume 2C, Notice to applicants

• QRD Product Information Template with explanatory notes
6. What legal status can I obtain for my medicinal product?

Rev. July 06

In accordance with Article 9(4)(b) of Regulation (EC) No 726/2004, the documents annexed to the CHMP favourable opinion to the granting of a Marketing Authorisation for a medicinal product shall include "details of any conditions or restrictions which should be imposed on the supply or use of the medicinal product concerned, including the conditions under which the medicinal product may be made available to patients, in accordance with the criteria laid down in Title VI of Directive 2001/83/EC, as amended".

The classification for the supply of the medicinal product to the patient is also referred to as 'Legal Status'.

6.1. Categories for the Legal Status of a medicinal product

At the first level, 'main categories', the medicinal product is classified either as:

- subject to medical prescription or
- not subject to medical prescription

To this end, the criteria laid down in Article 71(1) of Directive 2001/83/EC, as amended, should be taken into account.

For products subject to medical prescription, where applicable, there is a second level and the EMA may have to apply one of the following additional 'sub-categories', in accordance with Article 70(2) of Directive 2001/83/EC as amended:

- Medicinal product subject to special medical prescription
- Medicinal product on restricted medical prescription, reserved for use in certain specialised areas

To this end, the factors laid down in Article 71 paragraphs 2 and 3 should be taken into account.

Medicinal products, which meet the criteria for both above-mentioned 'sub-categories', will be subject to special and restricted medical prescription.

There is another 'sub-category' foreseen in Article 70(2) of Directive 2001/83/EC, as amended, i.e.: 'medicinal products on medical prescription for renewable or non-renewable delivery'. The definition and therefore also the implementation may vary in those Member States where the 'sub-category' exists. Therefore it has been decided that for centrally authorised products such 'sub-category' will not be explicitly mentioned in the Opinion/Decision, leaving for Member States the possibility of the implementation of the 'sub-category' in accordance with national measures and in compliance with the content of the SPC.

6.2. Implementation of the Legal Status in the CHMP Opinion

At the pre-submission stage applicants should include a proposed classification for the supply of the medicinal product in their "notification of intention to submit an application” to be sent to the EMA at least 7 months before submission. At the time of the submission of the application applicants should indicate their proposal for Legal Status in the section 2.3 of the Module 1 application form (available in the Notice to Applicants (NTA) Volume 2B - Application Form: Module 1.2 Application form).
The CHMP refers to the above-mentioned criteria and factors where it comes to take a decision on the Legal Status.

The Legal Status will be mentioned in the CHMP opinion and in the Commission decision.

In the CHMP opinion, the Legal Status will be reflected in the following annexes:

- Annex I of the CHMP opinion (Summary of Product Characteristics)

Wherever appropriate, the SPC will include in section 4.2 an explanation on how the medicinal product should be supplied to patients (e.g. to be administered in a hospital setting or prescribed by specialists only, or specific type of care during the treatment of a chronic disease).

- Annex II.B of the CHMP opinion (Conditions or restrictions regarding supply and use) should mention one of the categories below:
  - medicinal product not subject to medical prescription
  - medicinal product subject to medical prescription
  - medicinal product subject to special medical prescription
  - medicinal product subject to restricted medical prescription (See Annex I: Summary of Product Characteristics, section 4.2)
  - medicinal product subject to special and restricted medical prescription (See Annex I: Summary of Product Characteristics, section 4.2)

- Annex III.A of the CHMP opinion (Labelling)

The outer packaging should mention either “medicinal product not subject to medical prescription” or “medicinal product subject to medical prescription” (without specifying “restricted” and/or "special")

As regards mock-ups and specimens, the use of any 'sub-category' at national level (e.g. renewable/non-renewable) and the information required to express this, should be addressed in the blue box (see also "When shall I submit mock-ups and/or specimens?").

This information may concern either one, or more, 'sub-categories' listed in Article 70(2) of Directive 2001/83/EC as amended, or a specific way of conveying particular information about the Legal Status. Some Member States use symbols or expressions/specific wordings. Such symbols or expressions are set out in the Annex to the "guideline on the packaging information of medicinal products for human use authorised by the Community". The EMA strongly advises Applicants to follow this guideline since compliance with the guideline ensures compliance with Community legislation.

### 6.3. Change of Legal Status

According to Article 74 of Directive 2001/83/EC as amended, when new facts are brought to its attention, the EMA shall examine and, as appropriate, amend the classification of a centrally authorised medicinal product, by applying the criteria listed in Article 71 of that Directive.

The data requirements for an application to change the classification for the supply of a medicinal product from to prescription to non-prescription ("Switch") are outlined in Part 2 of the "Guideline on changing the classification for the supply of a medicinal product for human use".

In addition, according to Article 74a of the same Directive, a change of classification may benefit from one year of protection. This 1-year period of protection covers significant pre-clinical tests or clinical trials carried out for the purpose of substantiating an application for a change of classification.
Commission decisions authorising a change of classification will contain a clear statement of whether the change is based on significant pre-clinical tests or clinical trials. A change of classification authorised after 20 November 2005 may benefit from this year of protection.

Further information on Legal Status is provided in the “Guideline on Legal Status for the supply to the patient of centrally authorised medicinal products” (EMEA/186279/2006).

References

- Regulation (EC) No 726/2004
- Directive 2001/83/EC
- Guideline on Legal Status for the supply to the patient of centrally authorised medicinal products (EMEA/186279/2006)
- “Guideline on the packaging information of medicinal products for human use authorised by the Community”, the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2C
- “Guideline on changing the classification for the supply of a medicinal product for human use”, the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2C
7. What is the procedure for appointment of CHMP Rapporteur/Co-Rapporteur and their assessment teams?

Rev. March 13

7.1. General principles

For any scientific evaluation in respect of a procedure, a Rapporteur and if relevant a Co-Rapporteur shall be appointed from amongst the members of the Committee for Medicinal Products for Human Use (CHMP) and alternate members. In addition for activities covering all aspects of the risk management of the use of human medicinal products a Rapporteur and if relevant a Co-Rapporteur shall be appointed from amongst the members of the Pharmacovigilance Risk Assessment Committee (PRAC) and alternate members. For Advanced Therapy Medicinal Products (ATMP) a Rapporteur and if relevant a Co-Rapporteur shall be appointed from amongst the members of the Committee for Advanced Therapies (CAT) and alternate members. In addition two CHMP Co-ordinators will be appointed (one supporting the CAT Rapporteur assessment team and another supporting the CAT Co-Rapporteur assessment team).

The appointment of any Rapporteur/Co-Rapporteur is made on the basis of objective criteria, which will ensure the provision of objective scientific opinions and will allow the use of the best and available expertise in the European Economic Area (EEA) on the relevant scientific area.

7.1.1. Requesting the appointment of CHMP/PRAC/CAT Rapporteurs/Co-Rapporteurs and their assessment teams

Applicants shall request the appointment of CHMP/PRAC/CAT Rapporteurs/Co-Rapporteurs (in the following only described as (Co-) Rapporteurs) by sending a completed Pre-submission request form (selecting the indent “Intent to submit MA”) to pa-bus@ema.europa.eu. The pre-submission request form can be accompanied by a cover letter. This notification is also called the “letter of intent”.

We advise applicants to notify the EMA of their intent to submit and request assignment of (Co-) Rapporteurs 7 months prior to the intended submission date. Although applicants may submit the letter of intent earlier than 7 months prior to the intended submission date the (Co-) Rapporteurs appointment procedure will not be initiated prior to that date.

Intended MAA submission dates must be as realistic and accurate as possible as such information is crucial to the EMA and to the future appointed (Co-) Rapporteurs and their assessment teams for planning purposes.

The (Co-) Rapporteurs appointment procedure takes one month and applicants are notified about the outcome. It is the responsibility of the applicant to liaise with the EMA in due course to confirm its intended submission date and request (Co-) Rapporteurs appointment.

For submission deadlines for letters of intent see Q&A 7a.

Please be aware that separate pre-submission forms have to be submitted for requesting eligibility and the appointment of (Co-) Rapporteurs (selecting the corresponding indents on the first page of the pre-submission form), even if an applicant submits both requests in parallel.

Please note that the Applicant’s proposals/preferences are not considered for the appointment of (Co-) Rapporteurs.
7.2. Appointment of (Co-) Rapporteurs and their assessment teams for different application types / procedures

7.2.1. Full applications

In the pre-authorisation phase of a full Marketing Authorisation Application (MAA), two Rapporteurs (i.e. a Rapporteur and a Co-Rapporteur) are appointed. The two Rapporteurs are usually members/alternate members of the CHMP, except for ATMPs, where the Rapporteur and Co-Rapporteur are appointed amongst the CAT members/alternate members with two Co-ordinators appointed from the CHMP.

Furthermore a PRAC Rapporteur and a Co-Rapporteur will be appointed.

7.2.2. Generic/hybrid medicinal products

Due to the particularities of generic/hybrid applications (e.g. legal basis, data requirements), the following principles are considered for the appointment of CHMP/PRAC Rapporteurs/Co-Rapporteurs and their assessment teams:

- A CHMP Rapporteur is appointed for the scientific evaluation of a generic/hybrid medicinal product. For the scientific evaluation of a generic application there is usually no Co-Rapporteur required.
- For the scientific evaluation of a hybrid medicinal product the appointment of a Co-Rapporteur is considered on a case-by-case basis (depending on the particularity of the applied hybrid medicinal product).
- For a generic/hybrid medicinal product, a CHMP pharmacovigilance (PhV) Rapporteur is appointed. The CHMP PhV Rapporteur is the same CHMP member/alternate as the CHMP Rapporteur of the reference medicinal product as applicable.
- Furthermore a PRAC Rapporteur will be appointed.

7.2.3. Similar biological medicinal products

For the scientific evaluation of a similar biological medicinal product CHMP and PRAC Rapporteurs and Co-Rapporteurs will be appointed.

7.2.4. Non-prescription medicinal products

Due to the particularities of non-prescription medicinal products (e.g. self-care environment, data requirements), the following principles are considered for the appointment of CHMP/PRAC Rapporteurs/Co-Rapporteurs and their assessment teams:

- For the scientific evaluation of a non-prescription medicinal product CHMP and PRAC Rapporteurs and Co-Rapporteurs shall be appointed.
  - In the pre-authorisation phase the CHMP/PRAC Rapporteurs and Co-Rapporteurs shall be involved.
  - In the post-authorisation phase, when a change in legal status is foreseen (e.g. switch from prescription to non-prescription), a CHMP peer reviewer shall be appointed to work with the existing CHMP/PRAC Rapporteurs and Co-Rapporteurs already in place for the given medicinal product.
7.2.5. Re-examination of a CHMP opinion

In cases of re-examination of a CHMP opinion a CHMP/CAT Rapporteur and a Co-Rapporteur shall be appointed. In case a PRAC Rapporteur is deemed necessary, he/she will be appointed. For CHMP opinions where the CHMP/CAT Co-Rapporteur was not involved in the initial evaluation, no re-examination Co-Rapporteur needs to be appointed. A different CHMP/CAT Rapporteur and, where applicable, a different CHMP/CAT Co-Rapporteur from those appointed for the initial evaluation shall be appointed in order to adequately assess the grounds for the re-examination of the CHMP opinion. These Rapporteurs will coordinate the evaluation for the duration of the re-examination procedure only.

The Rapporteur, Co-Rapporteur (if applicable) appointment process will be initiated as soon as the EMA/CHMP receives written notice that the applicant/MAH wishes to request a re-examination of the CHMP opinion.

7.2.6. Ancillary medicinal substances or ancillary human blood derivatives incorporated in medical devices

The notified body is requested to submit the letter of intent at least 6 months before the expected date of submission.

A CHMP Rapporteur and Co-Rapporteur, if appropriate, will be appointed.

References

- PRAC Rules of Procedure (EMA/PRAC/567515/2012)
- Regulation (EC) No 726/2004
- Centralised Procedure, the Rules governing Medicinal Products in the European Community, Volume 2A, Notice to Applicants, Chapter 4
8. How should I notify a change in the intended submission date of my application? *NEW May 2015*

In case the previously indicated submission date of an upcoming application for marketing authorisation is changed, the applicant shall inform the EMA by re-sending to pa-bus@ema.europa.eu the completed Pre-Submission request form (pre-submission request form), where the scope of request should be selected as “notification of change” and the new intended date of submission indicated in the corresponding field. The text of the e-mail should also describe the type of the change requested.

The change in intended MAA submission date must be notified as soon as possible. Since this information is crucial to the EMA and to the appointed (Co-) Rapporteurs and their assessment teams for planning purposes the intended submission date should be accurate and realistic. In some cases, a change in the planned submission data could lead to re-appointment of one or several Rapporteurs, if the previously appointed Rapporteur(s) will not be able to perform the assessment according to the new timings. In such case the applicants will be informed accordingly.

**References**

- Pre-submission request form
9. What is the role of the EMA product team? NEW Apr 2015

An EMA 'Product Team' is set up for each medicinal product submitted through the centralised procedure. The Product Team is responsible for providing support to the evaluation activities of the EMA scientific committees. In particular this includes:

- Provision of procedural guidance concerning all pre authorisation activities directly preceding the application and liaison with the (Co-)Rapporteurs in the conduct of such activities;
- Provision of advice to (Co-)Rapporteurs/committee members/applicant concerning all questions of a regulatory or procedural nature;
- Provision of advice to the applicant in the technical preparation of the marketing authorisation application and subsequent validation of such applications;
- In collaboration with the (Co-)Rapporteurs assessment teams production of the List of Questions, List of Outstanding Issues, draft summary of product characteristics to support committees discussion/adoption:
- Supporting the (Co-)Rapporteurs with regulatory, technical advice in briefing / debriefing / clarification meetings with applicants;
- To support planning and conduct of oral explanations, ad-hoc expert groups, referral to Working Parties, Scientific Advisory Groups etc;
- Managing the timeframe of the procedure to ensure it remains within legal timeframe;
- Co-ordinating the linguistic check of product information to ensure consistency and high quality;
- Informing the (Co-)Rapporteurs on elements of regulatory and scientific consistency of the application of quality, safety, efficacy and guidelines in the conduct of the evaluation procedure;
- To prepare the committee assessment report and subsequent Summary of Opinion (SMOP) and European Public Assessment Report (EPAR).

The Product Team is established during the pre-submission phase of the initial marketing authorisation application and is in place post-authorisation. It ensures oversight of all elements of product knowledge through the complementary contributions of the various team members. The composition of the team is adapted over time depending on the complexity of the product and procedure as well as the type of issues raised during the product’s lifecycle. From an applicant’s perspective the following team members are particularly relevant:

- the **procedure manager**, or PM, to oversee all aspects of the management of specific procedures. Procedure managers ensure regulatory consistency at EMA and are responsible for managing the regulatory process for each application. The PM is supported by the **procedure assistant (PA)** in terms of administrative and secretarial aspects.
- the **EMA product lead**, or EPL, to maintain oversight of a medicine as it moves through the different stages of its lifecycle.

The applicant will be notified of the appointed PM, including their contact details via the eligibility outcome letter and of the EPL in the CHMP Rapporteur appointment letter. Any subsequent change to

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3 Based on the responsibilities as defined in Notice to Applicants Volume 2A, Chapter 4 “Centralised procedure”, April 2006.
the resource allocation for these functions will be communicated to the applicant/marketing authorisation holder.

Further team members assigned for each product are representing the functions of quality, risk management, labeling review and regulatory affairs. Specialised functions, like inspections and signal validation, will be involved as required.

Please see other relevant questions and answers in the EMA pre-authorisation guidance “Who is my contact at the European Medicines Agency during a marketing authorisation application (MAA) evaluation procedure?” and in the EMA post-authorisation guidance “Who is my contact at the European Medicines Agency during post-authorisation procedures?”, “Who is my contact at the European Medicines Agency during an application procedure for extension of indication?” and “Who is my contact at the European Medicines Agency during the post-authorisation phase outside any evaluation procedures?”. 
10. **Who is my contact at the European Medicines Agency during a marketing authorisation application (MAA) evaluation procedure? **NEW Apr 2015

In the context of an initial marketing authorisation application (MAA) evaluation in the centralised procedure, the procedure manager (PM) is the primary contact for the applicant prior to submission and throughout the procedure until the decision is granted by the European Commission.

The applicant will be notified of the allocated PM at time of confirmation of eligibility to the centralised procedure.

The PM will serve as the main liaison person between the EMA product team, the Rapporteurs and the applicant. The PM, in close co-operation with the EMA Product Lead (EPL) and the rapporteurs, will ensure that the applicant is kept informed of all aspects related to the MAA evaluation.

The applicant should contact the **PM** for all questions regarding the evaluation procedure, including:

- Requests for guidance in the pre-submission phase, such as the pre-submission meeting;
- Any type of procedural questions during the evaluation, such as availability of assessment reports and Opinion documents;
- Discussion on timetables including requests for extension of clock-stops;
- Any question where guidance related to the evaluation procedure is needed; in such cases the PM will address or liaise and redirect as appropriate.

Questions concerning the **validation** of the MAA, once submitted, will be dealt with by an assigned Validation Officer.

At **certain milestones** during the evaluation procedure, the **EPL** will contact the applicant for a direct exchange to facilitate the discussion on the scientific evaluation. These include:

- Preparation and conduct of clarification meetings (where applicant requests such meeting);
- Immediate feedback regarding scientific aspects from committee plenary discussions, where required;
- Expectations relating to the Oral Explanation, including topics to be addressed;
- Discussion of required post-authorisation measures;
- Late-stage revisions of the product information before adoption of the final Opinion.

These interactions occur in close co-operation with the Rapporteurs. Occasionally other members from the EMA Product team may contact the applicant directly to facilitate the discussion on specific aspects (e.g. quality, risk management, mock-up review).

Where the applicant is in direct contact with the EPL or another member of the EMA Product Team the PM should always be copied in the correspondence.

Please see other relevant questions and answers in the EMA pre-authorisation guidance “What is the role of the EMA product team?” and in the EMA post-authorisation guidance “Who is my contact at the European Medicines Agency during post-authorisation procedures?”, “Who is my contact at the European Medicines Agency during an application procedure for extension of indication?” and “Who is my contact at the European Medicines Agency during the post-authorisation phase outside any evaluation procedures?”.
11. Is my product eligible for an Accelerated Assessment

11.1. Legal basis and general principles

According to Articles 6(3) and 7c of Regulation (EC) No 726/2004, the maximum timeframe for the evaluation of a marketing authorisation application under the Centralised Procedure is 210 days, excluding clock stops when additional written or oral information is to be provided by the applicant in response to questions asked by the CHMP.

However, according to Recital 33 and Article 14(9) of Regulation (EC) No 726/2004, the applicant may request an accelerated assessment procedure in order to meet, in particular the legitimate expectations of patients and to take account of the increasingly rapid progress of science and therapies, for medicinal products of major interest from the point of view of public health and in particular from the viewpoint of therapeutic innovation.

Applicants requesting an accelerated assessment procedure should justify that the medicinal product is expected to be of major public health interest. Based on the request, the justifications presented, and the recommendations of the Rapporteurs, the CHMP will formulate a decision. Such a decision will be taken without prejudice to the CHMP opinion (positive or negative) on the granting of a marketing authorisation.

If the CHMP accepts the request, the timeframe for the evaluation will be reduced to **150 days**.

11.2. Request for an accelerated assessment: timing and justification

Any request for accelerated assessment should be made as early as possible before the actual submission of the marketing authorisation application. The request, including the justification, should be sent electronically to the EMA product team leader and all CHMP members at least 10 working days in advance of the CHMP meeting preceding the intended start of the centralised procedure. In practice, submission of the request will generally occur at least between 10 to 30 days before the intended start of the procedure.

Applicants requesting an accelerated assessment procedure should duly substantiate the request and in particular, justify their expectation that the medicinal product is of major public health interest particularly from the point of view of therapeutic innovation. There is no single definition of what constitutes major public health interest. This should be justified by the applicant on a case-by-case basis.

The justification should include the major benefits expected and present the arguments to support the claim that the medicinal product introduces new methods of therapy or improves on existing methods, thereby addressing to a significant extent the greater unmet needs for maintaining and improving public health.

The key items to be described in the justification, and the appropriate level of detail, should be evaluated on a case-by-case basis. The request should be presented as a short but comprehensive document (ideal length 5-10 pages). The following list of key items would normally be addressed in the justification:

- the unmet needs and the available methods of prevention, diagnosis or treatment
- the extent to which the medicinal product is expected to have major impact on medical practice, its major added value, and/or how it addresses the greater unmet needs
- a brief outline of the main available evidence on which the applicant bases its claim of major public health interest

When submitting an accelerated assessment request, the applicant should use the Pre-submission request form (Accelerated Assessment) and Applicant’s justifications, which should be sent, electronically, to: pa-bus@ema.europa.eu.

Alternatively, a paper version can be sent to the following mailing address:

Product and Application Business Support (PA-BUS)
European Medicines Agency EMA
30 Churchill Place
Canary Wharf
London, E14 5EU
UK

Following receipt of the request, the Rapporteurs will produce a briefing note including the Rapporteurs’ recommendations as to the appropriateness of an accelerated assessment. The CHMP will consider the request submitted by the applicant, the Rapporteurs’ recommendations and the views of other CHMP members, in order to conclude on the acceptability or not of the request. If necessary, the CHMP may request clarifications from the applicant about the request. The CHMP conclusions will be communicated to the applicant. The reasons for accepting or rejecting the request will also be summarised in the CHMP assessment report.

If a request for an accelerated assessment procedure is granted, the CHMP will take into consideration the standard timetable agreed for the accelerated assessment procedure (see Section 6 of the “Guideline on the procedure for accelerated assessment pursuant to Article 14(9) of Regulation (EC) No 726/2004” (EMEA/419127/05)).

References

- Regulation (EC) No 726/2004
- “Guideline on the procedure for accelerated assessment pursuant to article 14 (9) of Regulation (EC) No 726/2004” (EMEA/419127/05)
- “Centralised Procedure”, the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2A, Chapter 4
12. If I intend to submit multiple applications for a specific medicinal product? *Rev. Jul 14*

The EMA is regularly approached by applicants wishing to obtain, either simultaneously or successively, more than one Marketing Authorisation for a specific medicinal product, under different invented names.

According to Article 82(1) of Regulation (EC) No 726/2004, the Commission may authorise applicants to submit more than one application to the EMA, when there are objective verifiable reasons relating to public health regarding the availability of medicinal products to health care professionals and/or patients or for co-marketing reasons.

Therefore, applicants will be asked to explain and justify the motives behind multiple applications and their intentions as far as the exploitation of the marketing authorisation is concerned.

In the framework of the article 82(1) of the Regulation, a specific procedure has been agreed between the EMA and the European Commission. Under this procedure Applicants should, approximately **four months** prior to the anticipated date of submission, notify the Commission of their motives for submitting multiple applications and provide the necessary explanation and justification addressing the article 82(1) of the Regulation (EC) No 726/2004 criteria, with a copy to the EMA, addressing either public health reasons or co-marketing reasons.

Such notification should be sent to the following address:

European Commission  
D5 Unit for Medicinal products – authorisations, EMA  
DG Health and Consumers  
Rue Demot 24, DM24 02/128  
B-1049 Bruxelles  
Belgium  

The Commission will consider the situation, liaise with the Applicant(s) where appropriate and inform the Applicant(s) as to whether it would have specific objections to the granting of multiple Marketing Authorisations or not. The company will always need to include this Commission response as Annex 5.16 to the application form, as otherwise the Agency cannot validate such applications.

**Procedural aspects**

Multiple/duplicate applications for a specific medicinal product with an active substance(s) already under assessment via the centralised procedure have automatic access to the centralised procedure. Nevertheless, in all cases the eligibility of a medicinal product for evaluation via the centralised procedure needs to be requested by the applicant by submitting an eligibility request to the EMA. For details see Question 2. This has to be done prior to submission of any dossier and should also include the request for Rapporteur assignment.

For the assessment procedure, the objective is to ensure the adoption of a CHMP Opinion for a multiple application at the same time when the CHMP Opinion for the initial application is adopted. Therefore, for practical reasons, the EMA strongly recommends the following time points for the time for submission of the multiple application(s):

a. In parallel with the initial application submission (day 0)  
b. Submission before the adoption of the list of questions (before the day 120) for the initial application
c. Submission at the time of the response to list of questions (day 121) for the initial application

It should be noted that multiple applications are subject to a full validation as they are stand-alone applications. Therefore, the validation outcome may differ from the one of the original application. Following the positive outcome of the validation, the evaluation of the multiple application(s) will be aligned with that of the ongoing initial application, in case the above timeframes have been duly observed by the applicant. The submission of the multiple application(s) should be done in advance, to allow sufficient time for the validation to be completed by D120 or D121 of the ongoing initial application. The validation period between submission date and start date is 13 EMA working days. Please observe the EMA procedural timetables.

Relevant aspects of the Paediatric legislation should be considered as appropriate for each of the multiple applications submitted. The Risk Management Plans for multiple applications should be product specific and reflect the particulars of each specific application (e.g. product details including differences in indication(s) in case of patent issues, RMP version number and date).

Multiple applications can also be submitted after the Commission Decision on the initial application as stand-alone applications or Informed Consent applications. Again, requirements for eligibility and Rapporteur assignment remain. However, as a rule, an abridged timetable for assessment will be adopted in line with a 60 days procedure. Submission of the application(s) should be done in advance to allow the completion of the validation before the intended start date of the procedure.

Applicants are reminded that multiple applications of the same marketing authorisation holder will be covered by the notion of “global marketing authorisation”.

References

- Directive 2001/83/EC
- Regulation (EC) No 726/2004
- The Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2A, Chapter 1 and Chapter 4.
- EC communication on ‘Handling of Duplicate Marketing Authorisation Applications’ (March 2010) and Update 1 (October 2011)
- EMA procedural timetables

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4 Later submission of the dossier is not recommended due to difficulties with its alignment with the original application.
13. What aspects should I consider at time of submission of my application for marketing authorisation if my medicinal product has been designated as an orphan medicinal product? New Feb 13

If your medicinal product has been designated as an orphan medicinal product, you will have to consider the following points at the time of submission of your application for marketing authorisation:

- The applicant for the marketing authorisation application will have to be the same as the holder of the orphan designation; where necessary, the orphan designation will have to be transferred to the new sponsor in advance of the submission of the application for marketing authorisation. In case the sponsor remains the same person or legal entity but changes its name and/or address, a letter should be sent to the Agency indicating the new name and/or address details and confirming that the identity of the Sponsor remains the same.

- The therapeutic indication requested for your medicinal product will have to fall within the scope of the orphan designation, i.e. the therapeutic indication applied for cannot be broader than the orphan indication. Reference is made on this regard to Article 7(3) of Regulation (EC) No 141/2000 (“Orphan Regulation”), which provides that the marketing authorisation granted for an orphan medicinal product shall cover only those therapeutic indications which fulfill the criteria for designation set out in Article 3.

- It is not possible to combine within the same application for marketing authorisation orphan and non-orphan indications. However, this is without prejudice to the possibility of applying for a separate marketing authorisation for other indications which have not been designated as orphan, as provided for in the Orphan Regulation.

- You will have to submit at the same time as the submission of the initial application for marketing authorisation, a report on the maintenance of the orphan designation criteria, which will be reviewed by the COMP. This report should be addressed to the Head of the Orphan Medicines Section.

References

- Regulation (EC) No 141/2000 on orphan medicinal products
- Guideline on the format and content of applications for designation as orphan medicinal products and on the transfer of designations from one sponsor to another
- Sponsor’s report on the maintenance of the designation criteria at the time of marketing authorisation applications for a designated orphan medicinal product
14. What aspects should I consider if the designation for my orphan medicinal product is still pending at the time of submission of my application for marketing authorisation?

New Feb 13

When an application for orphan designation is still pending at time of submission of the application for marketing authorisation, it is nevertheless possible for the medicinal product to be authorised as an orphan medicinal product, provided that the orphan designation is granted and confirmed by the COMP before the granting of the marketing authorisation.

However, in such cases, the eligibility to the centralised procedure (which precedes the submission of the application for marketing authorisation) cannot be based on Article 3(1), Annex 4 – Orphan designated medicinal product. Similarly, a fee reduction will not be applicable, as it can only be considered if orphan designation has already been granted at the time of submission of the application for marketing authorisation.

References

- Explanatory note on fees payable to the European Medicines Agency
15. What aspects should I consider at time of submission of my application for marketing authorisation if there are orphan medicinal products designated or authorised for a condition related to my proposed therapeutic indication? Rev. Feb 15

In advance of submission of your application for marketing authorisation, irrespective of whether your medicinal product has been designated as orphan or not, you are advised to check the Community register of orphan medicinal products, for information on medicinal products designated as orphan which are under market exclusivity protection.

You will have to indicate in the application form (section 1.2.2) if any medicinal product has been designated as an orphan medicinal product for a condition relating to the therapeutic indication proposed in your application and, if applicable, specify the respective orphan designation number.

If any of the designated orphan medicinal products has been granted a marketing authorisation in the Union, and a period of market exclusivity is in force, you will have to provide in Module 1.7.1 a similarity report addressing the possible similarity between your medicinal products and the orphan medicinal product(s) which have received a marketing authorisation.

This legal requirement arises from Article 8(1) of the Orphan Regulation which provides that where a marketing authorisation in respect of an orphan medicinal product is granted, the Agency and the Member States shall not, for a period of 10 years, accept another application for a marketing authorisation, or grant a marketing authorisation or accept an application to extend an existing marketing authorisation, for the same therapeutic indication, in respect of a similar medicinal product.

Article 3 of Commission Regulation (EC) No 847/2000 defines similar medicinal product as a medicinal product containing a similar active substance or substances as contained in a currently authorised orphan medicinal product, and which is intended for the same therapeutic indication.

It also defines similar active substance as an identical active substance, or an active substance with the same principal molecular structural features (but not necessarily all of the same molecular features) and which acts via the same mechanism.

Based on the above mentioned definitions, the assessment of similarity between two medicinal products takes into consideration the following criteria

- Principal molecular structural features,
- Mechanism of action and
- Therapeutic indication.

If significant differences exist within one or more of these criteria, the two products will not be considered as similar. These criteria are explained in the Guideline on aspects of the application of Article 8(1) and 8(3) of Regulation (EC) No 141/2000: Assessing similarity of medicinal products versus authorised orphan medicinal products benefiting from market exclusivity and applying derogations from that market exclusivity.

For information on the procedure and timetable for assessment of similarity and, where applicable, derogation report vis-à-vis authorised orphan medicinal products, please refer to Q&A "What is the procedure and timetable for assessment of similarity and, where applicable, derogation report vis-à-vis authorised orphan medicinal products?".

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Please note that if the Agency identifies a possible similarity issue not addressed by the applicant before validation, the applicant will be asked to complete the application with information on similarity and, if applicable, on one of the derogations. Validation of the application will only proceed once the applicant has submitted a report justifying the lack of similarity or, if similar, additional information justifying one of the derogations in Article 8(3).

As considerable time may elapse between validation of an application and adoption of an opinion, if applicants become aware of medicinal products which have been authorised as orphans for a condition related to the therapeutic indication proposed in their application, this information should be communicated promptly to their Procedure Manager at the Agency in order to arrange for the submission of updated application form and modules 1.7.1 and 1.7.2, as applicable.

In any case, the Agency will check at certain milestones of the procedure, i.e. Day 120, Day 180 and prior to adoption of a CHMP opinion whether new orphan medicinal products have been authorised for the same condition.

References

- Regulation (EC) No 141/2000 on orphan medicinal products
- Regulation (EC) No 847/2000
- Community register of orphan medicinal products
- Guideline on aspects of the application of Article 8(1) and 8(3) of Regulation (EC) No 141/2000: Assessing similarity of medicinal products versus authorised orphan medicinal products benefiting from market exclusivity and applying derogations from that market exclusivity
16. What aspects should I consider if my medicinal product is considered similar to an orphan medicinal product? Rev. Feb 15

If your product is considered to be similar to any authorised orphan medicinal product, you will have to provide in Module 1.7.2 justification that one of the following derogations, laid down in Article 8(3) of the Orphan Regulation applies, i.e.

(a) the holder of the marketing authorisation for the orphan medicinal product has given his consent for submission of your application, in which case a signed letter from the MAH of the orphan medicinal product should be provided confirming the consent for submission of an application for marketing authorisation;

(b) the holder of the marketing authorisation for the orphan medicinal product is unable to supply sufficient quantities of the medicinal product, in which case the applicant should provide a report including details of the supply shortage and justify that patients’ needs in the orphan indication are not being met;

(c) the applicant can establish that their product, although similar to the orphan medicinal product already authorised, is more effective, safer or otherwise clinically superior, in which case a critical report justifying clinical superiority to the authorised product must be provided.

For information on the procedure and timetable for assessment of derogation report vis-à-vis authorised orphan medicinal products, please refer to Q&A "What is the procedure and timetable for assessment of similarity and, where applicable, derogation report vis-à-vis authorised orphan medicinal products?"

Please note that if the Agency identifies a possible similarity issue not addressed by the applicant before validation, the applicant will be asked to complete the application with information on similarity and, if applicable, on one of the derogations. Validation of the application will only proceed once the applicant has submitted a report justifying the lack of similarity or, if similar, additional information justifying one of the derogations in Article 8(3).

As considerable time may elapse between validation of an application and adoption of an opinion, if applicants become aware of medicinal products which have been authorised as orphans for a condition related to the therapeutic indication proposed in their application, this information should be communicated promptly to their Procedure Manager at the Agency in order to arrange for the submission of updated application form and modules 1.7.1 and 1.7.2, as applicable.

In any case, the Agency will check at certain milestones of the procedure, i.e. Day 120, Day 180 and prior to adoption of a CHMP opinion whether new orphan medicinal products have been authorised for the same condition.

References

- Regulation (EC) No 141/2000 on orphan medicinal products
- Regulation (EC) No 847/2000
- Community register of orphan medicinal products
• Guideline on aspects of the application of Article 8(1) and 8(3) of Regulation (EC) No 141/2000: Assessing similarity of medicinal products versus authorised orphan medicinal products benefitting from market exclusivity and applying derogations from that market exclusivity
17. What is the procedure and timetable for assessment of similarity and, where applicable, derogation report vis-à-vis authorised orphan medicinal products? New Feb 13

The assessment of similarity and, where applicable, of the derogation report vis-à-vis authorised orphan medicinal products will be conducted by the CHMP Rapporteur and Co-Rapporteur in charge of assessing the quality, safety and efficacy of your medicinal product.

This assessment of similarity is conducted in parallel to the evaluation of the application for marketing authorisation or extension of the marketing authorisation, as applicable, and normally follows a 60 day timetable. This assessment includes the consultation of the Quality Working Party or the Biologicals Working Party, as appropriate, for the aspects concerning the similarity of the molecular structures of the products.

Where necessary, a list of questions will be adopted by the CHMP on Day 60 and a timetable of 30 days applies, normally, for assessment of the responses to the questions raised.

Where the outcome of the CHMP assessment is that the medicinal products are considered similar, the applicant will be requested to provide a justification that one of the derogations in Article 8(3) is fulfilled. This assessment will follow also a 60 day timetable with a possibility for raising questions to the applicant.

Where the CHMP concludes that the application for marketing authorisation is not similar to an authorised orphan medicinal product or, if similar, that one of the derogations claimed by the applicant applies, this will not prevent the granting of the marketing authorisation / extension to the marketing authorisation, provided that the quality, safety and efficacy of the medicinal product are demonstrated.

Should the CHMP conclude that the product which is the subject of the application for marketing authorisation is considered similar to an authorised orphan medicinal product and none of the derogations provided for in Article 8(3) of the Orphan Regulation applies, the CHMP will adopt an opinion recommending the refusal of the granting of the marketing authorisation/extension to the marketing authorisation, irrespective of the demonstration of the quality, safety or efficacy of the medicinal product.

References

- Regulation (EC) No 141/2000 on orphan medicinal products
- Regulation (EC) No 847/2000
- Guideline on aspects of the application of Article 8(1) and 8(3) of Regulation (EC) No 141/2000: Assessing similarity of medicinal products versus authorised orphan medicinal products benefiting from market exclusivity and applying derogations from that market exclusivity
18. What fee do I have to pay and how is the appropriate fee for my application calculated? Rev. Dec 10

Fees for obtaining and maintaining a Community authorisation to market medicinal products for human use are levied in accordance with Regulation (EC) No 297/95.

Since 1 December 2005, the EMA will issue an invoice on the date of the notification of the administrative validation to the applicant and fees will be payable in EURO within 45 calendar days of the date of the said notification. The invoice is sent to the billing address indicated by the Applicant, and will contain clear details of the product and procedures involved, the type of fee, the amount of the fee, the bank account to where the fee should be paid and the due date for payment.

Where more than one procedure is processed in a given month a summary invoice or statement is issued at the end of each month for payment within 30 days of the end of the month.

If the application cannot be validated, the EMA will issue an invoice on the date of the notification of the administrative non-validation to the applicant for an administrative charge to cover administrative costs.

Further details on EMA fees can be obtained in the "Explanatory Note on fees payable to the European Medicines Agency" (EMA/348317/2010)

Where an applicant disagrees on the classification by the EMA of an application under one of the fee categories described in the ‘Fee Regulation’, the following procedure may apply:

- Any disagreement should be sent to the Executive Director accompanied by the appropriate justification, at the latest two weeks after receipt of the invoice indicating the fees payable to the EMA.

- The Executive Director will take a decision following consultation with the competent committee.

The EMA contacts point for queries on Fees, Procedures or Application numbers, are:

Product and Application Business Support (PA-BUS) or e-mail address: pa-bus@ema.europa.eu

References

- Regulation (EC) No 297/95

- "Centralised Procedure", the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2A, Chapter 4

- "Explanatory Note on fees payable to the European Medicines Agency"

- European Medicines Agency Management Board "Rules for the implementation of Regulation (EC) No 297/95 as amended on fees payable to the European Medicines Agency and other measures" (EMA/MB/818152/2009)
19. What definition of strength is used for the calculation of fees? Rev. Oct 12

The "Guideline on the categorisation of New Applications versus Variation Applications" describes the agreement reached as to the use of the same definitions of strength in case of applications submitted through the Centralised Procedure and the Mutual Recognition Procedure.

This definition will be taken into account for the calculation of fees as well as for the numbering system used by both the EMA and the Commission (see "Payment of fees" and "Management of applications").

The following definitions therefore apply:

- For single-dose preparations, total use, the strength is defined as the amount of active substance per unit dose
- For single-dose preparations, partial use, the strength is defined as the concentration expressed as the amount of active substance per ml, per puff, per drop, per kg, per m², in percentage as appropriate
- For multi-dose preparations, the strength is defined as the concentration expressed as the amount of active substance per ml, per puff, per drop, per kg, per m², as appropriate
- For powder for reconstitution (powder for oral solution or suspension, powder for solution for injection, etc.) the strength is defined as the concentration after dissolution or suspension (reconstitution) to the volume and liquid recommended
- For concentrates for solutions (for injection or for infusion) the strength is defined as the concentration of the concentrate before dilution
- For transdermal patches, the strength is defined as the amount of active substance released from the patch in 24h

Please note that no additional strengths or presentations can be applied for by the applicant after the validation of the application and payment of the fee. Such changes can be introduced after the marketing authorisation has been granted through a variation procedure.

References

- "Guideline on the categorisation of New Applications versus Variation Applications", the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2C
20. What is the fee for a GMP/GCP inspection? Rev. March 13

For all inspections requested by the CHMP in respect of an application under the Centralised Procedure fees are payable by the applicant under Regulation (EC) No 297/95, as amended.

For information on the fee applicable for inspections, please refer to the Explanatory note on fees payable to the European Medicines Agency.

Invoices are issued within 20 days of the confirmation of the inspection dates by the relevant inspectors and are sent by registered post to the applicant. Applicants for marketing authorisations under the centralised procedure are required to pay the total fee charged within 45 days from the date on which the inspection is carried out. (Important: Invoice reference number to be mentioned with each payment).

For inspections outside the EEA/European Union the applicant is also required to pay the travel and accommodation expenses of the Inspector(s) and any Experts or Rapporteur involved in carrying out the inspection(s). These expenses are to be paid directly by the applicant to the inspector’s Authorities.

In the case of concurrent multiple applications for exactly the same medicinal product, the applicants may agree between themselves that one of them will be regarded as the “lead” applicant for the purpose of inspections. The identity of the “lead” applicant must be notified in writing to the EMA at the pre-submission stage. In such a case, the multiple applications will be treated as a single one and the total fee(s) will be charged to the “lead” applicant.

Where an inspection that has been formally notified to the applicant (and an invoice has been issued) is cancelled due to the withdrawal of or change to an application at any stage in the processing of the application, the applicant will be liable for 50% payment of inspection fee(s) as follows:

- Applicant decides to withdraw the application.
- In the context of GMP inspection, change to manufacturing arrangements by the manufacturer necessitating cancellation of the inspection, agreed at any time before the inspection is carried out.
- In the context of GCP inspection, change to the scope of the application or submitted data, or access to, ownership of, or location of facilities or data necessitating cancellation of the inspection, agreed at any time before the inspection is carried out.

Where the cancelled inspection was to take place outside the EEA/European Union, the applicant will be liable for any travel expenses already incurred by the inspectors at the date of cancellation for which they are not able to obtain reimbursement.

For more information on inspection fees, please refer to the EMA website: Regulatory\Human medicines\Fees

References

- Regulation (EC) 297/95, as amended
- Explanatory note on fees payable to the European Medicines Agency
- Rules for the implementation of Council Regulation (EC) No 297/95 on fees payable to the European Medicines Agency and other measures
21. When could a fee waiver / fee reduction be granted?

Rev. Oct 14

Applicants may benefit from fee incentives if at the time of the administrative validation the application or the applicant itself meets the criteria for fee reduction or deferral. Any changes which may take place after validation, would not retrospectively affect the levied fee.

Under article 7(2) of Regulation (EC) No 141/2000 on orphan medicinal products, total or partial fee exemptions may be granted by the EMA, for medicinal products designated as “orphan” by the European Commission on recommendation from the Committee on Orphan Medicinal Products. This includes fees for pre-authorisation activities such as protocol assistance (scientific advice), and for products using the centralised procedure: the application for marketing authorisation, inspections and post-authorisation activities such as variations, annual fees, etc.

Each year funds are made available by the EU Budgetary Authority to grant fee exemptions for designated orphan medicinal products. Subject to the availability of funds, the Executive Director will decide at the beginning of each year on the percentage of fee reductions to be granted that year.

Fees incentives for orphan medicinal products are automatically granted and sponsors of orphan medicinal products do not need to apply for such incentives.

It should be noted that fee reductions can only be granted once a decision on orphan medicinal product designation has been adopted by the European Commission. In addition, the application should fall within the scope of the orphan condition. The applicant or marketing authorisation holder requesting the fee reduction must be the sponsor of the designation. If this is not the case, the sponsorship of the designation should be transferred prior to submitting the request.

Further information on the applicable fee reductions for an orphan medicinal product is provided in the EMA Public Statement on Fee reductions for Designated Orphan Medicinal Products (EMEA/622074/2013).

Applicants which meet the definition of a micro, small or medium-sized enterprise (SMEs) as set out in Commission Recommendation 2003/361/EC of 6 May 2003, are eligible for certain fee reductions from the EMA. This includes fee reductions for scientific advice, pre- and post-authorisation inspections, scientific services, and a full fee waiver for administrative services (with the exception of parallel distribution).

Deferral of the fee payable for the application for marketing authorisation or related inspections may also apply.

It should be noted that fee reductions and deferrals can only be considered once the applicant has been assigned SME status by the EMA. SME applicants wishing to receive a written confirmation of fee incentive should address an e-mail to the EMA’s SME Office (sme@ema.europa.eu).

Fee reductions may also be granted by the EMA Executive Director in exceptional circumstances and for imperative reasons of public or animal health, after consultation of the competent committee, in accordance with Article 9 of Regulation (EC) No 297/95, as amended. In such circumstances applicants should liaise with the Agency.

References

- Regulation (EC) No 297/95
• Regulation (EC) No 141/2000
• Regulation (EC) No 2049/2005
• Commission Recommendation 2003/361/EC
• Fees payable to the European Medicines Agency
• EMEA Public Statement on Fee Reductions for Designated Orphan Medicinal Products (EMEA/622074/2013)
22. When shall I submit mock-ups and/or specimens?

Rev. Feb 15

Mock-ups and specimens of the outer and immediate packaging together with the package leaflet must be submitted by the applicant/MAH to the EMA for review, before commercialisation of the medicinal product.

A "Mock-up" is a copy of the flat artwork design in full colour, presented so that, following cutting and folding where necessary, it provides a replica of both the outer and immediate packaging, so that the three dimensional presentation of the labelling text is clear.

A "Specimen" is a sample of the actual printed outer and immediate packaging materials and package leaflet (i.e. the sales presentation).

The checking process of mock-ups and specimens in the Centralised Procedure is based on the following general principles:

- The European medicines Agency (EMA), through the translations checking policy, will ensure that high-quality product information in all EU languages, as prepared by the MAH and checked by the Member States prior to the granting of the MA, is included in Commission Decisions on centrally authorised medicinal products;
- MAHs are responsible for the correct implementation of the agreed product information texts in their printed packaging materials, in line with the Commission Decision and relevant EU legislation;
- The EMA will not perform a detailed linguistic check of mock-ups and specimens, but rather a general check from the viewpoint of readability in order to contribute to the safe use of medicines;
- The EMA can, at any time, request specific specimens from the MAH for review (e.g. further to a safety-related or product defect issue).

Based on the above, EMA will not check the national requirements included in the blue box. However, the fact that the mock-up has to be a real example of the sales presentation implies that the mock-up should indicate how the information specifically required by Member States (such as price, reimbursement, legal status, identification and authenticity) will be presented in the 'blue box'. This means that if at the time of submission of the mock-ups this specific information is not yet known, at least an indication should be given of the way in which this information will be presented in the ‘blue box’ on the outer packaging i.e. the blue outline of the ‘blue box’ should be displayed to show the location of the ‘blue box’ on the outer carton.

Details on the ‘blue box’ content, for each Member States, are given in the Annex of the "Guideline on the packaging information of medicinal products for human use authorised by the Union" as published by the European Commission in the Notice to Applicants, Volume 2C.

The inclusion of a national barcode on the labelling would normally be viewed as a Member State driven requirement located within the ‘blue box’ on the outer carton. However, EMA can also accept the inclusion of a national barcode on the immediate packaging (e.g. for traceability purposes), where space and readability permit.

Applicants should provide the EMA with mock-ups and/or specimens for new applications in accordance with the following requirements:

Mock-ups
• At the time of submission of the application, one English colour full-size mock-up and one multi-lingual colour full-size mock-up ("worst-case") of the outer and immediate packaging for each pharmaceutical form in each container type in the smallest pack-size must be included in Module 1.3.2 of the application. Mock-ups of the package leaflet may be included (optional).

• By day 121, revised mock-ups of the labelling and package leaflet should be submitted within Module 1.3.2 as part of the answers to the list of questions, in case of comments or in case the applicant changes the overall design.

• By day 181, further mock-ups may need to be submitted if there any outstanding comments made at Day 150 to be solved prior to the opinion.

• The applicant will liaise with the EMA by e-mail to muspecimens@ema.europa.eu to resolve any outstanding comment on the mock-ups of labelling and package leaflet prior to the adoption of the opinion.

• Submission of further mock-ups for review is not required after adoption of the Opinion. However, EMA would be willing to perform an additional review of updated mock-ups in the post-opinion phase, if requested by applicants prior to specimen printing.

Specimens

• At the latest 15 working days before marketing, one set of relevant specimens examples of the outer and immediate packaging and package leaflet for each strength or for each different total content per total volume (when the strength is expressed as concentration per unit volume (x mg/ml)) and each pharmaceutical form in each container type need to be provided to the EMA (using the "Specimen Submission Form" (see EMEA/305821/2006):
  – before first marketing in the EU,
  – before first marketing as a multi-lingual pack (if different from the first specimens sent to the EMA),
  – when any other multi-lingual pack is marketed with a higher number of languages than the multi-lingual pack(s) previously reviewed.

The EMA will perform a general check from the viewpoint of readability within 15 working days, and will check if any previous comments on mock-ups/specimens have been duly implemented. The applicant will be informed about the outcome of the check.

For any questions on the checking process or to discuss upcoming mock-up/specimen submissions please contact the EMA on: muspecimens@ema.europa.eu.

References

• Directive 2001/83/EC, as amended
• Checking Process of Mock-Ups and Specimens of outer/immediate labelling and package leaflets of human medicinal products in the Centralised Procedure (EMEA/305821/2006)
• "Guideline on the packaging information of medicinal products for human use authorised by the Community", the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2C
• "Centralised Procedure", the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2A, Chapter 4 and Chapter 7
• The Linguistic Review Process of Product Information in the Centralised Procedure (EMEA/5542/02)
23. Do I have to submit samples together with my application? *Rev. May 06*

Samples for testing the proposed medicinal product are not required at time of submission of the application.

The CHMP may however request the testing of samples of the medicinal product and/or its ingredients during the assessment of the application in accordance with the provisions of Article 7 (b) of Regulation (EC) No 726/2004.

In this case the Rapporteur and/or Co-Rapporteur will specify a test protocol (type of samples, number of samples, number of batches, testing to be performed and methods and specifications to be used) and agree with the EMA which Official Medicines Control Laboratory (OMCL) or other laboratories designated for this purpose by a Member State will carry out the required testing.

Sampling and testing will be co-ordinated by the EMA in collaboration with the European Directorate for the Quality of Medicines and Healthcare (EDQM).

The results of the tests are reported to the EMA, Rapporteur and Co-Rapporteur and the CHMP for consideration in finalising the CHMP Assessment Report.

**References**

- Regulation (EC) No 726/2004
- The Rules governing Medicinal Products in the European Community, Volume 2A, Chapter 4 and Chapter 7

The Marketing Authorisation Holder (MAH) is the person who holds the authorisation to place a medicinal product on the market and is legally responsible for marketing the medicinal product. The granting of a marketing authorisation by a competent authority does not discharge the holder from civil and criminal liability as provided for by the Union law.

The MAH may be a natural or legal person.

The MAH of a centralised marketing authorisation must be established within the EEA (Norway, Iceland, Liechtenstein and the Member States of the European Union).

In order to fulfil this requirement the MAH must have a permanent legal structure which is formed in accordance with the law of an EEA Member State and which allows the concerned holder to assume the duties and responsibilities as well as to perform the tasks laid down by Union law.

Companies or firms formed in accordance with the law of a Member State and having their registered office, central administration or principal place of business within the EEA will be treated in the same way as natural persons who are nationals of Member States. An applicant should demonstrate that it is duly established in the EEA. A proof of establishment from the applicant company is required by the Agency in order for an application to be validated (e.g. in the United Kingdom, a certificate of incorporation issued by the Registrar of Companies, and in France, an extrait du registre du commerce et des sociétés). This proof of establishment should be included in annex 5.3 of the application form.

It should be emphasised that while the MAH may delegate certain activities to third parties, the MAH remains responsible for assuring all the obligations imposed on MAHs by the European legislation and by national law, as applicable.

References:

- Regulation (EC) No 726/2004
- The Rules governing Medicinal Products in the European Community, Volume 2A, Notice to Applicants, Chapter 1
- Directive 2001/83/EC
25. What information relating to the manufacture and batch release should be provided as part of my application? Rev. 

Feb 12

The EMA requires the applicant to provide background information in support of the application relating to the manufacture (including packaging), batch testing and batch certification (batch release) by the Qualified Person in the European Economic Area (EEA). This should be sent to the EMA along with the application dossier.

The EEA includes European Union Member States plus Iceland, Norway and Liechtenstein. Switzerland is not part of the EEA.

Once validated, it is normally not permitted to add a new site or to change the steps of manufacture/batch release described under Module 1.2 (i.e. Application Form) of the application during the 210-day review period. Any additional site or change in the manufacturing or batch release arrangements should be submitted as a variation after the granting of the Marketing Authorisation.

The information on manufacturing/batch release sites submitted in Module 1.2 of the application must be consistent with module 3. All the manufacturing/batch release sites mentioned in module 3 must be listed in Module 1.2 and the activities carried out at each site must be described in Module 1.2 consistently with the information provided in module 3.

25.1. Manufacturing sites

All sites involved in the production of the finished medicinal product and of the active substance must be described (name and detailed address, including building reference) in Module 1.2 of the application for a marketing authorisation together with a description of the steps performed. This should include:

- active substance manufacture and packaging
- any contract laboratories used for testing the active substance (including ongoing stability monitoring)
- bulk medicinal product manufacture
- diluent/solvent manufacture (if any)
- manufacture of any other associated medicinal product (if any)
- finished product manufacture and packaging
- batch release
- any contract manufacturing sites
- any contract laboratories used for testing the finished product
- Official Medicines Control Laboratory (OMCL) for blood products/vaccines if "Official Batch Release" is a requirement for the product in question.

For third country manufacturers, information about any previous EEA inspection in the last 2-3 years and/or any planned EEA inspection(s) should be provided and should include details of the inspection dates, product category inspected and the name of the inspecting competent authority.
25.2. **Documents to be attached to Module 1.2 of the application**

The following documents should be attached to Module 1.2 of the application:

- For all sites in the EEA, other than active substance manufacturers, copies of the "Manufacturing Authorisation" authorising the sites involved in the manufacture, importation, control and /or testing and Qualified Person release of batches of the medicinal product. Alternatively, a reference can be made to the appropriate entry in the EudraGMP database.

(Note: for sites in the EEA, GMP Certificates are not an acceptable alternative to a Manufacturing Authorisation. However, GMP certificates can be useful additional information. Also, particular attention should be paid that the scope of the Manufacturing Authorisation for a given manufacturer covers the activities proposed as part of the Marketing Authorisation Application).

- For all sites other than active substance manufacturers, located in third countries where a Mutual Recognition Agreement or other relevant agreement is in place, MRA certificate, not older that 3 years, from the local competent authority that carried out the inspection and/ or GMP certificate from the EEA inspecting competent authority if the site has been inspected by an EEA competent authority in the last 2/3 years. Where the MRA partner has placed the certificate in the EudraGMP a reference to the entry will suffice. For the countries which have operational Mutual Recognition Agreements (MRA) with the EU, please consult the EMA website on Mutual Recognition Agreements.

- For all sites other than active substance manufacturers, located in third countries with no Mutual Recognition Agreement, GMP certificate from the EEA inspecting competent authority if the site has been inspected by an EEA competent authority in the last 2/3 years. Alternatively, a reference can be made to the appropriate entry in the EudraGMP database.

- In addition to the above, copy of the registration or other document analogous to a manufacturing authorisation from the local competent authority demonstrating that the site is authorised for manufacture of the product/pharmaceutical form and details of any inspection performed other than by EEA authorities (e.g. GMP certificate or similar statement from the competent authority which carried out the inspection).

- A flow-chart describing all the main steps involved in the manufacture of the active substance and finished product.

- For each active substance, a declaration from the Qualified Person(s) of all the finished product manufacturer(s) located in the EEA listed in Module 1.2 where the active substance is used as a starting material and from the Qualified Person(s) of the batch release site(s) in the EEA that the active substance manufacturer(s) listed in Module 1.2 operate in compliance with the detailed guidelines on Good Manufacturing Practice.

25.3. **Contact person in the EEA for product defects/recalls**

A proposed contact point/person in the EEA for Quality problems and defective batches of product must also be provided in Module 1.2 of the application (name, full address, 24 hour emergency phone and fax numbers + e-mail address, and mobile phone number if available).

**References**

- Directive 2003/94/EC
- Directive 2001/83/EC
26. What batch release arrangements in the EEA are required for my medicinal product? Rev. Feb 12

26.1. Importing site/Supervisory Authority

According to Article 51(1) of Directive 2001/83/EC, each batch of a medicinal product must be certified by a Qualified Person prior to release to the market in the EEA.

In the case of products imported from a third country, and for the purpose of Article 51(1)(b) of Directive 2001/83/EC, the site where the certification of batches by the Qualified person occurs is considered to be the importing site in the EEA (and not necessarily the site through which the batch first physically enters the EEA).

The EEA includes European Union Member States plus Iceland, Norway and Liechtenstein. Switzerland is not part of the EEA.

In accordance with the provisions of Article 18 of Regulation (EC) No 726/2004 the Supervisory Authority(ies) shall be the competent authorities of the Member State or Member States which granted the manufacturing authorisation provided for in article 40(1) of Directive 2001/83/EC in respect of the manufacture of the medicinal product. In the case of products imported from third countries, the Supervisory Authority(ies) shall be the competent authority(ies) of the Member State(s) which granted the manufacturing authorisation provided in Article 40(3) of Directive 2001/83/EC to the importer, unless a Mutual Recognition Agreement (MRA) or other relevant agreement covering GMP for the product under consideration is operating with the country where the medicinal product is manufactured.

In the exceptional circumstances where a valid manufacturing authorisation is not in place at the time of the marketing authorisation submission for any finished product manufacturer/importer/batch release site located in the EEA, EMA will consult the Supervisory Authority and a request for inspection may be triggered. The marketing authorisation procedure will require the inspection outcome before opinion and in particular confirmation of the grant of the manufacturing authorisation.

For any finished product manufacturer that is not in possession of a GMP certificate at the time of the marketing authorisation submission located in third countries with no Mutual Recognition Agreement, a request for inspection will normally be triggered. The marketing authorisation procedure will require the inspection outcome before opinion.

26.2. Batch testing upon importation

For medicinal products imported from third countries, retesting of each batch within the EEA upon importation is required unless a Mutual Recognition Agreement (MRA) or other relevant agreement covering GMP for the product under consideration is operating with the country where the medicinal product is manufactured. If such MRA is in operation, batch controls/tests carried-out in the country where the product is manufactured are acceptable.

It should be noted that MRAs cover batch control/testing and do not cover batch release. Batch release must take place in the EEA territory for every production batch released to market in the EEA, regardless of if a MRA with the exporting country is in place or not.

For the countries which have operational Mutual Recognition Agreements (MRA) with the EU, please consult the EMA website on Mutual Recognition Agreements.
Batch release of an imported medicinal product from a third country without re-testing is a serious failure of a qualified person’s legal obligations. According to Article 52 of Directive 2001/83/EC, it is expected that Member States’ Supervisory Authorities will launch appropriate administrative measures and may withdraw the product concerned from the market (Article 117(1)(e) of Directive 2001/83/EC).

26.3. Contracting out of certain controls

The provisions of Article 20(b) of Directive 2001/83/EC allows certain of the controls required under the provisions of Article 51(1) of Directive 2001/83/EC, to be contracted out to third parties, if justified, and provided that the laboratories have been verified by the Competent Authorities. Laboratories used for contract testing upon importation of medicinal products manufactured in third countries may be located in any EEA country.

The Qualified Person of the Manufacturing Authorisation Holder named in the Application is however responsible for certifying that any contract laboratory used carries out the controls in accordance with Good Manufacturing Practice, as applicable and with the requirements of the Marketing Authorisation, once granted.

References

- Directive 2001/83/EC
- Regulation (EC) No 726/2004
- Annex 16 to GMP Certification by a Qualified person and Batch Release (July 2001), Volume 4 of the rules governing medicinal products in the European Union
27. How shall I submit an Active Substance Master File (ASMF)? *Rev. Dec 13*

Annex I to Directive 2001/83/EC describes the concept of an open and closed Active Substance Master File (ASMF) and specifies that:

"For a well-defined active substance, the active substance manufacturer or the applicant may arrange for the:

i) Detailed description of the manufacturing process

ii) Quality control during the manufacture, and

iii) Process validation

to be supplied in a separate document directly to the competent authorities by the manufacturer of the active substance as an Active Substance Master File.

In this case, the manufacturer shall, however, provide the applicant with all of the data, which may be necessary for the latter to take the responsibility for the medicinal product...”

It should be emphasized that the concept of the ASMF shall only apply to a well-defined active substance and cannot be used for excipients, finished products and biological active substances. The information related to excipients, finished products and biological active substances shall be provided within the Marketing Authorisation Application (MAA) by the applicant.

In case an application under the Centralised Procedure includes the submission of an Active Substance Master File (previously referred to as European Drug Master File (EDMF)), applicants should be aware of the fact that, as mentioned in the Guideline on Active Substance Master File Procedure (CPMP/QWP/227/02), an ASMF consists of 2 parts:

- An ASMF Applicant’s Part, also referred to as Open Part, which shall be at the disposal of the applicant.

- An ASMF Restricted Part, also referred to as Closed Part, which is a confidential document closed to the applicant.

Both parts need to be separated and follow the structure of the Module 3.2.S of the CTD.

The content requirements as described in the above mentioned Guideline should be followed.

It is necessary for the applicant and the ASMF holder to liaise to ensure that the ASMF including all necessary documents are synchronized to arrive at ideally the same time as the planned MAA to be submitted by the applicant, although an interval of some days may be allowed. Note that the marketing authorisation application cannot be validated until all the necessary documents are received in a satisfactory form. This also applies to the ASMF-related responses to Day 120 LoQ and Day 180 LoOI. Applicants should be aware that the procedure cannot re-start until the responses from the ASMF holder are received by the Agency.

**Non applicability of ASMF concept to biological active substances**

Further to clarifications from the European Commission on the interpretation of Directive 2001/83/EC as amended, and the subsequent announcement in the October 2004 CHMP Monthly report, the ASMF concept is not acceptable for biological medicinal products.
The characterisation and determination of biological active substances requires not only a combination of physico-chemical and biological testing, but also extensive knowledge of the production process and its control.

The MAH/applicant for a biological medicinal product could therefore not comply with the requirement to 'take responsibility for the medicinal product' without having full and transparent access to these quality-related data. The use of an ASMF would prevent such access, and should therefore not be allowed for biological active substances.

In addition, active substances, which are present in certain medicinal products such as vaccines or cell therapy medicinal products, do not fit with the concept of a 'well-defined' active substance.

Non applicability of ASMF concept of open and closed parts to Vaccine Antigen Master File (VAMF) and Plasma Master File (PMF)

The concept of the ASMF does not apply to blood derived medicinal products and vaccine antigens. In this context, the manufacturer can submit a PMF or a VAMF.

Regarding the VAMF, the legislation specifies that the VAMF holder cannot differ from the MAH/applicant for the concerned medicinal product: there is hence no rationale for an open /closed parts system.

For VAMF linked MAs, if a particular MAH name and address are not identical to the name and address of the proposed VAMF certificate holder, a relevant declaration should be provided attached to the application form, stating that the MA applicant and the MAH belong to the same mother group of companies, which share the same data package.

For the PMF the legislation specifies that where the MAH/applicant differs from the holder of the PMF, the PMF shall be made available to the MAH/applicant for submission to the competent authority.

References

- Annex I to the Directive 2001/83/EC, as amended
- Guideline on Active Substance Master File Procedure (CPMP/QWP/227/02)
- CMD(h) – Overview of Biological Active Substances of non-recombinant origin
- Guideline on requirements for Vaccine Antigen Master File (VAMF) certification (EMEA/CPMP/4548/03/Final/Rev1)
- Guideline on requirements for Plasma Master File (PMF) certification (CPMP/BWP/4663/03)

27.1. What data should be submitted by the ASMF holder? Rev. Dec 13

In the first submission of an ASMF with an allocated EMEA ASMF reference number, the ASMF holder is required to submit:

- ASMF dossier (Applicant’s part, Restricted part, Quality Overall Summary and Expert’s *curriculum vitae*);
- Letter of Access (Annex 2 of the ASMF Guideline);
• Submission Letter and Administrative Details (Annex 3 of the ASMF Guideline);

• A commitment to inform the applicant and the EMA of any change in the ASMF to be provided either as a separate letter or within the Letter of Access (Annex 2 of the ASMF Guideline).

In later marketing authorisation application or variations submissions referencing to an already submitted ASMF with an allocated EMEA/ASMF number, the ASMF holder is only required to submit:

• Letter of Access (Annex 2 of the ASMF Guideline);

• Submission Letter and Administrative Details (Annex 3 of the ASMF Guideline);

• The relevant revised sections of the ASMF dossier reflecting changes to the previously accepted version, as applicable;

• A commitment to inform the applicant and the EMA of any change in the ASMF to be provided either as a separate letter or within the Letter of Access (Annex 2 of the ASMF Guideline).

The Letter of Access and the Letter of Commitment to inform the applicant about any changes in the ASMF should be included both in the application form submitted by the applicant (Annexes 5.10 and 5.11) and also in the ASMF submitted by the ASMF holder.

The contact details of the ASMF holder contact person (including contact email address) must be the same in the Cover Letter of the ASMF, in the Letter of Access and the Application Form (Module 1.2 of the eCTD or Nees submission).

ASMF holders are reminded that any initial submission and update to an ASMF should be accompanied by the Submission Letter and Administrative Details (Annex 3 of the ASMF Guideline) duly filled as detailed in the instructions provided in the Additional guidance on documents relating to an active substance master file. Either an EMEA/ASMF/xxxxx number or EU/ASMF/xxxxx number has to be entered.

New MAA or MAV should always reference the last version of the ASMF submitted. This will be subject to compliance checks during validation of the MAA and MAV.

The ASMF dossier and any subsequent updates should only be submitted once.

27.2. What data should be submitted by the applicant or MAH? Rev. Dec 13

In all cases, the applicant (in the context of a MAA) or MAH (in the context of a MAV) should submit:

• MAA and MAV application form stating the correct EMEA ASMF reference number;

• Copy of the Letter of Access (Annex 2 of the ASMF Guideline), as applicable;

• Copy of the complete current version of the Applicant’s part of the ASMF in Module 3 or its revised sections, as applicable;

• Copy of the commitment from the ASMF holder to inform the applicant and the EMA of any change in the ASMF to be provided either as a separate letter or within the Letter of Access (Annex 2 of the ASMF Guideline).

The Letter of Access and the Letter of Commitment to inform the applicant about any changes in the ASMF should be included both in the application form submitted by the applicant (Annexes 5.10 and 5.11) and also in the ASMF submitted by the ASMF holder.
The latest version of the ASMF submitted in the context of a previous centralised procedure will be considered the current version of that ASMF. The current version of the ASMF should correspond to the version of the ASMF Applicant’s part declared in a new MAA or MAV form and included in Module 3. This will be subject to compliance checks during validation of the MAA and MAV.

**Example:**

The version of EMEA/ASMF/12345 (EMA/ASMF/reference number) currently held at the Agency is: AP January 2012/RP April 2013.

If the version of the ASMF included in the Module 3 of the MAA and referenced in the application form is AP November 2011, the applicant will be requested to update Module 3 and the application form according to the current version of the EMEA/ASMF/12345.

Equally, if the version of the ASMF included in the Module 3 of the MAA and referenced in the application form is AP December 2012, the ASMF holder will be requested to submit the latest version of the EMEA/ASMF together with the Annex 3 of the ASMF Guideline.

ASMF Holders are reminded of their responsibility to inform the MAHs of any changes to their ASMFs. Similarly, MAHs are reminded of their legal obligation to submit the applicable variation to their MAs when changes are proposed to the ASMF, i.e. when an updated version of the ASMF is submitted, the MA(s) linked to that ASMF will only integrate the ASMF update once the applicable variation is submitted and positively concluded.

The applicant should submit the Applicant’s Part of the ASMF. It should be included in their application within Module 3.2.S of the dossier. The ASMF holder should only submit updates to the Applicant’s Part (which should be identical to the one provided by the applicant) and/or Restricted Part, as applicable, in the context of a MAA or MAV. Both parts should be the latest versions available of the ASMF. It is recommended to include a table summarising those changes made to the ASMF compared to the previous version.

Applicants should note that the ASMF constitutes an integral part of the dossier and therefore it should be always made available to the EMA and CHMP Members.

The applicant is responsible for the submission of all necessary documents to the EMA.

It should be noted that although the ASMF procedure is developed to keep intellectual property confidential, it is also permitted to use the procedure when the applicant is also the manufacturer of the active substance.

Additional information on the ASMF procedure can be found in the ASMF WG webpage.

### 27.3. What is the EMEA/ASMF reference number? Rev. Dec 13

From 1 September 2013, ASMF holders submitting their ASMF dossiers relating to a Centrally Authorised Product are asked to send it to the Agency and Committee Members only once.

According to the new ASMF submission rules the Agency will assign a reference number on request prior to submission of the ASMF that can cover multiple CAPs.

The EMEA/ASMF/XXXXX number is an internal reference number sequentially assigned by the EMA to enable an appropriate data lifecycle management of ASMFs used in one or more centralised Marketing Authorisation.
The EMEA ASMF reference number does not replace the responsibility of the ASMF holders to version control their ASMF (in accordance with GMP) nor replaces their own ASMF numbering system.

27.4. Who should request an EMEA ASMF reference number? Rev. Dec 13

The EMEA ASMF reference number should be requested by the ASMF holder for:

- new ASMFs submitted for MAAs and MAVs as of 1 September 2013. From this date, reference to an EMEA ASMF number will be checked at validation,
- ASMFs previously submitted to the EMA when referenced in a new MAA or MAV. The request for the EMEA ASMF reference number should be made before submission of a new MAA or MAV to update the ASMF,
- ASMFs submitted in relation to a variation application.

For previously submitted ASMFs, in cases where the ASMF is used in more than one MA the ASMF Holder should only request one EMEA ASMF reference number, when applicable. The allocated EMEA ASMF reference number should be communicated to the applicant or MAH, so that reference to the EMEA/ASMF/XXXXX number is made in all future submissions.

27.5. When and how to request an EMEA ASMF reference number?

Up to two weeks before submitting a complete ASMF, or an update to an already submitted ASMF, the ASMF holder should request the EMEA ASMF reference number. The request should be sent to PA-BUS@ema.europa.eu.

Agency ASMF reference numbers are allocated sequentially. A request form is available.

The EMEA ASMF reference number allocated by the Agency should be referenced in all subsequent communications (e.g. in response to a validation issue, List of Questions, List of Outstanding Issues, upcoming variation) and should always be included in the following documents:

- MAA (in the field of the National ASMF number) or MAV application form (in the Present and Proposed field);
- Letter of Access (Annex 2 of the ASMF Guideline);
- Submission Letter and Administrative Details (Annex 3 of the ASMF Guideline)

It is the responsibility of the ASMF holder to inform the applicant of a MAA or MAV of the allocated EMEA ASMF reference number. Failure to state a valid EMEA ASMF reference number on the MAA or MAV form will trigger validation questions and may delay the start of procedure.

27.6. EMA ASMF or EU ASMF reference number? Rev. Dec 13

The EU/ASMF reference number allows for the identification by all Competent Authorities (National Competent Authorities and EMA) of ASMFs used in centralised and national (Decentralised and Mutual

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6 Example: substantially different route of synthesis/manufacturing process which results in changes to important quality characteristics of the active substance, e.g. bioavailability of the active substance, may result in the allocation of two different EMEA/ASMF numbers.
Recognition) MAAs or MAVs, and therefore enabling the ASMF Assessment Report Work Sharing (ASMF AR WS) procedure.

More information on the ASMF AR WS will be made available closer to the start of the pilot phase (1 December 2013) in the dedicated ASMF WG webpage.

ASMF holders should either have an EMEA/ASMF/ reference number or an EU/ASMF/ reference number before submitting an ASMF. Both numbering systems will run in parallel as of the start of the pilot phase of the ASMF AR WS. ASMF holders are encouraged to request an EU/ASMF/reference number if the ASMF is expected to be used in centralised and national applications.

27.7. **Which format and submission channel should be used for submitting ASMFs?** Rev. Aug 14

Under the new ASMF submission requirements, the following formats are accepted for ASMF submissions:

- Electronic Common Technical Document (eCTD)
- Non-eCTD electronic submission (NeeS)

Guidance on the above formats can be found on the eSubmission website. Additional, please take a note of the EMAs statement of intent of the mandatory use of eSubmission Gateway and Webclient.

**Submission requirements for the different Committee (Co-) Rapporteurs**

ASMF holders must submit the application to all (Co-)Rapporteurs, otherwise there may be a delay in the start of the procedure due to the time lapse between the validation by the Agency and the confirmation from the (Co-)Rapporteurs that they have received the dossier.

For a full overview of the submission requirements for the different Committee (Co-)Rapporteurs see: Dossier requirements for Centrally Authorised Products (CAPs).

The above method and requirements also apply to the submission of responses to List of Questions / List of Outstanding Issues.

27.8. **How to proceed if the ASMF was previously submitted in paper format?** Rev. Dec 13

The ASMF holder of ASMFs previously submitted in paper format should request an EMEA ASMF reference number as indicated above.

After the reference number is allocated the ASMF holder should submit the ASMF in an accepted electronic format (Electronic Common Technical Document (eCTD) or Non-eCTD electronic submission (NeeS)).
27.9. How to proceed if there is an existing eCTD life-cycle for the ASMF?

NEW Dec 13

ASMF holders need to request the EMEA/ASMF number by filling the request form. The EMA will provide the requestor with the new number within 3 working days. Please note that this number is NOT equivalent of EU/ASMF number and should never be inter-changed.

If the ASMF holder already has more than one eCTD life-cycle filed for the given substance, they will need to select one of these (informing the EMA in the cover letter which one it will be) and follow the eCTD life-cycle of the selected ‘product’ only. This, selected life-cycle will, then receive a new EMEA/ASMF/01xxx number covering all listed CAPs.

Once the ASMF holder is submitting an update or new version to the ASMF, they have to do so with this new number. The ASMF holder will have to prepare a new sequence (increasing by one) in which (module 1, cover letter) they declare that the previously submitted ASMF version has not been modified since it was last submitted.

If there have been modifications (new version) since the last ASMF submission, additionally the relevant modules within this new eCTD sequence will have to be updated.

ASMF holders have to inform all MAH(s) about the new EMEA/ASMF/xxxxx number and if an update is submitted to an ASMF related to their Centrally Authorised Product (the MAH should then submit the relevant variation application)
28. What shall I submit if my medicinal product contains or consists of genetically modified organisms (GMOs)?

Potential applicants are advised to discuss their future applications which consist or contains of GMOs well in advance (6 months – 1 year) of their submission with the EMA.

Applicants may also find it useful to apply for scientific advice during the development of their medicinal product. For any scientific advice questions relating to the Environmental Risk Assessment (ERA), the necessary consultations will be held with the designated GMO Competent Authorities (CAs).

With the letter of intent to submit an application for a Marketing Authorisation under the Centralised Procedure for a medicinal product containing or consisting of Genetically Modified Organisms (GMOs) within the meaning of Article 2 of Directive 2001/18/EC, the applicant will be required to provide a confirmation that all obligations have been complied with. It is necessary to ensure the traceability at all stages of the placing on the markets of GMOs as or in products authorised under part C (article 12) of the above-mentioned Directive.

Article 6 of Regulation (EEC) No 2309/93 specifies the documents to be presented in Module 1.6.2 for a Marketing Authorisation Application (MAA) for a medicinal product consisting of or containing GMO(s):

- A copy of the CA’s written consent to the deliberate release into the environment of the GMOs for research and development purposes. Although already appearing in Modules 1 (annex to the application form), this information should be repeated in Module 1.6.2.

- The technical and scientific information on the GMO specified in Annexes III and IV to Directive 2001/18/EC. As the Directive qualifies this point with a statement to the effect that not all listed points may be applicable to particular GMOs or GMO categories, the list in these Annexes should be understood to be a compilation of points to consider which is subject to justified deletions and/or additions, depending on the nature of the medicinal product. The information also needs to take into account, inter alia, the diversity of sites of use of the GMO and the results of research and trials already completed on the GMO.

- The ERA dossier. The content of this dossier should follow the order of headings and requirements specified within Annex II to Directive 2001/18/EC and expanded upon in Commission Decision 2002/623/EC.

- The results of any investigations performed for the purposes of research or development.

In addition and in analogy with the requirements of Article 6 of Regulation (EEC) 2309/93, it is recommended to complete M1.6.2 with the following:

- Information on the proposed product information (including proposed conditions of use and handling) and on the packaging of the product. Although already appearing elsewhere in the MAA, this information should be repeated in Module 1.6.2 for the benefit of the lead consulted CA which will not receive the full MAA dossier.

- A plan for monitoring, in accordance with Council Decision 2002/811/EC, during the period of use and beyond, of the product, or a justification for the omission of such a plan.


- Bibliographical references.
The module section 1.6.2, presenting all these particulars, should be bound separately from the remainder of the dossier. Moreover, there is no provision for a summary to be included in Module 2 of the dossier.

The fundamental dossier requirements for ERAs for GMOs proposed to be placed on the market as or in products are included in Directive 2001/18/EC and in Commission Decision 2002/623/EC.

Technical and scientific information presented in the ERA will overlap with items of information presented in other sections of Module 1, and other Modules of the MA application dossier. Applicants are reminded to ensure full consistency of all data throughout the dossier, bearing in mind that variability, reflecting different origins (medicinal product regulatory versus environmental regulatory texts) may occasionally be encountered in the official terminology describing GMO attributes.

This Directive 2001/18/EC shall not apply to GMOs as or in products as far as they are authorised by the Council Regulation (EEC) No 2309/93 provided a specific environmental risk assessment is carried out in accordance with the principles set out in Annex II to this Directive and on the basis of the type of information specified in Annex III to this Directive.

Council Regulation (EEC) 2309/93, as amended, requires that the Rapporteur hold necessary consultations with the Competent National Authorities under Directive 2001/18/EC, where the medicinal product contains or consists of GMOs.

To accelerate the consultation process, the CHMP rapporteur may appoint one of the national GMO CAs to act as lead consulted CA. This lead consulted CA will liaise with its fellow GMO CAs on the review of the documentation forwarded to it by the applicant.

The assessment report on the module 1.6.2 data, prepared by the lead consulted CA and including any comments received from the fellow CAs, will be sent to the Rapporteur for CHMP consideration. The CHMP Members will subsequently have the opportunity to comment on all aspects of the scientific assessment. The environmental assessment is an integral part of the assessment report, and is done accordingly to the same timelines.

It should be highlighted that current experience under the Centralised Procedure with medicinal products containing or consisting of GMOs is rather limited. Once more experience is accumulated, an SOP will be prepared by the EMA for discussion and adoption by the CHMP.

References

- The Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2B, Presentation and content of the dossier
- Environmental Risk Assessments for Medicinal products containing, or consisting of, Genetically Modified Organisms (GMOs) (Module 1.6.2) (EMEA/CHMP/BWP/135148/2004) (CHMP released for consultation on EMA website, January 2005)
29. What information shall I provide if my medicinal product contains or uses in the manufacturing process materials of animal and/or human origin? Rev. July 06

The applicant must comply with the Part I Module 3.2 (9) “Content: basis and principle” of the Annex I to Directive 2001/83/EC, as amended, which requires that “The applicant must demonstrate that the medicinal product is manufactured in accordance with the Note for Guidance on minimising the risk of transmitting animal spongiform encephalopathy agents via medicinal products (…)” and its updates.

Demonstration of compliance with “the Note for guidance on minimising the risk of transmitting animal spongiform encephalopathy agents via medicinal products” can be done by submitting Certificates of Suitability from the European Directorate for the Quality of Medicines (EDQM) (in Annex 6.12 of the Application form), or by inclusion in module 3.2 of the dossier of scientific data to substantiate this compliance. In the latter situation, this data should be reviewed in Module 2.3 (expert reports).

For all applications, the table A on ’Materials of animal origin covered by the Notice for Guidance on minimising the risk of transmitting animal spongiform encephalopathy agents via medicinal products’ should be completed and included in Module 3.2.R.

For materials from animals not covered by the Notice for Guidance on minimising the risk of transmitting animal spongiform encephalopathy agents via medicinal products and the Annex I to Directive 2001/83/EC as amended, applicants are requested to complete the table B on ’Other materials of animal origin’, and include it in Module 3.2.R.

Materials of human origin

If an application relates to a medicinal product, which contains or uses in the manufacture materials of human origin, applicants are requested to complete the table C ’on albumin and other human tissue derived materials’ and include it in Module 3.2.R.

References

- Directive 2001/83/EC
- Note for guidance on minimising the risk of transmitting animal spongiform encephalopathy agents via medicinal products (EMEA/410/01)
30. Where on my medicinal product information can I mention a local representative? **Rev. Jan 06**

Some Holders of Community Marketing Authorisations have requested that there be a contact point identified in the Package Leaflet and on the label. This would normally be the Holder of the Community Marketing Authorisation. However, a Marketing Authorisation Holder may wish to add the name of another (local) contact point, the "local representative".

"Local representative" shall be taken to mean: any private or legal person established in the Community charged, through a civil contract with the Marketing Authorisation Holder, with representing him in a defined (geographical) area; this contract excluding any transfer of any responsibility imposed on the Marketing Authorisation Holder by Community law and by national law, regulation and administrative action implementing such Community law.

The "local representative" may be indicated:

- In the Package Leaflet, under heading 6 as detailed in the QRD Product information Template, by name, telephone number and electronic e-mail address (optional) only. Postal address may be added space permitting, and
- By name in the blue box on the label, as long as not interfering with the legibility of the EU text on the outer packaging, and if mentioned in the leaflet.

All telephone numbers should be accessible when dialled from abroad (e.g. when a toll free number is given which is not accessible from abroad, an alternative international number may have to be added).

Reference to website addresses or to e-mails linking to websites are not allowed neither for the marketing authorisation holder nor for the local representative.

Designation of a local representative cannot be a requirement but, when the Holder of a Community Marketing Authorisation wishes to identify a local representative in the Leaflet, all of the Community must be covered so that the consumer in each Member State and EEA country has equivalent access to a local representative. A local representative may be designated for more than one Member States or EEA country and may be also the Marketing Authorisation Holder when no other local representative is indicated.

Moreover it is reminded that, in principle, only one local representative should be indicated per Member State or EEA country. Local representatives should be able to address queries in the local official EEA language(s) of the country for which he or she is designated.

There has been some confusion with regard to terms such as 'exploitant', 'technical director', 'distributor' etc. Since there is neither a commonly agreed understanding of these terms nor equivalent legal definitions of these terms amongst the Member States, and in the absence of any reference or definition in Community law, reference to such terminology will not be accepted for a medicinal product authorised by the Community.

It must be recalled that Member States may not require that a local representative of the Marketing Authorisation Holder be appointed for their territory. Therefore, the arrangements outlined above are **purely optional** for Holders of the Community Marketing Authorisations.
References

- “Guideline on the packaging information of medicinal products for human use authorised by the Community” the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2C
- QRD Templates with Explanatory Notes

In order to fulfil EU dossier requirements applicants must submit new Marketing Authorisation Applications (MAA) as follows:

Languages to be used

All applications have to be submitted in English.

Format of submission

From 1 January 2010, eCTD is the only acceptable electronic format for all applications and all submission types in the context of the centralised procedure (e.g. new applications, variations, renewals). Any other electronic format, including NeeS, will be automatically rejected and the submission receipt will not be acknowledged. Additionally, if the eCTD submission results in an invalid Technical Validation the submission will not be accepted.

The latest version of the ICH M2 eCTD specification can be found at http://www.ich.org/products/electronic-standards.html, and the current version of the eCTD EU Module 1 specification can be found in the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2B or the eSubmission website with related documents.

Where applications are amended during the agency’s review, such as e.g. responses to the lists of questions or a withdrawal, a new or consolidated eCTD sequence should be provided in order to maintain the eCTD life-cycle. Replacement sequences of a previously submitted eCTD application (e.g. following corrections) are not acceptable. Any modification of an eCTD application must be reflected in a new eCTD sequence.

For further information regarding the e-submission requirements in the context of the Centralised Procedure, please refer to the TIGes Harmonised Guidance for eCTD submissions.

The use of the electronic Application Forms (eAFs) is strongly recommended for Centralised Procedure. Information on the electronic Application Form electronic application form can be found on the eSubmission eAF webpage.

Cover letter

The European Medicines Agency is standardising the administrative information required in cover letters for any submission concerning centralised procedures. This is in line with changes to the internal financial system and quality improvements to distribution workflows. The Summary Table should be incorporated in the cover letter of each submission in the Centralised Procedure (see explanatory notes in the template).

Please refrain from sending additional and separate copies of cover letters as they will create delays in processing.

Product Information (PI)

As per eCTD requirement, the Product Information (SmPC, PIL and labelling) has to be submitted within the module 1 of the eCTD structure in PDF format. Additionally, this information should also be submitted in Word format outside the eCTD structure but in the same eSubmission Gateway /
eSubmission Web Client package within a folder called “xxxx_working documents”, where the number (xxxx) equals the sequence number.

**Active Substance Master File (ASMF)**

In cases where an Active Substance Master file (ASMF) exists, the applicant should ensure that the Active Substance Master File is or has been submitted by the ASMF holder to the Agency, (see also question “How should I submit an active-substance master file (ASMF)?”), in order to proceed with the validation of the dossier.

**Submission to the EMA**

From 1 March 2014 the use of the eSubmission Gateway or Web client is mandatory for all electronic Common Technical Document (eCTD) submissions through the centralised procedure. The European Medicines Agency (EMA) no longer accepts submissions on CD or DVD. This applies to all applications for human medicines.

More information on how to register and connect to the Gateway / Web Client can be found in the eSubmission website and detailed information on the required naming conventions and file formats can be found in the European Medicines Agency eSubmission Gateway: Questions and answers relating to practical and technical aspects of the implementation and the eSubmission Gateway web client: Guidance for applicants. Applicants must not send duplicate submissions electronically or via CD-ROM or DVD as this might lead to delays in the handling of applications.

An automated ‘acknowledgement’ e-mail is sent from the system confirming whether their submission has passed the relevant technical validation criteria and has been uploaded to the agency’s review tool and made available via the Common Repository. There is no need to send any accompanying hard media or separate paper cover letters for these submissions, as the cover letter will be in the relevant part of eCTD module 1 in PDF format.

**Submission requirements for the different Committee (Co-) Rapporteurs**

One electronic copy should be submitted to the (Co-) Rapporteurs after the eSubmission Gateway/Web Client confirmation of a technically valid submission to the EMA if the relevant NCA is not using the Common Repository. Otherwise there may be a delay in the start of the procedure due to the time lapse between the validation by the Agency and the confirmation from the (Co-)Rapporteurs that they have received the dossier. For a full overview of the submission requirements for the different Committee (Co-)Rapporteurs see: Dossier requirements for Centrally Authorised Products (CAPs).

The above method and requirements also apply to the submission of responses to List of Questions / List of Outstanding Issues.

**Validation of the application**

In the event that the Agency requires additional data, information or clarification in order to complete its validation of the dossier, it will contact the applicant requesting to supply this information within a specific time limit. When supplying the Agency with this information, the applicant should also send a copy of this information to the (Co-)Rapporteurs, if necessary in accordance with the published document “Dossier requirements for Centrally Authorised Products (CAPs)”. In this case, the validation can only be completed after receipt and verification of the information submitted. The submission of responses to validation supplementary information (VSI) should be sent in accordance with eCTD requirements including validation supplementary information (VSI) related to the ASMF part of the dossier, when applicable.
In order to start the procedure by the targeted start date, the applicant is required to provide the information requested within a given deadline. If the applicant is unable to respond within the deadline, the Agency is able to accept the responses up to 2 months from the VSI letter. The published submission timetable applies. If no response is received within 2 months the validation outcome will be considered negative and the application closed. An invoice for the relevant administrative fee will follow.

If the (Co-)Rapporteurs have not received their copy of the dossier and/or supplementary validation information on the day the dossier is validated by the Agency, the start of the procedure may be delayed until the procedural starting date of the next month.

**Submission requirements for the other Committee members**

After validation of the application, the Agency will notify the applicant accordingly in writing. The same notification will also be sent to the (Co-)Rapporteurs.

Upon receipt of this notification, the applicant should forthwith send one electronic copy of the application to the other Committee Members who are not using the Common Repository, including any additional data or information supplied during the validation phase as appropriate.

There are number of NCAs that have access to centrally authorised product submissions directly via the Common Repository after submission to the Agency. Please note that these NCAs do not need additional submissions. Please refer to the "Dossier requirements for Centrally Authorised Products (CAPs)" document to see if an electronic copy should also be sent to other Committee members after the validation phase for evaluation, to maintain the life cycle of the eCTD dossier.

It is essential that identical eCTD sequences are circulated to Committee Members. Any minor changes that affect the ‘md5 checksum’ will lead to inconsistency and possibly result in future technical invalidity.

**References**

- “Centralised Procedure”, the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2A, Chapter 4
- The Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2B, Electronic Technical Document (eCTD)
- Official Website for ICH
- eSubmission Website
- eSubmission Gateway and the Web Client
- Electronic Application Form
- Dossier requirements for centrally authorised products
Initial marketing authorisation (MA) applications submitted to the European Medicines Agency (EMA) as part of the centralised procedure are subject to a validation process. The objective is to make sure all essential regulatory elements required for scientific assessment are included in the MA application prior to the start of the procedure. Initial MA validation has been centralised and is now being performed by a dedicated service within the Agency.

There are two elements to validation:

1. The first is technical validation which takes place once an electronic application has been received by the Agency. This ensures that the structure of the submission is compliant with the EU Module 1 Specification.

1. The second element is regulatory and administrative content validation, which can only commence once the application has successfully passed technical validation.

**What to expect once an Initial Marketing Authorisation Application has been submitted to the EMA?**

Once submitted to the European Medicines Agency in the agreed standard format, the Agency performs a technical validation. The outcome of this technical validation is immediately notified to the applicant when the application is received via eSubmission / Web Client.

If the dossier is technically invalid and the replacement sequence is not delivered by the intended submission deadline, the start of the procedure is automatically postponed to the next month, as only technically valid and complete applications can be subject to the validation process. This also applies to the Active Substance Master File (ASMF) submissions (see "How to avoid most common quality validation issues – Active Substance Master File (ASMF)").

The Agency will inform the applicant of the start of the regulatory and administrative content validation. If any issues are found during validation then the Agency will issue a Validation Supplementary Information (VSI) request to the applicant. Applicants will have to respond to this request in order to resolve any validation issues before the procedure can start. Any response to this VSI request has to be sent as a new sequence.

The Agency will communicate to the applicant the outcome of the validation. A positive outcome means that the scientific evaluation will start on the next available starting date according to the Agency timetables and the applicant will be invoiced the relevant fee. A negative outcome means that the applicant will have to re-submit a new application and will be invoiced a negative validation administrative fee.

**What are the potential scenarios when validating an Initial Marketing Authorisation Application?**

There are four potential scenarios:

**Scenario one (valid first time, no supplementary information requested)**

1. The applicant submits a complete application according to the Agency’s guidance (see below: *What are the main principles that my application should follow in order to pass validation successfully?*)

2. The Agency does not require any additional information.
3. The Agency will confirm the positive validation to the applicant via a positive validation letter.

4. The scientific evaluation will start on the next available start date according to the EMA timetables.

**Scenario two (validation supplementary information requested)**

5. The applicant submits an application that is not in accordance with the Agency’s guidance (see below: *What are the main principles that my application should follow in order to pass validation successfully?*).

6. The Agency will ask the applicant via a request for Validation Supplementary Information (VSI) to submit the additional information, clarifications or corrections.

7. The applicant provides the above additional information within the validation timeline. If the additional information submitted is as requested, the Agency will confirm the positive validation to the applicant.

8. The scientific evaluation will start on the next available start date according to the Agency timetables.

**Scenario three (suspension of validation)**

9. The applicant submits an application that is not in accordance with the Agency’s guidance. (see below: *What are the main principles that my application should follow in order to pass validation successfully?*).

10. The Agency will ask the applicant via a request for Validation Supplementary Information (VSI) to submit the additional information, clarifications or corrections.

11. However, if the additional information is not provided as requested and within the validation timeline, the validation will be suspended and the applicant informed accordingly. The applicant will have up to two months from the date of the initial Validation Supplementary Information (VSI) request to provide the additional information, clarifications or corrections.

12. Within the two month period and according to the EMA timetables, the Agency will confirm the positive validation if all pending issues have been addressed otherwise a negative validation will be generated.

**Scenario four (negative validation)**

13. In the case of non-compliance with applicable legal and regulatory requirements within the above mentioned 2 months, the Agency will issue a negative validation.

14. In that case, the Agency will confirm the negative validation to the applicant via a negative validation letter and invoice the administrative fee.

**What are the timelines of initial Marketing authorisation validation?**

Validation takes place according to the Agency procedural timetable. Applications received on or before a quoted submission date will undergo validation by the Agency. The application must receive a positive validation outcome in order for a procedure to start on the next available start date.

**What are the main principles that my application should follow in order to pass validation successfully?**

The agreed standard format should be used. An eCTD structure according to the TIGes Harmonised Guidance for eCTD Submission should be sent and the format should strictly follow Volume 2B of the Notice to Applicants.
The use of the Electronic Application Form is also highly recommended.

The application form and the different parts / modules of the dossier should be consistent (i.e. the composition is the same in the application form and in module 3 and SmPC).

**How to avoid common validation issues?**

The following sub-paragraphs provide guidance on how to avoid common issues found during validation that will help to submit an application valid first time.

**How to avoid most common Good Manufacturing Practice (GMP) validation issues**

Manufacturing authorisations or equivalent for MRA partners (GMP certificates)

- Please make sure that the scope covers the activities that a given applicant is seeking registration for in the marketing authorisation application.

QP declaration(s)

- Please make sure that the Qualified Person (QP) declaration (annex 5.22) is in accordance with NTA requirements. For human medicinal products, please make sure the date of the last audit of the active substance manufacturer(s) is included.

Consistency / missing information

1. Please make sure that the following sections are consistent:
   1.1. Sections 2.5.1, 2.5.2 and 2.5.1.2 of the application form versus section 3.2.P.3.1 of module 3;
   1.2. Section 2.5.3 of the application form versus section 3.2.S.2.1 of module 3;
   1.3. Flow-chart indicating all manufacturing and control sites involved in the manufacturing process of the medicinal product and the active substance (annex 5.8) versus the application form.

2. Please make sure that the following documents are not missing:
   2.1. Documents equivalent to manufacturing authorisation (Article 8.3(k) Dir. 2001/83) for non-EU/EEA manufacturers. While not explicitly a validation issue, applicants should bear in mind that active substance manufacturers located in third countries should be registered with their local authorities in order to facilitate the import of active substances, in accordance with the Falsified Medicines Directive (Directive 2011/62/EU).

**How to avoid most common Good Clinical Practice (GCP) validation issues**

1. The pivotal studies to support the application in module 2.5 must be identified as such.

2. Please make sure that a list of inspection(s) conducted or planned by other regulatory authorities, related to the product and trial sites involved, is provided, preferably attached to the application cover letter. Alternatively provide confirmation that no inspections have taken place nor are planned.

3. Ensure the list of investigators (name, address, country), preferably in a tabular form, showing the number of patients enrolled by each site, and the total number of sites is included.

4. Please make sure that a table with the number of patients enrolled per country is included. These should be identified in the clinical study report of each study, for instance in section 10.1 or appendix 16.1.4.
5. Please make sure that a clear description of the study administrative structure (clear identification of the sponsor and of the parties who have performed the monitoring, data management, statistics, laboratory assessments, randomization, site(s) of manufacture, other applicable activities and the location of the trial master file) preferably in a tabular form and indicating name and address of the site where each activity was performed, responsibilities and scope of each activity is included. These should be identified in the clinical study report of each study, for instance in section 6, or appendix 16.1.4.

**How to avoid most common Paediatrics validation issues**

1. When applicable, please make sure that the full Paediatric Investigation Plan (PIP) decision (with all annexes) is submitted and not the Opinion. Please make sure that the compliance letter is not missing (sometimes applicant submits only the Compliance report).

2. In the application form, please make sure that the decision number (P/0xxx/YY) is submitted and not the PIP procedure number (EMEA-000xxx-PIPxx-YY). In case of class waiver, please make sure that not only the applicability letter is submitted, but also the last class waiver decision (currently CW/1/2011).

**How to avoid most common Quality validation issues**

- Where permitted, valid justifications for the absence of documents in Module 3 must be provided in module 2.3, e.g. for novel excipients or material of human and animal origin.

- Latest versions of Certificates of Suitability (CEP) for the active substance (chemical) or TSE must be provided.

- Please make sure that the following sections are consistent with one another:
  
  2.2. Section 2.6.1, of the application form versus section 3.2.P.1 of module 3 versus Product Information (composition)

  2.3. Section 2.6.2 of the application form versus module 3.2.A.2 versus module 3.2.R (materials of human or animal origin)

  2.4. Section 2.2.3.2 of the application form versus module 3.2.P.8.1 versus Product Information (shelf-life)

- Please make sure that the following information is included:

  2.5. Composition of coating mixtures, printing inks and gelatine capsules (both in application form section 2.6.1 and module 3.2.P.1)

  2.6. Description of analytical methods/validation – information for all methods should be present in 3.2.S.4 and/or 3.2.P.5

**Active Substance Master Files (ASMF):**

- Active Substance Master Files are often submitted to the EMA in a non-compliant and incomplete format. Applicants should provide the ASMF Holder with the requested information in advance of their submission and ensure that the EMA ASMF Submission Rules are always followed. Non-compliant or incomplete submissions will result in failed technical validation.

- An ASMF is an integral part of the MAA dossier. If the ASMF procedure is used by the applicant as indicated in section 2.5.3 of the application form, the ASMF should be delivered to the Agency well before the submission deadline, in coordination with the related dossier. If the ASMF is technically invalid or arrives to the Agency after the submission deadline for the given month, the dossier is
considered incomplete. In this case, the applicant will be informed of this fact and the validation process would be suspended until a complete dossier is presented. The applicant should liaise with the ASMF holder in order to provide a valid ASMF as soon as possible.

- The validation will be resumed upon receipt of the valid ASMF. Once the validation is finalised, the options are:
  - No validation issues. The procedure starts at the next available date
  - Validation issues identified. A VSI letter will be issued and deadline for responses identified. The following steps are as above (scenario 2, 3 or 4)

- Check there are no missing expert statements in the Active Substance Master File (ASMF)
- The ASMF holders are encouraged to use the eSubmission tool, i.e. Gateway/Web Client portal. This prevents delays related to courier services, holidays and technical validation.

**Other frequently encountered issues:**

- Application for the new active substance status should be presented as annex 5.23 to the Application Form (required as of 3 June 2013).
- Name of the active substance in the application form:
  - 2.7. Title page, section 2.2.1, section 2.6.1 – active substance including salt/hydrate
  - 2.8. 2.1.2 – INN
- Strength of the dosage form in the dossier and Product Information should be expressed per active moiety (i.e. base), corresponding to the actual salt – e.g. exemplain 120 mg (corresponding to 135 mg exemplain hydrochloride); see also here.
- Section 2.6.2 of the application form and modules 3.2.A.2. and 3.2.R should contain ALL materials of human and animal origin, including those used during manufacture of the active substance (e.g. components of fermentation media), while module 3.2.P.4.5 relates only to excipients.
- Where applicable, justification for not using a standard term for pharmaceutical form should be placed in module 1.3.
- Excipients should be tested according to Ph. Eur. monograph where available. Where no Ph. Eur. monograph exists, other pharmacopoeias may be used as supportive but description and validation of the methods used should be provided. For detailed requirements see also here.
- Ph. Eur. analytical methods should be used whenever possible (e.g. for heavy metals, loss on drying)

**How to avoid most common Non-clinical and Clinical validation issues**

- When documents in Modules 4 or 5 are absent, a valid justification must be provided:
  - The non-clinical overview and summaries should be structured following the indents of module 4.
  - The clinical overview and summaries do not mandate explicitly addressing some indents of module 5 (most commonly: 5.3.1.2. Comparative Bioavailability (BA) and Bioequivalence (BE), 5.3.1.3 *In vitro - In vivo* Correlation, 5.3.5.1 Study Reports of Controlled Clinical Studies and 5.3.5.2 Study Reports of Uncontrolled Clinical Studies). Justifications for the absence of studies in any indent of module 5, even when it is considered self-explanatory, should be included in the clinical overview.
• Please make sure that a valid justification for empty eCTD modules/indents is provided, i.e. a statement that relevant studies have not been conducted or are not available is not considered a valid justification. Even if it is straightforward or a matter of development strategy (e.g. reliance of outstanding results from uncontrolled studies to support marketing authorisation application in the absence of controlled data), the reason for ultimately not submitting certain studies should be explained.

• Please make sure that justifications are easy to locate/identify: Validation does not entail extensive review of submitted materials, but only identification of legally required elements. Providing a necessary justification for empty eCTD modules/indents within a discussion may prevent the validation officer from identifying it. One may want to place such justifications under relevant subsections of the (Non-) Clinical Overview or provide them as an Annex at the end of the (Non-) Clinical Overview making sure of the use of the correct terminology of the indent.

**How to avoid most common pharmacovigilance validation issues**

1. Please make sure that a statement indicating that “the applicant has the necessary means to fulfil the tasks and responsibilities listed in Title IX of Directive 2001/83/EC” is included.

2. Please make sure there is a reference to the location where the pharmacovigilance system master file for the medicinal product is kept.
33. When to submit the Marketing Authorisation Application?

Rev. Aug 14

In the same way as it is important for applicants to plan their application strategies for an efficient use of their resources, it is important for the European Medicines Agency, Committee members and Experts to be able to plan and allocate their workload efficiently. If the actual submission date is several months after the date originally indicated, (Co-)Rapporteurs may find it difficult to provide the necessary expertise and re-appointment could be necessary.

The European Medicines Agency advises applicants to consider the date of submission very carefully and to notify the Agency and (Co-)Rapporteurs of a 'real' submission date.

At least **seven months** before submission, applicants should notify the European Medicines Agency of their intention to submit a MAA and provide the intended date of submission. This should be done by using the Pre-submission request form Pre-submission request form (Intent to submit MA); selecting as a scope of request: *Centralised Procedure-Intent to submit a MAA*; this should be sent electronically to pa-bus@ema.europa.eu. The appointment procedure for (Co-)Rapporteurs will be initiated 7 months prior to the Marketing Authorisation Application intended submission date (see question "What is the procedure for appointment of CHMP Rapporteur/Co-Rapporteur and their assessment teams?").

Furthermore applicants are requested to notify the European Medicines Agency and (Co-)Rapporteurs as soon as possible when the previously notified submission date cannot be met, by re-sending an updated Pre-submission request form, selecting as a scope of request: *Notification of change-applicant/contact person details*.

Applicants are finally requested, if they no longer wish to pursue the submission of their application, to notify the European Medicines Agency of their intention to withdraw the request for submission of a MAA. This should be done by using the Pre-submission request form, selecting as a scope of request: *Withdrawal of request*; this should be sent electronically to pa-bus@ema.europa.eu. Please note that this will close the case procedure and the whole pre-submission history.

The submission deadlines and full procedural detailed timetables are published as a generic calendar (see submission deadlines and full procedural timetables). The published timetables identify the submission, start and finish dates of the procedures as well as other interim dates/milestones that occur during the procedure. Applicants should ensure that a technically valid eCTD submission is received by the EMA before the submission deadline. Any technically invalid sequence will result in non-acceptance that may cause a delay in the start of the procedure.

In order to accelerate and facilitate the procedure, one electronic copy should be submitted to the (Co-) Rapporteurs after the eSubmission Gateway/Web Client confirmation of a technically valid submission to the EMA if the relevant NCA is not using the Common Repository (refer to the published "Dossier requirements for Centrally Authorised Products (CAPs)"). Please note that the EMA requires only one eSubmission Gateway/ Web Client submission without any paper cover letter.

After the notification of a valid application, the Agency will send an invoice to the Applicant. The fees should be paid within 30 days of the receipt of this invoice. For more information regarding the applicable fee, see question "What fee do I have to pay and how is the appropriate fee for my application calculated?".

For more information on the complete set of documents that need to be submitted and for the addresses of Committee members for submission of the application, see question "How and to whom shall I submit my dossier?"
References

- The Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2A, Chapter 4 and Chapter 7
- Dossier requirements for Centrally Authorised Products (CAPs)
34. How shall my application be evaluated (timetable)? Rev. Aug 14

Once the application is validated and provided the Rapporteurs have confirmed that they have received the dossier, the EMA starts the procedure at the monthly starting date published on the EMA website. If, within a month from the start of the procedure, any other Committee member has not received the requested parts of the dossier from the applicant, the EMA will stop the clock until confirmation is received that each member has been delivered the requested documentation. It is therefore important that applicants are able to provide a proof of delivery to Rapporteur, Co-Rapporteur and to Committee members (upon request) to the EMA.

A timetable is prepared by the EMA in consultation with Rapporteur and Co-Rapporteur. This timetable is then proposed to the CHMP for adoption.

In order to allow the CHMP to adopt a timetable at the first CHMP meeting after submission of the valid MAA, applicants are advised to submit the MAA accordingly to the published EMA calendar (See "Dates for CHMP meetings").

The submission deadlines and full procedural detailed timetables are now published as a generic calendar on the EMA website (see: "submission deadlines and full procedural timetables").

The published timetables identify the submission, start and finish dates of the procedures as well as other interim dates/milestones that occur during the procedure.

The EMA shall ensure that the opinion of the CHMP is given within 210 days (less any clock-stops for the applicant to provide answers to question from the CHMP) in accordance with the following standard timetable, which can be shortened in exceptional cases (see Request for accelerated assessment).

<table>
<thead>
<tr>
<th>DAY</th>
<th>ACTION</th>
</tr>
</thead>
<tbody>
<tr>
<td>1*</td>
<td>Start of the procedure</td>
</tr>
<tr>
<td>80</td>
<td>Receipt of the Assessment Report(s) or critique from Rapporteur and Co-Rapporteur(s) by CHMP members (which includes the peer reviewers) and EMA. EMA sends Rapporteur and Co-Rapporteur Assessment Report/critique to the applicant making it clear that it only sets out their preliminary conclusions and that it is sent for information only and does not yet represent the position of the CHMP.</td>
</tr>
<tr>
<td>87</td>
<td>PRAC Rapporteur circulates the RMP assessment report and proposed RMP LoQ</td>
</tr>
<tr>
<td>100</td>
<td>(Co-)Rapporteurs, other Committee members and EMA receive comments (including peer reviewers).</td>
</tr>
<tr>
<td>101-104</td>
<td>PRAC adopts PRAC RMP Assessment Overview and Advice for D120 LoQ</td>
</tr>
<tr>
<td>115</td>
<td>Receipt of draft list of questions (including the CHMP recommendation and scientific discussions), from CHMP (Co-)Rapporteurs, as discussed with the peer reviewers, together with the PRAC RMP Assessment Overview and Advice by CHMP members and EMA</td>
</tr>
<tr>
<td>120</td>
<td>CHMP adopts the LoQ as well as the overall conclusions and review of the scientific data to be sent to the Applicant by the EMA. Clock stop. At the latest by Day 120, adoption by CHMP of request for GMP/GLP/GCP inspection, if necessary (Inspection procedure starts).</td>
</tr>
<tr>
<td>121*</td>
<td>Submission of the responses, including revised SmPC, labelling and package leaflet texts in English.</td>
</tr>
</tbody>
</table>
After receipt of the responses, the CHMP will adopt a timetable for the evaluation of the responses. In general the following timetable will apply:

<table>
<thead>
<tr>
<th>DAY</th>
<th>ACTION</th>
</tr>
</thead>
<tbody>
<tr>
<td>150</td>
<td>PRAC Rapporteur circulates the RMP assessment report and proposed LoOI</td>
</tr>
<tr>
<td>157</td>
<td>Joint Response Assessment Report from CHMP (Co-) Rapporteurs received by CHMP members and the EMA. EMA sends this joint Assessment Report to the applicant making clear that it is sent for information only and does not yet represent the position of the CHMP. Where applicable inspection to be carried out. EMA/QRD sub-group meeting for the review of English product Information with participation of the applicant (optional) around day 165.</td>
</tr>
<tr>
<td>167</td>
<td>PRAC adopts PRAC RMP Assessment Overview and Advice for D180 LoOI</td>
</tr>
<tr>
<td>170</td>
<td>Deadline for comments from CHMP Members to Rapporteur and Co-Rapporteur, EMA and other CHMP members</td>
</tr>
<tr>
<td>180</td>
<td>CHMP discussion and decision on the need for adoption of a list of outstanding issues (LoOI) and/or an oral explanation by the Applicant. If an oral explanation is needed, the clock is stopped to allow the Applicant to prepare the oral explanation. Submission of final inspection report to the EMA, Rapporteur and Co-Rapporteur by the inspection team (at the latest by day 180).</td>
</tr>
<tr>
<td>181</td>
<td>Restart of the clock and oral explanation (if needed).</td>
</tr>
<tr>
<td>183</td>
<td>PRAC Rapporteur circulates the RMP assessment report</td>
</tr>
<tr>
<td>181 to 210</td>
<td>Final draft of RMP, English SmPC, labelling and package leaflet sent by Applicant to the Rapporteur, Co-Rapporteur, PRAC Rapporteur, EMA and other CHMP members.</td>
</tr>
<tr>
<td>197</td>
<td>PRAC adopts the final PRAC RMP Assessment Overview and Advice</td>
</tr>
<tr>
<td>By 210</td>
<td>Adoption of CHMP Opinion + CHMP Assessment Report. Adoption of a timetable for the provision of product information translations</td>
</tr>
</tbody>
</table>

After adoption of a CHMP opinion, the preparation of the annexes to the Commission Decision is carried out in accordance with the following timetable:

<table>
<thead>
<tr>
<th>DAY</th>
<th>ACTION</th>
</tr>
</thead>
<tbody>
<tr>
<td>215 at the latest</td>
<td>Applicant provides to the EMA the product information and Annex A in the 25 languages (all EU languages including Icelandic and Norwegian) and the &quot;QRD Form 1&quot; by Eudralink*.</td>
</tr>
<tr>
<td>229</td>
<td>Member States will send linguistic comments on the product information by e-mail with a copy to the EMA together with QRD Form 1</td>
</tr>
<tr>
<td>235 at the latest</td>
<td>Applicant provides EMA with final translations of SmPC, Annex II, labelling and package leaflet and Annexes IV and 127a if applicable in the 25 languages (+ &quot;QRD Form 2&quot; and “PDF checklist”) by Eudralink.</td>
</tr>
<tr>
<td>237</td>
<td>Transmission of Opinion and Annexes in all EU languages to applicant, Commission, and Members of the Standing Committee, and Norway and Iceland.</td>
</tr>
<tr>
<td>239-261</td>
<td>Draft Commission Decision</td>
</tr>
</tbody>
</table>
DAY | ACTION
--- | ---
Standing Committee Consultation
By 277 | Finalisation of EPAR in consultation with Rapporteur, Co-Rapporteur, CHMP and Applicant (the latter for confidentiality aspects)
277 | Final Commission decision

*By e-mail: qrd@ema.europa.eu

References

- Regulation (EC) No 726/2004
- Centralised Procedure, the Rules governing Medicinal Products in the European Community, Volume 2A, Notice to Applicants, Chapter 4 and Chapter 6
- The linguistic review process of product information in the centralised procedure – human (EMEA/5542/02)
35. How is an EMA Application Number attributed? 

On submission of an application, details of the product are entered into SIAMED, the EMA tracking system.

### 35.1. Procedures

The name and the active substance(s) of the product primarily identify applications for the granting of a Community Marketing Authorisation (MA) for a medicinal product. However, for administrative purposes, each application is also given a core number, EMEA/H/C/xxxxxx, where H stands for Human and C for Centralised Procedure, with the remainder corresponding to a sequentially allocated and unique number identifying the whole of the application. This core number, which is provided after the submission of the application and communicated to the applicant at the start of the procedure, is retained throughout the life cycle of the product.

In every case of an administrative procedure relating to the product, an additional marker denoting the nature of the procedure is appended to this core number, i.e. for the first application for the granting of the MA, any extension, variation, transfer or renewal of MA. A sequential number is added, too. The markers currently used are as follows:

<table>
<thead>
<tr>
<th>Marker</th>
<th>Procedure</th>
<th>Example</th>
</tr>
</thead>
<tbody>
<tr>
<td>/0000</td>
<td>First new application</td>
<td>EMEA/H/C/000789/0000</td>
</tr>
<tr>
<td>N/xxxx</td>
<td>Notification Art. 61(3)</td>
<td>EMEA/H/C/000789/N/0001</td>
</tr>
<tr>
<td>IA/xxxx</td>
<td>Type IA variation</td>
<td>EMEA/H/C/000789/IA/0002</td>
</tr>
<tr>
<td>IB/xxxx</td>
<td>Type IB variation</td>
<td>EMEA/H/C/000789/IB/0003</td>
</tr>
<tr>
<td>II/xxxx</td>
<td>Type II variation (regardless of procedural length)</td>
<td>EMEA/H/C/000789/II/0004</td>
</tr>
<tr>
<td>X/xxxx</td>
<td>Annex II application</td>
<td>EMEA/H/C/000789/X/0005</td>
</tr>
<tr>
<td>S/xxxx</td>
<td>Annual Re-assessment</td>
<td>EMEA/H/C/000789/S/0006</td>
</tr>
<tr>
<td>T/xxxx</td>
<td>Transfer of MA</td>
<td>EMEA/H/C/000789/T/0007</td>
</tr>
<tr>
<td>R/xxxx</td>
<td>Renewal of MA</td>
<td>EMEA/H/C/000789/R/0008</td>
</tr>
<tr>
<td>Z/xxxx</td>
<td>(Renewal of) Suspension of MA</td>
<td>EMEA/H/C/000789/Z/0009</td>
</tr>
</tbody>
</table>

These numbers are used as a reference by the EMA and should be used by the Applicant in all correspondence relating to a certain procedure.

### 35.2. Presentations

In addition, the numbering system covers all presentations (pharmaceutical forms, strengths and pack sizes) of the product. This is mainly relevant during evaluation of the procedure and for the purpose of identifying single presentations in lists such as the Annex A to the opinion. (For correspondence, it is sufficient to indicate the procedural number as above.)
A sequential three-digit number for each presentation is added to the procedural number (core number plus procedural marker). An example is given below for a product consisting of three different presentations, with two ensuing procedures creating new presentations:

<table>
<thead>
<tr>
<th>Procedure</th>
<th>Example</th>
<th>Numbers in Annex A</th>
</tr>
</thead>
<tbody>
<tr>
<td>First new application</td>
<td>EMEA/C/H/000789/0000</td>
<td>EMEA/H/C/000789/0000/001&lt;br&gt;EMEA/H/C/000789/0000/002&lt;br&gt;EMEA/H/C/000789/0000/003</td>
</tr>
<tr>
<td>Type II variation creating three new presentations</td>
<td>EMEA/C/H/000789/II/0004</td>
<td>EMEA/H/C/000789/II/0004/004&lt;br&gt;EMEA/H/C/000789/II/0004/005&lt;br&gt;EMEA/H/C/000789/II/0004/006</td>
</tr>
<tr>
<td>Annex II application creating a further three new presentations</td>
<td>EMEA/C/H/000789/X/0005</td>
<td>EMEA/H/C/000789/X/0005/007&lt;br&gt;EMEA/H/C/000789/X/0005/008&lt;br&gt;EMEA/H/C/000789/X/0005/009</td>
</tr>
</tbody>
</table>

NB: For all procedures creating new presentations, this numbering system is superseded after MA by the EU numbers, which would from then onwards appear in the Annex A to opinions. The EU number is allocated independently of the EMA number, but retains the principle of identifying each single presentation by ending in a three-digit sequential number.
36. How are the European Union MA numbers assigned for new marketing authorisations? *New Feb 13*

The European Commission is responsible for assigning the EU main marketing authorisation number for new marketing authorisation (e.g. EU/1/04/276).

At the time of the adoption of a CHMP opinion for a new marketing authorisation, the Agency will liaise with the European Commission in order to include the EU sub-numbers for each presentation (e.g. EU/1/04/276/01, EU/1/04/276/02, etc.) in the Annex A of the medicinal product, which will be transmitted to the Marketing Authorisation Holder together with the CHMP Opinion and respective annexes.

The Marketing Authorisation Holder should include the assigned EU sub-numbers in all language versions of the Annex A and in all applicable sections of the product information, which are submitted following the CHMP opinion for linguistic review.

The inclusion of the EU sub-numbers in the Annex A transmitted to the Applicant is without presumption as to the outcome of the procedure, which requires the issuance of the Commission decision granting the marketing authorisation.
37. **Which information do I need to provide in my marketing authorisation application regarding GCP Inspections and GLP Compliance? New Mar 15**

Applicants are requested to provide the following information as annexes to the Cover Letter in their marketing authorisation applications:

**Regarding GCP Inspections:**

A list of GCP inspection(s) conducted or planned by any regulatory authority at clinical trial sites for all clinical trials included in the dossier. In case of BE trials a list of the inspections conducted at the clinical and analytical facility where the study was conducted.

Alternatively, a confirmation that no inspections had been requested nor taken place and that no inspection are planned.

Please also refer to Question 35 "When can I expect a pre-approval GCP inspection and how are they conducted?" for more information on GCP Inspections and the information to include in the application regarding GCP compliance.

**Regarding GLP Compliance:**

A summary table, listing the non-clinical studies and indicating for each study:

- study title,
- study code (*Unique identifier assigned to the study*),
- date of completion of the Final Report,
- test facility and test sites in which the study was conducted,
- complete address of the test facility (and test sites where applicable),
- period in which the test facility(ies) and/or test site(s) was(were) used indicating if in that period they were part of an European Union (EU) or an Organisation for Economic Co-operation and Development (OECD) Mutual Acceptance of Data (MAD) accepted GLP monitoring programme.

Regarding GLP compliance, as per Notice to Applicant, Volume 2B, there should be a comment in Module 2.4 Nonclinical Overview and Module 2.6 Nonclinical Summary on the GLP status of the studies submitted in the application.

**References**

38. When can I expect a pre-approval GCP inspection and how are they conducted? Rev. Feb 15

Clinical trials included in any marketing authorisation application (MAA) in the EU and in any subsequent application to the initial one are required to be conducted in accordance with Good Clinical Practices (GCP). GCP inspections are conducted in accordance with Article 15 of Directive 2001/20/EC. The requirements which apply for the conduct of clinical trials included in a MAA are set out in Recital 16 and Article 6(1) of Regulation (EC) No 726/2004 as well as in Annex I to Directive 2001/83/EC, as amended (Introduction and general principles - sections 4 and 8 - and Part 1 - Module 5). Requirements for the conduct of clinical trials and GCP inspections are published in Volume 10 of the Rules governing Medicinal Products in the European Community.

The EMA relies for the scientific review of centralised applications for marketing authorisations for medicinal products on the expertise located in the Member States. The same approach exists in the area of inspections, where inspections are conducted by Member States’ inspectorates if requested by the CHMP. These inspections are co-ordinated by the EMA if they pertain to centralised applications and in the case of GCP inspections, they are conducted by Member States’ inspectorates in accordance with Article 15 of Directive 2001/20/EC. There is a GCP Inspectors Working Group, composed of GCP inspectors from the Member States, which meets quarterly at the EMA.

EMA inspection sector reviews all new applications for evidence of GCP compliance and other validation aspects. All new applications are examined to assess the need for GCP inspection(s). The EMA Inspections Sector liaises closely with the Procedure Manager, Rapporteur and Co-Rapporteur during the pre-submission phase and in the period during and after validation to discuss the need to request GCP inspection(s). A need for inspection(s) may be identified at this stage, based on previous relevant experience of the Inspections Sector and the Member States’ national inspectorates. In addition, a need for GCP inspection(s) may also be identified during the review by the assessors, in particular during the initial assessment phase up to day 120.

GCP inspection issues are usually addressed in the List of Questions (although the inspection may commence earlier once adopted by CHMP), and therefore are usually adopted at Day 120. The GCP inspection(s) of the concerned site(s) can then take place in parallel with the “clock stop” period. However, GCP inspection(s) may be requested by CHMP at any stage of the assessment.

It should be noted that clinical data submitted as a result of specific obligations/follow-up measures, or within variations, extensions or other information received after the initial authorisation (e.g. in relation to safety updates, risk management plan etc…) may also trigger a GCP inspection request.

The Reporting Inspector appointed is usually from the inspectorate of the Member State of the CHMP Rapporteur or Co-rapporteur unless the site(s) to be inspected are located in a single EEA state (or small number (3 or less) of EEA states), in which case that Inspectorate is usually designated as the Reporting Inspectorate.

In addition to the Reporting Inspector, one Lead Inspector is designated per site to be inspected. The Lead Inspector is usually from the Inspectorate of the Member State where the site to be inspected is located (for inspections in the EEA). The Reporting Inspector may also be the Lead Inspector for one or more sites.

In the case of third country inspections, the Reporting Inspectorate and the inspectors are usually from the Rapporteur/Co-Rapporteur country inspectorates.
The applicant is asked to provide information in the application in order to facilitate the review and
where needed the preparation of GCP Inspections. This information should be provided in the
Individual Clinical Study Reports and their Appendices (Module 5) in line with the “Note for Guidance
on the Inclusion of Appendices to Clinical Study Reports in Marketing Authorisation Applications”
(CHMP/EWP/2998/03), and the “Note for Guidance on Structure and Content of Clinical Study Reports”
(CPMP/ICH/137/95). Some of the key information to be provided for each study are listed below with
the specific references to the section numbers given in the “Note for Guidance on Structure and
Content of Clinical Study Reports” (CPMP/ICH/137/95):

- A clear description of the study administrative structure (clear identification of the sponsor and of
  the parties who have performed the monitoring, data management, statistics, laboratory
  assessments, randomization, site(s) of manufacture, site of release in Europe, medical writing,
  other applicable activities and the location of the trial master file) preferably in a tabular form
  and indicating name and address of the site where each activity was performed, responsibilities and
  scope of each activity. These should be identified in the clinical study report of each study, for
  instance in section 6, or appendix 16.1.4.

- A list of investigators (name, address, country), preferably in a tabular form, showing the number
  of patients enrolled by each site, and the total number of sites. In addition a table with the number
  of patients enrolled per country should be included. These should be identified in the clinical study
  report of each study, for instance in section 10.1 or appendix 16.1.4.

- Audit certificates (indicating the sites audited, the dates of audit, the type of audit and the
  auditor). These should be identified in the clinical study report of each study, for instance in
  appendix 16.1.8.

- Signature of the principal or coordinating investigator(s) according to Annex I to Directive
  2001/83/EC as amended and in line with the "Note for Guidance on Structure and Content of
  Clinical Study Reports" (CPMP/ICH/137/95), and not only the signature of the sponsor’s
  responsible medical officer. These should be identified in the clinical study report of each study,
  for instance in appendix 16.1.5.

A list of inspection(s) conducted or planned by other regulatory authorities, related to the product and
trial sites involved, should also be provided, preferably attached to the Application cover letter.

Each clinical study report should contain a statement indicating whether the study was performed in
compliance with Good Clinical Practices (GCP), including the archiving of essential documents.

According to the Notice to Applicant, Volume 2B, the clinical overview (Module 2), should assess the
quality of the design and performance of the studies and also include a statement regarding GCP
compliance.

In addition, in accordance with Article 6(1) of Regulation (EC) No 726/2004, a statement to the effect
that clinical trials carried out outside the European Union meet the ethical requirements of Directive
2001/20/EC should be provided, where applicable, in Module 1.9. This statement should indicate that
"clinical trials carried out outside the European Union meet the ethical requirements of Directive
2001/20/EC" together with a listing of all trials (protocol number) and countries (outside the EU)
involved.
References

- “Centralised procedure” the Rules governing Medicinal Products in the European Community, Volume 2A, Notice to Applicants, Chapter 4
- Directive 2001/20/EC
- Directive 2001/83/EC, as amended
- Regulation (EC) No 726/2004
- “Note for Guidance on the Inclusion of Appendices to Clinical Study Reports in Marketing Authorisation Applications” (CHMP/EWP/2998/03)
- “Note for Guidance on Structure and Content of Clinical Study Reports” (CPMP/ICH/137/95)
- “Clinical trials”, The Rules governing Medicinal Products in the European Community. Notice to Applicants, Volume 10
39. When can I expect a pre-authorisation GMP inspection and how are they conducted? Rev. Feb 15

39.1. Legislative Basis

Directive 2001/83/EC as amended states that Manufacturing Authorisation Holders are obliged to comply with the Good Manufacturing Practice (GMP) for medicinal products and to use as starting materials only active substances that have been manufactured in accordance with the detailed guidelines on Good Manufacturing Practice for starting materials.

The principles and guidelines for GMP for medicinal products for human use are stated in Directive 2003/94/EC. Compliance with these principles and guidelines is mandatory within the European Economic Area (EEA), interpretation of these requirements is provided in part I of the Guide to Good Manufacturing Practice, published in Volume 4 of Eudralex. Part II of this guide provides for the detailed guidelines on Good Manufacturing Practice for active substances used as starting materials. These guidelines are supplemented by a series of Annexes. Part III of the guide includes other related guidance.

Inspections will follow "The compilation of Community procedures on Inspections and exchange of information" which is published by EMA on behalf of the European Commission (http://www.ema.eu.int/Inspections/GMPhome.html).

39.2. Pre-submission notification

In their notification of intention to submit, Applicants should mention:

- The name and the address of the proposed manufacturer(s) of the active substance(s) and finished product
- The name and address of the proposed site(s) in the EEA responsible for batch release of the medicinal product
- If the medicinal product is imported from a third country, it should also include information on GMP inspections of the site(s) concerned carried out in the last 2-3 years by EEA competent authorities and/or by competent authorities of countries where a Mutual Recognition Agreement (MRA) is in operation, where this is applicable.
- Final manufacturing and batch release arrangements will have to be provided when submitting the application
- A description of the roles of all different sites involved. A flow chart is recommended for complex operations.

The manufacturing sites mentioned should be in compliance with Good Manufacturing Practice (GMP) and hence be “inspection ready” at the time of submission of the application and throughout the assessment.

Manufacturing sites in third countries should be aware of European Union GMP requirements as mentioned below.

Once the application is received, it is normally not permitted to add a new site or to change the steps of manufacture/release described in the dossier during the 210-day assessment procedure. Any additional site should be submitted as a variation after the granting of the marketing authorisation.
**39.3. Submission**

On receipt of the application, EMA reviews the information provided on the GMP status of the manufacturing sites involved and determines together with Rapporteur and co-Rapporteur whether to recommend that CHMP makes a request for inspection of the manufacturer of either the active substance or the medicinal product in order to complete the assessment. In addition an inspection request may be triggered by specific issues and questions raised during the assessment of the application.

The performance of these inspections by the EEA competent authorities will be co-ordinated by EMA.

**39.4. Inspection Team**

The inspection team will be drawn from the inspection services of the Supervisory and/or other competent authorities of the EEA. On the advice of the Rapporteur and/or Co-Rapporteur the Inspection Team may include scientific experts and/or a Rapporteur for the Inspection as referred to in the provisions of Article 8 of Regulation (EC) No 726/2004.

**39.5. Type of inspection**

Inspections may be carried out to verify compliance with European Community Good Manufacturing Practice principles and guidelines and/or to cover product or process related issues arising from the assessment of the application. Inspections may cover the following activities:

**39.5.1. Manufacture of the Active Substance**

The detailed guidelines on Good Manufacturing Practice adopted by the EEA for the manufacture of the active substance are contained in part II of the EU Guide to Good Manufacturing Practice (Good Manufacturing Practice for Active Pharmaceutical Ingredients) in "The Rules Governing Medicinal Products in the European Union - Volume 4".

**39.5.2. Manufacture of the Medicinal Product**

The GMP principles and guidelines applying to the manufacture of medicinal products for the EEA are laid down in Commission Directive 2003/94/EC, which are restated along with part I of the EU Guide to Good Manufacturing Practice in "The Rules Governing Medicinal Products in the European Union - Volume 4".

Where a manufacturing site is located in the EEA it is normally not necessary to request an inspection to confirm its GMP status as it is required by the above-mentioned Directive to be regularly inspected by the relevant authorities by virtue of holding a manufacturing authorisation.

An inspection will normally be requested to confirm the GMP compliance status of manufacturing sites in third countries unless satisfactory information is available from an inspection of the same or similar category of product carried out during the last 2-3 years by an EEA competent authority or by the competent authority of a country where a MRA is in operation, when applicable.

In all cases (for sites in the EEA and third countries), an inspection may be requested to cover product or process related issues arising from the assessment of the application. In this case the Rapporteur and/or Co-Rapporteur will provide the Inspection Team with a list of questions/issues, which should be addressed during the inspection.
39.5.3. Importing Site - Site located in the EEA

Importing sites in the EEA are required by the provisions of title IV of Directive 2001/83/EC as amended, to hold a manufacturing authorisation. Inspections of importing sites to confirm their GMP compliance status are not normally requested in connection with applications for marketing authorisations. Inspections may however be requested to cover product or process related issues arising from the assessment of the application. In this case the Rapporteur and/or Co-Rapporteur will provide the Inspection Team with a list of questions/issues, which should be addressed during the inspection.

39.6. Timetable for Inspections

Inspection(s) requested in connection with an application for a marketing authorisation must be carried out and the final report(s) sent to EMA and submitted to the CHMP in accordance with the 210 day time limit for the evaluation of the application by the CHMP.

Once an inspection request is adopted by the CHMP EMA will write to:

- the applicant explaining that an inspection(s) will take place, giving details (target date for carrying out the inspection, inspection team, scope of the inspection, contact person in the relevant authority responsible for arranging the inspection)
- the Rapporteur and Co-Rapporteur for information.

The Inspection Team will contact the Company to agree inspection dates within the agreed target date. Inspections usually take place in parallel with the “clock stop” period and will approximately be conducted within two months from the adoption of the inspection request.

Regarding the fees to be paid, details can be found in the question “What is the fee for a GMP/GCP inspection?”.

39.7. Inspection Reports

Inspectors will send the draft Inspection Report to the manufacturer within fifteen days of the Inspection for comments on major factual errors, point of disagreement or remedial actions. Where necessary, the manufacturer should respond within a further fifteen days to provide comments and, if necessary an action plan with a timetable for implementation. This will be considered during the finalisation of the Inspection Report.

The timing of any discussions, further actions and/or the provision of additional information arising from the inspection will be agreed with the Inspectors and communicated by the Inspectors to the Rapporteur, the Co-Rapporteur and EMA.

Inspectors will finalise the report and send it to EMA by Day 180 at the latest and the Rapporteur, Co-Rapporteur will receive a copy. In case of a non-satisfactory inspection outcome, a non-compliance statement may be issued and it will not be possible to have a positive opinion until the relevant issues have been resolved.

39.8. Documents for inspection

A site master file for use in preparing and carrying out the inspection will be necessary. The preferred format is given in Part III of the GMP guide and is the same as that recommended by the Pharmaceutical Inspection Co-operation Scheme (PIC/S). The Applicant should supply this document
directly to the Inspection Team when requested by it. The site master file is not required to be submitted to EMA.

References

- Regulation (EC) No 726/2004
- Directive 2003/94/EC
- Directive 2001/83/EC
- The rules governing medicinal products in the European Community, Good Manufacturing Practice, Volume 4
- The Rules Governing Medicinal Products in the European Community, the Notice to Applicants, Volume 2A, Chapter 4
40. Which tools are used by the EMA to facilitate the streamlining of the European Decision making process? What is the QRD product information?  

The Quality Review of Documents group (QRD) was established in June 1996 and operates under the mandate adopted by the EMA Management Board on 3 December 1997.

The QRD Group is composed of representatives of the Member State’s national authorities with experience in regulatory affairs and product information and representatives of the EMA (which also chairs the Group and provides secretariat facilities). The European Commission as well as observers from candidate EU countries and the Commission “Centre de Traduction” are invited to participate.

The main task of this group is to ensure clarity, consistency and accuracy of the medicinal product information (summary of product characteristics (SPC), labelling and package leaflet) and of its translations, which will be attached to scientific CHMP opinions. The mandate sets out a series of other tasks, namely:

- Verification of terminology used in translations of Opinions and their consistency with the original version of documents
- Ensuring linguistic and other formal coherence and consistency between different terminology used in scientific Opinions, and promotion of initiatives towards the standardisation of terminology
- Review and update of Opinion templates
- Promotion of legibility of patient information and verification of specimens of sales presentations/mock-ups in all EU official languages
- Consideration of issues which could lead to delays in the Commission’s decision-making process and possible development, on request, of advice (particularly with a view to contribute to the development of common understanding on the implementation of legislation and guidelines)

The mandate also provides that “the Group shall develop its own working methods” and will consider “how best it may be associated with the different stages of the evaluation and Decision-making process”.

In this regard, a New Linguistic Review Process of Product Information has been developed and adopted, providing a more streamlined and more efficient review of the Product Information in all EEA languages.

The new process can be summarised as follows:

**Pre-opinion:**

Before Day 210 two reviews of the English Product Information are performed.

Between Day 80 and 110, a first review is done by the EMA Product Information Quality Group (PIQ) followed by a second review by the QRD group between Day 121 and 165.

The new process also foresees the possibility for one or two Applicant representative(s) to participate to a meeting around Day 165 to discuss the comments and the English Product Information with representatives from the EMA and the QRD.
Post-opinion:

Between Day 215 and 229, a detailed review of all translations of the Product Information is made by the Member States coordinated by the national QRD members concerned.

Between Day 232 and 237, the PIQ reviews the implementation of Member States comments made by the applicants in the final texts.

By Day 237, the final translations are sent to the European Commission to start the external Standing Committee consultation.

As part of a Marketing Authorisation Application, Applicants must submit proposals for SPC, Labelling and Package Leaflet texts in module 1.3.1. using the QRD Product Information Templates.

The Templates:

• are intended to provide applicants with practical advice on how to draw up the product information, but without prejudice to any final position of the EMA, CHMP and European Institutions as to the contents of the document
• set out the standard headings and indicate the most commonly used standard phrases and terms in the 20 official EU languages (with addition of Icelandic and Norwegian)
• define the format and layout for Summary of Product Characteristics (SPC); labelling and Package Leaflet (see also “Convention” to be followed for QRD templates in order to ensure absolute consistency between all language versions)
• provide useful guidance as to the content of the information to be supplied, in the QRD template with explanatory notes

In addition, QRD Reference Documents provide more detail guidance on various aspects concerning terminology and style.

While the templates and guidance notes aim to provide practical hints to the applicants, in particular in relation to how to address common problem areas, they are by no means a comprehensive guide to the information required to be included in the product literature. Thus applicants must also refer to the current EU legislation, guidelines, CHMP notes for guidance etc, when drawing up their drafts in order to be able to fully comply with the legal requirements in respect to product information.

For more details, please visit the QRD Website for all information relating to Product Information and all useful References Documents.

References

• The new Linguistic Review Process of Product Information in the Centralised Procedure (EMEA/5542/02)
• QRD Templates with Explanatory Notes
41. How is a MAA pre-submission meeting conducted at the EMA? Rev. Nov 14

41.1. General principle

The pre-submission meetings represent important points in the product development and regulatory approval process, and relate to the preparatory steps in advance of submitting a request for marketing authorisation application (MAA). Successful pre-submission meetings should enable applicants to submit applications, which are in conformity with the legal and regulatory requirements and which can be smoothly evaluated. These meetings will also enable applicants to establish contact with the EMA Product Team Members who will be closely involved in the centralised evaluation procedure of their medicinal product.

41.1.1. Purpose/scope of meeting

a. MAA pre-submission meetings are aimed at providing applicants with information that will assist them in the finalisation of their upcoming marketing authorisation application. Such meetings typically address product-specific legal, regulatory and scientific issues in order to facilitate subsequent validation and assessment of the application. Pre-submission meetings can be especially helpful to SMEs / other companies that may have limited experience of interaction with the EMA or are unfamiliar with the centralised procedure. However, experience has shown the usefulness of pre-submission meetings even for applicants that already have experience with the centralised procedure, to address issues specific to their upcoming application in view of the constantly evolving regulatory framework and its application.

b. The MAA pre-submission meeting request form provides an overview of the most relevant topics (checklist) that applicants are advised to consider when preparing their upcoming application, and which will be discussed at a MAA pre-submission meeting. For each topic, a reference is included to the corresponding ‘question and answer’ in the EMA Pre-Submission Guidance for Users of the Centralised Procedure (PSG), which is available on the EMA Website. The PSG addresses a number of questions, which users of the centralised procedure may have, together with hyperlinks to relevant legislative documents and procedural guidelines which further complement the advice given in the PSG. The EMA considers that the information provided answers the majority of applicants’ queries. As EMA commits to keeping the pre-submission guidance document updated, there should not be a need to check or confirm the answers given in the PSG document at a pre-submission meeting. A topic should therefore only be proposed for discussion at a pre-submission meeting, in case the applicant’s questions are not fully answered by the PSG or other available guidance documents, due to certain particularities of the upcoming application and/or nature of the product. In that case, applicants are advised to clearly describe the issues in the ‘comments’ box under the topic concerned, and to provide relevant background information. Other topics not listed in the form may be added.

41.2. Timing of MAA pre-submission meetings

Pre-submission meetings for marketing authorisation applications (MAA) usually take place 6-7 months before submission. The MAA pre-submission meeting request form should be sent at least 6 weeks before the proposed meeting date or 3 months in advance if sent together with the Request of Eligibility, so that the meeting can be set-up at a mutually agreed date taking into account availability.
of EMA participants and meeting rooms. The meeting will start with the applicant’s 20-30’ presentation followed by a discussion on the presentation and the topics ticked in the pre-submission request form. The total meeting duration should not exceed 2 hours.

41.3. Who is involved in a MAA pre-submission meeting?

EMA participants at MAA pre-submission meetings are the Procedure Manager (PM) and the EMA Product Lead (EPL) together with the EMA Quality, Risk Management and Regulatory Affairs Product Team Members (PTM). Depending on the topics to be discussed, other EMA staff from the following services/offices and departments may attend parts of the meeting: Orphan Medicines, SME, Paediatric Medicines, Labelling Review and Standards, Scientific Advice, Manufacturing and Quality Compliance, Clinical and Non-clinical Compliance, Product and Application Business Support and Specialised Disciplines Department (Non-clinical, Biostatics, Clinical pharmacology). CHMP/PRAC Rapporteurs and/or assessment team members may also participate in the meeting.

Please note that the PM will be chairing the meeting and will remain the primary contact point between the applicant and the rapporteurs during the procedure.

Applicant’s representatives should not exceed 7 to 8 participants. If needed, additional participants can join via teleconference.

41.4. Documents to be prepared for a MAA pre-submission meeting

- The MAA pre-submission meeting request form needs to be filled in electronically and send to PA-BUS at pa-bus@ema.europa.eu. This form includes topics and questions to be addressed at the pre-submission meeting.

- One of the key-documents to be provided with the MAA pre-submission meeting request form is an overview of the product and its development programme (quality, non-clinical and clinical) together with a draft Table of Contents of the Application, listing the studies performed for each EU-CTD heading and the draft product information.

- Applicants will need to provide a number of documents in relation to the product and the application with the MAA pre-submission meeting request form. In addition, depending on the topics to be discussed, the applicant should provide additional topic-specific information (e.g. draft justification for accelerated review).

- Another important document to be provided is a draft MA Application Form (EU-CTD Module 1.2), which should be completed as far and accurate as possible. The form will provide important information on the product and the type of application (e.g. legal basis, reference product details, manufacturing sites, conditional approval) in relation to the topics to be discussed at the meeting. It will also allow EMA to identify topics, other than those requested by the applicant, for discussion/clarification at the meeting, and thereby preventing issues to be raised at validation. In order to avoid duplication of information, the topics in the pre-submission meeting request form will not require the inclusion of the detailed elements which are already to be provided in the application form (e.g. tick-boxes for legal basis, eligibility for centralised procedure).

- Following receipt of the pre-submission meeting request form and annexed documents, the EMA Procedure Manager will review the topics proposed for discussion. He/she may consider that certain proposed topics would not need to be discussed at the meeting, as they are sufficiently
addressed in existing guidance documents or as they could be easily clarified by phone or e-mail, in order to focus the meeting on particular product-specific issues. The applicant will be informed accordingly in advance of the meeting.

**Note:** Applicants must in all cases comply with all requirements of Community Legislation. Provisions, which extend to EEA countries (i.e. the EU member states, plus Norway, Iceland and Liechtenstein) by virtue of the EEA agreement, are outlined in the relevant sections of the text.

### 41.5. How are MAA pre-submission meetings conducted?

At the start of the meeting, the applicant will be invited to give a 20-30’ presentation on the product development. The applicant’s presentation should include the following topics:

- Company’s participants and contact points during the evaluation
- Brief description of the product
- Brief summary of the dossier content
- Particular EU guideline deviations

On the basis of the information provided, EMA participants will discuss with the applicant the appropriateness of the chosen legal basis in view of the available data, highlight elements to be specifically addressed in the CTD Overviews (e.g. missing data, deviations from scientific advice), will provide an EMA view on the possibility for requesting approval under exceptional circumstances or conditional approval if applicable, etc. EMA may also draw attention to relevant scientific and regulatory guidelines, in particular the CHMP ‘clock-stop’ rules in case of a potential premature submission, recommend (further) scientific advice and suggest improvements to the product information.

The MAA pre-submission meeting request form will serve as the agenda for the remaining of the meeting. The topics listed in the pre-submission meeting request form are grouped according to the following areas:

- Quality + GMP
- Non-clinical + Clinical + GLP + GCP + paediatric + orphan
- Pharmacovigilance
- Regulatory + procedural
- Product information
- Transparency
- Administrative
- Other

It is envisaged that the issues will be addressed in this order at the pre-submission meeting. This will allow a sequential discussion of all the applicant’s questions on topics related to the same area, with involvement of relevant EMA staff with expertise in the area concerned (e.g. Labelling Review and Standards Staff members will attend the discussion on the topics dealing with product information and transparency etc).
Note: Applicants wishing to meet with their appointed (Co-) Rapporteur and assessment teams at national level should also inform the EMA Procedure Manager so that relevant EMA staff from the Product Team could participate in such a meeting via teleconference. In any case, minutes of such meetings should be provided to the EMA PM.

41.6. Follow up of MAA pre-submission meetings

Detailed meeting minutes should be prepared by the applicant and provided to the EMA PM within 2 weeks after the meeting. EMA Product Team Members will subsequently review the minutes within 2 weeks and agree the final (amended) minutes with the applicant.

41.7. Flow-chart summary

References:
- MAA Pre-submission meeting request form
- Human pre-submission Q&A
- Pre-submission procedural advice for users of the centralised procedure
42. Is my medicinal product eligible for approval under exceptional circumstances? *New Jan 06*

42.1. Legal basis and Criteria

The legal basis for the marketing authorisation (MA) under exceptional circumstances is the Article 14 (8) of the Regulation (EC) No 726/2004, and the relevant documentation for applications in exceptional circumstances are laid down in Part II of Annex I of Directive 2001/83/EC, as amended.

Products for which the applicant can demonstrate in this application that he is unable to provide comprehensive data on the efficacy and safety under normal conditions of use, because:

- the indications for which the product in question is intended are encountered so rarely that the applicant cannot reasonably be expected to provide comprehensive evidence, or
- in the present state of scientific knowledge, comprehensive information cannot be provided, or
- it would be contrary to generally accepted principles of medical ethics to collect such information,

may be eligible for marketing authorisation under exceptional circumstances.

Consequently, the authorisation under exceptional circumstances is granted subject to a requirement for the applicant to introduce specific procedures, in particular concerning the safety of the medicinal product, notification to the competent authorities of any incident relating to its use, and action to be taken.

42.2. Prior to submission

As early as possible during drug development, the applicant is encouraged to seek scientific advice from the EMA about the justification for applying for a marketing authorisation under exceptional circumstances, especially on the inability to provide comprehensive data.

Any further discussion on the appropriateness should preferably occur in the context of the presubmission meeting.

42.3. Timing of the submission and Documentation to be supplied

- First of all, the applicant should submit a statement on the appropriateness of the granting of a marketing authorisation under exceptional circumstances in the notification to the EMA of their intention to submit a marketing authorization application (at least 6 months before submission).

- Then, if the applicant considers that the grounds for approval under exceptional circumstances should apply, the applicant should tick the box 1.5.2 of the application form and include its justification in module 1, covering the following aspects:
  1. A claim that the applicant can show that he is unable to provide comprehensive non-clinical or clinical data on the efficacy and safety under normal conditions of use
  2. A listing of the non-clinical or clinical efficacy or safety data that cannot be comprehensively provided
  3. Justifications on the grounds for approval under exceptional circumstances
4. Proposals for detailed information on the specific procedures/obligations to be conducted (Safety procedures, programme of studies, prescription or administration conditions, product information).

The proposals for detailed information on the specific procedures/obligations to be conducted shall also be written in accordance with the “Guideline on risk management systems for medicinal products for human use”.

42.4. **Assessment of the justification for exceptional circumstances**

The Rapporteur, Co-Rapporteur and the other CHMP members will assess the justification/data submitted for exceptional circumstances as part of the overall assessment of the benefit/risk of the application.

It is up to the CHMP, during the review, to ultimately decide on the type of the marketing authorisation.

42.5. **Differences between Exceptional circumstances and conditional marketing authorisation**

<table>
<thead>
<tr>
<th>Conditional Marketing Authorisation</th>
<th>Marketing Authorisation under Exceptional Circumstances</th>
</tr>
</thead>
<tbody>
<tr>
<td>Demonstrate positive benefit-risk balance, based on scientific data, pending confirmation</td>
<td>Comprehensive data cannot be provided (specific reasons foreseen in the legislation)</td>
</tr>
<tr>
<td>Authorisation valid for one year, on a renewable basis</td>
<td>Reviewed annually to reassess the risk-benefit balance, in an annual re-assessment procedure-</td>
</tr>
<tr>
<td>Once the pending studies are provided, it can become a “normal” marketing authorisation</td>
<td>Will normally not lead to the completion of a full dossier and become a “normal” marketing authorisation</td>
</tr>
</tbody>
</table>

A marketing authorisation under exceptional circumstances should not be granted when a conditional marketing authorisation is more appropriate. A conditional marketing authorisation is for example granted in the absence of comprehensive clinical data when it is likely that the applicant will be in the position to provide such data in a short timeframe, whereas the fulfilment of any specific procedures/obligations imposed as part of the marketing authorisation under exceptional circumstances is aimed at the provision of information on the safe and effective use of the product and will normally not lead to the completion of a full dossier.

42.6. **Particularities of the marketing authorisation under exceptional circumstances**

- It should be noted that designated orphan products are eligible for approval under exceptional circumstances only if the criteria considered for the approval under exceptional circumstances are fulfilled.
- The summary of product characteristics and package leaflet should mention that a marketing authorisation has been granted subject to certain specific obligations to be reviewed annually.
The renewal of the marketing authorisation of a medicinal product under exceptional circumstances follows the same rules as a "normal" marketing authorisation. After 5 years, the marketing authorisation will then be renewed under exceptional circumstances for an unlimited period, unless the competent authority decides, on justified grounds relating to pharmacovigilance, to proceed with one additional five-year renewal. (See the renewal guidance).

References

- Regulation (EC) No 726/2004
- Guideline on procedures for the granting of a marketing authorisation under exceptional circumstances, pursuant to Article 14 (8) of Regulation (EC) No 726/2004
- Guideline on risk management systems for medicinal products for human use (EMEA/CHMP/96268/2005)
43. Do I need to perform User Consultation? / When and how to submit information on User Consultation? New July 06

Articles 59(3) and 61(1) of Directive 2001/83/EC, as amended, require that the package leaflet reflects the results of consultations with target patient groups ('user consultation') to ensure that it is legible, clear and easy to use and that the results of assessments carried out in cooperation with target patient groups are provided to the competent authority.

A user consultation is always required in the following situations:

- First authorisation of a medicinal product with a new active substance,
- Medicinal products which have undergone a change in legal status,
- Medicinal products with a new presentation,
- Medicinal products with particular critical safety issues.

However, reference to already approved package leaflets may be acceptable where appropriate, based on a sound justification by the applicant. Examples of when this may be considered acceptable as well as the considerations to be taken into account when choosing the types of ‘reference’ package leaflets are detailed in the “Guidance concerning consultations with target patient groups for the package leaflet”.

If user consultation has been performed on a package leaflet in the old QRD templates, there is no need to be retested when updating according to the new QRD templates. However, it should be noted that compliance with the QRD templates does not exempt from the obligation to undertake a user test or other form of user consultation. See also “What is the QRD product information?”

The package leaflet should be legible, clear and easy to read in all EEA languages, but it is normally sufficient to undertake user consultation in one EEA language. However, results of user consultation should be presented in English in order to allow assessment.

43.1. Methods of user consultation

The legislation does not define a precise method to be used for user consultation.

One of the possible ways of complying with the new legal requirement is by performing a ‘user testing’ of the package leaflet, i.e. to test the readability of a specimen with a group of selected test subjects. It is a development tool which is flexible and aims to identify whether or not the information as presented, conveys the correct messages to those who read it. Testing itself does not improve the quality of the information but it will indicate where there are problem areas which should be rectified.

Other methods than user testing may be acceptable provided that the outcome ensures that the information is legible, clear and easy to use so that patients can locate important information within the package leaflet, understand it and enables the user to act appropriately. Such alternative methodology will have to be justified by the applicant and will be considered on a case-by-case basis.

An example of a method for user testing of a package leaflet is provided in the Annex 2 of "A Guideline on the Readability of the Label and Package Leaflet of Medicinal Products for Human Use".
Further guidance on one way of user testing is also provided in the "EFPIA General Recommendations for Readability User Testing of Package Leaflets for Medicinal Products for Human Use Submitted or Approved under the European Centralised Procedure" and its Annexes (www.efpia.eu).

43.2. Submission and assessment of information on user consultation

During the pre-submission phase the applicant may discuss how to address ‘user consultation’ with EMA and (Co-) Rapporteur, if necessary. This discussion may indicate whether new ‘user consultation’ would be necessary or whether a justification for its absence or ‘focused’ user testing could be acceptable.

At the time of submission of the application, information regarding the ‘user consultation’ performed together with a presentation of its results, or a justification for not performing such consultation, is to be included in Module 1 (Section 1.3.4) of the dossier. The presentation of results should be shortened to a summary explaining how the consultation was executed and how the resulting package leaflet accommodated any need for change. The recommended structure of such a summary is provided in the “Guidance concerning consultations with target patient groups for the package leaflet”.

In their assessment reports, the (Co-) Rapporteur will include the assessment of the results of user consultation or of the justification for its absence as well as a conclusion on the overall readability of the package leaflet. It should be noted that, if not included in the initial submission, the results of user consultation or any further clarification, as requested, will have to be submitted as part of the answers to the list of questions at Day 121.

The user consultation results and the (Co-) Rapporteur’s assessment will also be forwarded to QRD Group, as useful information when reviewing the draft product information.

Further details on the assessment of information on user consultation can be found in the EMA Operational Procedure on Handling of “Consultation with target patient groups” on Package Leaflets (PL) for Centrally Authorised Products for Human Use (EMEA/277378/2005).

References

- Directive 2001/83/EC, as amended
- “Guideline on the Readability of the Label and Package Leaflet of Medicinal Products for Human Use”, the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2C
- “Guidance concerning consultations with target patient groups for the package leaflet”, the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2C
- EMEA Operational Procedure on Handling of “Consultation with target patient groups” on Package Leaflets (PL) for Centrally Authorised Products for Human Use (EMEA/277378/2005)
44. Do I need to include Braille on the Packaging of my Medicinal Product? *New July 06*

Braille is the internationally widespread reading and writing system for blind and partially sighted people. It consists of arrangements of dots which make up the letters of the alphabet, numbers and punctuation marks.

The revised legislation requires that the name of the medicinal product is expressed in Braille format on the packaging of the medicinal product. In addition, Marketing Authorisation Holders must ensure that the package leaflet is made available on request from patients’ organisations in formats appropriate for the blind and partially sighted.

These new requirements apply to new marketing authorisations with Commission Decisions as of 20 November 2005. Nevertheless, companies are encouraged to apply the provision to all centrally authorised medicinal products as soon as possible.

### 44.1. Packaging requirements

The *(invented)* name of the medicinal product followed by its *strength* should be put in Braille on the packaging of the product. The uncontracted Braille system should be used. For medicinal products authorised only in a single strength, it is acceptable that only the invented name in Braille is put on the packaging.

The name in Braille should only appear on the outer/secondary packaging (usually a carton). In case where there is no secondary packaging, it is possible to fix an adhesive Braille label around the bottle.

On a volunteer basis, the name in Braille can be expressed on all packaging components.

It is also possible for companies to include, on a voluntary basis, further information in Braille on bigger volume packages (e.g. pharmaceutical form, expiry date, etc).

In case of multilingual packaging, the name in Braille has to be printed in all the different languages concerned.

It should be noted that there is no need to put the name in Braille on the packaging of products which are only intended for administration by health care professionals.

In case of small volume packages (up to 10 ml) with limited space capacity, alternative means of providing Braille information may be considered, e.g. use of contracted Braille system or certain defined abbreviations or addition of a supplementary “tab” label.

At the time of submission of the application, applicants should address in Module 1 - section 1.3.6 of the application dossier the proposed implementation of the Braille requirements on the packaging of the medicinal product. In addition, the information that will appear in Braille on the printed outer packaging should be mentioned, if applicable, as normal text in section 16 of the outer packaging labelling (Module 1 - section 1.3.1 – Annex IIIA) and, where applicable and feasible, should be indicated with dots on the mock-ups (Module 1 – section 1.3.2).

### 44.2. Package leaflet for blind and partially sighted

On request the package leaflet should be provided for partially sighted people in a suitable print, taking into consideration all aspects determining the readability. For blind people the text has to be provided in an appropriate format, e.g. perceptible by hearing (CD-ROM, audiocassette, etc...) or in Braille.
Choice of the appropriate medium should be made by the MAH in consultation with representatives of organisations for the blind and partially sighted.

Further guidance on the implementation of the requirements for Braille and the requirements for the package leaflet for the blind and partially sighted is provided in the European Commission "Guidance concerning the Braille requirements for labelling and the package leaflet (Article 56a of Directive 2001/83/EC, as amended)". Please note that this guidance will be included in the Commission ‘Guideline on the Readability of the Label and Package Leaflet of Medicinal Products for Human Use’, after finalisation of the revision of the guideline.

References

- Article 56a of Directive 2001/83/EC, as amended
- “Guidance concerning the Braille requirements for labelling and the package leaflet (Article 56a of Directive 2001/83/EC, as amended)"
- “Guideline on the Readability of the Label and Package Leaflet of Medicinal Products for Human Use”, the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2C
45. What is the period of protection for my medicinal product? *New July 06*

45.1. Data exclusivity and market exclusivity period for reference medicinal products

A reference medicinal product is a medicinal product, which has been granted a marketing authorisation by a Member State or by the Commission on the basis of a complete dossier, i.e. with the submission of quality, pre-clinical and clinical data in accordance with Articles 8(3), 10a, 10b or 10c of Directive 2001/83/EC, as amended, and to which the marketing authorisation application for a generic, hybrid or similar biological medicinal product (i.e. application under Articles 10(1), 10(3) or 10(4) of the same Directive) refers (see also "What is the legal basis for my application?").

45.1.1. Submission of the Marketing Authorisation Application (MAA) before 20 November 2005: previous periods of protection

Reference medicinal products authorised through the centralised procedure for which the initial submission was made before 20 November 2005, continue to benefit from the previous periods of protection which are 10 years, (and 10 years for all medicinal products authorised following an opinion of the CHMP in accordance with Article 4 of Directive 87/22/EEC (ex-concertation procedure)).

According to Article 89 of Regulation (EC) No 726/2004, the new periods of protection do not apply to those reference medicinal products for which the initial application for authorisation (date of submission of the application and not validation) was submitted before 20 November 2005.

45.1.2. Notion of global marketing authorisation / Particular case of “Fixed combinations”

The *global marketing authorisation* contains the initial authorisation and all variations and extensions thereof, as well as any additional strengths, pharmaceutical form, administration routes or presentations authorised through separate procedures and under a different name, granted to the marketing authorisation holder of the initial authorisation.

In accordance with Article 6(1) of Directive 2001/83/EC, as amended, all these presentations of a given product shall be considered as part of the same marketing authorisation for the purposes of applying the rules on data and marketing protection.

This means that for a reference medicinal product, the start of the data and market exclusivity periods is the date when the first marketing authorisation was granted in the Community. New additional strengths, pharmaceutical form, administration routes, presentations as well as any variation and extensions do not restart or prolong this period. This will apply even if the new presentation has been authorised to the same marketing authorisation holder through a separate procedure and under a different name.

The “*fixed combinations*” are not considered part of the global marketing authorisation and will benefit from an independent period of protection.
45.1.3. Submission of the MAA after 20 November 2005: new periods of protection

Directive 2001/83/EC, as amended, and Regulation (EC) No 726/2004 have introduced new rules concerning the periods, from the initial marketing authorisation of the reference product, during which generic, hybrid or similar biological medicinal products’ applicants cannot rely on the dossier of the reference product for the purposes of submitting an application, obtaining a marketing authorisation or placing the product on the market.

Applications for generic, hybrid or similar biological medicinal products can be submitted after a so-called “data exclusivity” period of 8 years from initial authorisation of the reference medicinal product. Generic, hybrid or similar biological medicinal products authorised in this way can be placed on the market after a so-called “market exclusivity” period of 10 years from initial authorisation of the reference medicinal product.

45.2. One year period of protection for new indications of well-established substances

According to Article 10(5) of Directive 2001/83/EC as amended, “where an application is made for a new indication for a well-established substance, a non-cumulative period of one year of data exclusivity shall be granted, provided that significant pre-clinical or clinical studies were carried out in relation to the new indication.”

The data exclusivity period refers exclusively to the data concerning the new indications.

Commission Decisions authorising new therapeutic indications for well-established substances will contain a clear statement of whether the new indication is based on significant pre-clinical or clinical studies.

A well-established substance is an active substance included in the relevant medicinal product which can be shown to have a well-established use in accordance with the requirements of indent (a) in section 1 (“Well established medicinal use”) of Part II of the Annex to Directive 2001/83/EC as amended. This does not however mean that the medicinal product concerned must have been authorised under the legal basis of the well-established use procedure.

A new indication submitted after 20 November 2005 may benefit from this year of protection.

45.3. One-year period of protection for data supporting a change of classification

According to Article 74a of Directive 2001/83/EC as amended reads: “Where a change of classification of a medicinal product has been authorised on the basis of significant pre-clinical tests or clinical trials, the competent authority shall not refer to the results of those tests or trials when examining an application by another applicant for or holder of marketing authorisation for a change of classification of the same substance for one year after the initial change was authorised.”

The 1-year period of protection covers significant pre-clinical or clinical trials carried out for the purpose of substantiating an application for a change of classification. The interpretation by competent authorities of the notion of significant pre-clinical tests or clinical trials under Article 74a will be without prejudice to the interpretation of that phrase under Article 10(5) of the Directive.

When adopting a decision authorising a change of classification of a medicinal product, the competent authority must assess whether the change is based on significant pre-clinical tests or clinical trials. In the case of products authorised in accordance with Regulation (EC) No 726/2004, Commission
Decisions authorising a change of classification will contain a clear statement of whether the change is based on significant pre-clinical tests or clinical trials (see also “Guideline on changing the classification for the supply of a medicinal product for human use”).

A change of classification authorised after 20 November 2005 may benefit from this year of protection.

**45.4. Extension of the ten-year period of marketing protection in the case of new therapeutic indications (8 + 2 +1)**

In accordance with Article 14(11) of Regulation (EC) No 726/2004, the ten-year period of marketing protection (8+2) may be extended by 1 year in the event of authorisation of new therapeutic indications but only if:

- The new application represents a significant clinical benefit in comparison with existing therapies,
- The new indication is granted during the first eight years since the initial marketing authorisation.

This additional year of marketing protection applies to the global marketing authorisation for the reference medicinal product. Generic, hybrid or similar biological medicinal products, with or without the new therapeutic indication, may not be placed on the market until expiry of the eleventh year.

The overall period of protection cannot exceed eleven years. Therefore, this provision can be used only once per ‘global marketing authorisation’ within the meaning of Article 6(1) of Directive 2001/83/EC as amended.

Commission Decisions authorising new therapeutic indications will contain a clear statement of whether the new indication represents a significant clinical benefit in comparison with existing therapies.

*This year of protection shall apply only to those reference medicinal products for which the initial application for authorisation is submitted after 20 November 2005.*

Detailed information on market exclusivity for orphan medicinal products is provided in the "Communication from the Commission on Regulation (EC) No 141/2000 on orphan medicinal products" (section D) and in the draft "Guideline on aspects of the application of Article 8 of Regulation (EC) No 141/2000".

**References**

- Regulation (EC) No 726/2004
- Directive 2001/83/EC, as amended
- The Rules Governing Medicinal Products in the European Community, Notice to Applicants, Volume 2A, Chapter 1
- "Guideline on elements required to support the significant clinical benefit in comparison with existing therapies of a new therapeutic indication in order to benefit from an extended (11 years) marketing protection period”
- "Guideline on changing the classification for the supply of a medicinal product for human use”, the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2C
- Communication from the Commission on Regulation (EC) No 141/2000 on orphan medicinal products
European Commission Guideline on aspects of the application of Article 8 of Regulation (EC) No 141/2000: Assessment of similarity and/or clinical superiority of orphan medicinal products when assessing marketing authorisation applications and variations
46. How shall I submit my EU Risk Management Plan as part of my application? **Rev. March 13**

46.1. Description of the risk management system

Article 8(3) ((iaa)) of Directive 2001/83/EC requires that a marketing authorisation application (MAA) shall include the risk management plan describing the risk management system which the applicant will introduce for the medicinal product concerned, together with a summary thereof.

A Risk Management Plan (RMP) is a detailed description of the risk management system that in itself is a set of pharmacovigilance activities and interventions designed to identify, characterise, prevent or minimise risks relating to medicinal products including the assessment of the effectiveness of those activities and interventions.

The aim of a risk management system is to ensure that the benefits of a particular medicine (or a series of medicines) exceed the risks by the greatest achievable margin for the individual patient and for the target population as a whole.

46.2. When should I submit my RMP?

An initial RMP or an update may need to be submitted at any time during a product’s lifecycle, i.e. during both pre- and post-authorisation phases. Detailed information when an initial or updated RMP should be submitted can be found in Guideline on good pharmacovigilance practices (GVP) Module V – Risk management systems.

For initial marketing authorisation applications the RMP should be submitted together with the application dossier in eCTD module 1.8.2. A RMP or an update of the RMP is also expected together with an application involving a significant change to an existing marketing authorisation (extension of indication, line extension, new manufacturing process of a biotechnologically-derived product); and at the request of the Agency or national competent authority.

46.3. When should updates of the RMP be submitted? **Rev. July 2013**

The MAH should submit updates of the RMP if information becomes available that has an impact on the benefit-risk profile of the medicinal product(s) included in the RMP. E.g. when new information is received that may impact on the current Safety Specification, Pharmacovigilance Plan, assumptions regarding efficacy or risk minimisation measures; or within 60 days of an important (pharmacovigilance or risk minimisation) milestone being reached. An update of the RMP should also be submitted at the request of a Competent Authority.

Updates to the RMP should also be submitted with an application involving a significant change to an existing marketing authorisation (extension of indication, line extension, new manufacturing process of a biotechnologically-derived product) if there is already an existing RMP in place.

Since February 2013, the CHMP Opinion will not include fixed time schedules for the submission of routine updates of the RMP (e.g. ‘annually until first renewal’). If routine updates of the RMP are still mentioned in the Annex II (no update to latest QRD template has taken place yet), this provision is no longer applicable.

In specific cases, the CHMP and PRAC may provide a specific timeline for the update of a RMP and this will be reflected in the relevant Opinion. GVP Module V will reflect this new position in the next update.
Applicants are generally encouraged to contact the EMA prior to submitting new applications to discuss RMP related questions.

At any stage, but in particular during the pre-authorisation phase, a MAA/MAH may request advice on the development or content of an EU-RMP through the scientific advice procedure.

Whether or not the scientific advice procedure has been used, discussion on any questions relating to the EU-RMP for a medicinal product seeking a new authorisation through the centralised procedure should take place at the pre-submission meeting.

**46.4. When can I submit a RMP within a procedure? Rev. Feb 2014**

A RMP can be submitted as part of an initial marketing authorisation application. Furthermore, a RMP or RMP update can be submitted as part of a renewal, or a regulatory procedure involving a change to an existing marketing authorisation (e.g. extension of indication, line extension, new manufacturing process of a biotechnologically-derived product).

Also, if a change to the RMP is necessary based on a safety variation to update the Summary of Product Characteristics, Labelling or Package Leaflet, the RMP can be submitted within that variation procedure.

If final study results are submitted for assessment through a variation, and the finalisation of the study leads to the need to update the RMP, this RMP update can be submitted as part of that variation. However, if interim results of a study lead to the need for an updated RMP (addition/deletion of safety specifications) a stand-alone variation for the RMP update should be submitted.

A RMP update can only be submitted together with a PSUR od a single centrally authorised medicinal product when the changes to the RMP are a direct result of data presented in the PSUR.

As an interim measure, submission of RMP updates cannot be accepted together with the PSURs of medicinal products (centrally and/or nationally authorised) subject to a PSUR EU single assessment (PSUSA). MAHs should update their RMP through another upcoming procedure affecting the RMP or alternatively, through a separate variation which can be submitted after finalisation of the PSUR single assessment procedure.

Any (update of the) RMP is provided outside another regulatory procedure should be submitted through a separate stand-alone variation.

**46.5. When is my RMP a stand-alone variation? New July 2013**

If a new/initial RMP is submitted outside of another regulatory procedure, this RMP should be submitted as a stand-alone variation.

A stand-alone variation for updates of the RMP is foreseen when safety concerns are added or deleted outside another procedure. For instance, if interim results of a study lead to the need to add or to delete safety specifications a stand-alone variation for the RMP update should be submitted.

A further submission of an updated RMP as a stand-alone variation is expected when the MAH proposes changes to already previously agreed category 3 studies in Part III.4.3 of the RMP. This applies also when the MAH provides an updated / amended protocol that changes the previously accepted protocol with an impact on the Part III.4.3.
46.6. Can I submit a RMP within my PSUR/PSUSA procedure? Rev. Feb 2014

- PSUR for single centrally authorised medicinal product

A RMP update of a single centrally authorised medicinal product can only be submitted together with a PSUR when the changes to the RMP are a direct result of data presented in the PSUR. In this case no stand-alone RMP variation is necessary. Should only the timing for submission of both documents coincide, but not be related to each other, the RMP submission should be handled as a stand-alone variation.

- PSUR EU single assessment (in reference to the EURD list)

As an interim measure, submission of RMP updates cannot be accepted together with the PSURs of medicinal products (centrally and/or nationally authorised) subject to a PSUR EU single assessment (PSUSA). MAHs should update their RMP through another upcoming procedure affecting the RMP. Alternatively MAHs should submit a separate variation to update their RMP, after finalisation of the PSUR single assessment procedure.

For nationally authorised medicinal products, RMP updates should be submitted to the National Competent Authority for assessment.

46.7. How shall I present my RMP?

Guidance on the format and content of the RMP as outlined in GVP module V has been made available in the Pharmacovigilance section of the Agency’s website. The submitted RMP should follow the RMP guidance.

- All new procedures submitted after the 10th of January and subsequently starting containing a (updated) RMP as well as stand-alone RMP submissions should follow the guidance and be submitted in the new format.

- All initial MAAs and line extensions, having been submitted before the 10th of January, but having their D121 or D181 responses submitted after the 10th of April will have to comply with the new RMP format, i.e. the responses will always have to contain an updated RMP according to the new guidance.

- All other post-authorisation procedures containing a RMP or stand-alone RMP submissions, having been submitted before the 10th of January will not have to provide within responses an updated RMP in line with the guidance.

In certain circumstances certain parts or modules of the RMP may be omitted, unless otherwise requested by the competent authority. Specific details can be found in Guideline on good pharmacovigilance practices (GVP) Module V – Risk management systems, paragraph V.C.3.1.

For updates of existing RMPs, clean and track change versions should be submitted along with a cover letter detailing the changes since the last submitted version.

For versioning of the RMP the numbering should be consecutive and without added text. The numbering is independent of whether the RMP was endorsed by the CHMP or not. The new version of the RMP should be dated.

46.8. How do I submit the EU RMP in eCTD?

The revised RMP should be provided in CTD section 1.8.2
Until further notice companies have to send in all parts and modules of the RMP in one single PDF-file so that a complete RMP is provided to the Agency. A cover letter stating which parts and/or modules of the RMP have actually been updated should be provided. Part I of the RMP also presents this information and should always be updated.

46.9. Should I provide documents with tracked changes highlighted to facilitate review? Rev. Aug 2014

Only clean versions of documents in PDF format should be managed within the eCTD lifecycle. If additional formats are required by any authority to facilitate the assessment (e.g. tracked changes versions for SmPCs, Risk Management Plans or other documents as specified by the agency), these should be provided in Word format in the separate folder ‘XXXX-working documents’. Further details can be found in section 2.9.9 of the TIGes Harmonised Guidance for eCTD Submissions in the EU.

When submitting an existing RMP for the first time in the new format, track change should not be used. However, a content change overview should be provided to identify which parts/sections of the RMP have been updated as a result of new data – ie those parts which would have subject to track changes if the RMP had been submitted in the old format.

46.10. Can more than one draft RMP be submitted for regulatory assessment?

Different sections of the approved version of the RMP can be under review as part of different procedures. Any submitted version of the RMP should be based on the latest approved version and should be seen as a draft, until approved. Details of the RMP approval status should be provided in the Module I of the document.

In line with current guidance (GVP Module V) an RMP update and submission for regulatory review should only be considered when significant new safety information becomes available, unless otherwise requested. For example, the increasing number of exposed patients post-marketing on its own would not represent significant new information for submission of an updated RMP.

If two or more draft RMPs are under evaluation in the context of overlapping procedures (e.g. an RMP update is submitted before the assessment of the RMP previously submitted in the context of another procedure is concluded), at the opinion of the procedure that is finalised last, the MAH should ensure that the approved RMP version includes all the amendments approved in the draft RMPs previously assessed.

Companies are strongly encouraged to streamline RMP amendments and submissions, in co-operation with the EMA (or Reference Member State for non-CAPs), in order to facilitate RMP assessments throughout the product life-cycle.

46.11. When should study progress reports be submitted?

The timelines of the progress reports should be pre-specified and indicated in the protocol. These progress reports may include available interim results, but there is no obligation or recommendation to include interim results in PSURs and RMPs unless required as part of an agreed pharmacovigilance plan.
46.12. **How long after the European Commission decision should Annex 1 of the RMP be submitted to EudraVigilance?**

There is 30 days to submit the Annex 1 of the RMP to EudraVigilance.

46.13. **Do I need to submit an RMP for my traditional herbal medicinal product?**

The submission of a risk management plan is not required for an application for a traditional-use registration.

For other herbal medicinal products not falling within the scope of the traditional-use registration, an RMP will be required for any initial marketing authorisation applications.

46.14. **What are the requirements for a RMP for a new application of an established generic product?**

Marketing authorisation applications for a generic medicinal product under Article 10(1) of Directive 2011/83/EC submitted after 2nd July 2012 will have to include a RMP in the application dossier, however as outlined in the Good Pharmacovigilance Practice (GVP) module V on risk management systems some parts or modules of the RMP for a generic may be omitted (see GVP V page 54-55).

46.15. **If there is no RMP in place for a reference medicinal product, how should module SVIII ‘summary of the safety concerns’ be populated for a generic medicinal product?**

The company of the generic medicinal product should use the (E)PAR and the SmPC of the reference medicinal product to obtain the safety concerns to be included in module SVIII of the RMP. Companies may also discuss with the relevant competent authority what safety concerns should be included.

46.16. **Should I include all of my ongoing studies in the RMP?**

If any or all of the safety issues are safety concerns in the RMP, then a study should be included in the PhV plan, even if, e.g. it would be an observational study using insurance claims data to characterize safety issues. According to the new format, it should be included in:

III.1. Safety concerns and overview of planned pharmacovigilance actions

And its category should be proposed as

III.4.3. Additional pharmacovigilance activities required by the CHMP/PRAC to address specific safety concerns or to measure effectiveness of risk minimisation measures

In addition, if some particular off-label use is a safety concern – either because it is a contraindication and/or use is likely outside of the approved indication – then studies outside of the target population or with a different dose, investigating the safety concern are appropriate in the PhV Plan.

Studies in the PIP should not be routinely included in the PhV Plan but any PIP recommendations for long term follow up of safety or efficacy issues should be specifically discussed in section SVI.6 of the RMP. Where use in children is a safety concern it may be appropriate to include individual activities aimed at providing further safety information in the PhV Plan. The aim here is to allow the safety concern to be investigated, not to provide studies reflecting the development plan for a paediatric indication.
46.17. **How is the assessment of an educational program as additional risk minimisation handled?**

The outlines of the educational program (i.e. the key elements) are part of Annex II.D of the marketing authorisation for centrally authorised medicinal products. Assessment of the educational program incorporating these key elements is done at the Member State level since GVP Module V chapter V.C.7 states that Member States have the responsibility for ensuring that the key elements described in the conditions and/or restrictions are implemented by the marketing authorisation holder in their territory.

46.18. **Can the internet be used as additional risk minimisation measure (e.g. website with educational material/video)?**

Use of websites should not be proposed in the RMP as a means of communicating information on additional risk minimisation measures. Mention of a specific medicinal product on a website is regarded as promotional in some Member States and may not be permissible. However, in some Member States it is possible that use of the internet may be permitted as part of the national communication plan agreed at Member State level.

46.19. **Should I submit a summary of the RMP and will it be published?**

Yes, the summary of the RMP is a mandatory element of any RMP submission and the Agency will make it publically available.

Guidance on the format and content of the RMP for MAA/MAH, including the RMP summary, is published on the Agency’s website. Part VI contains summarised information on the RMP, which will be prepared by the MAH and assessed as part of the normal evaluation process.

A RMP summary is required for all medicines authorised. The current approach is that subsequent updates in the new format will be used to ensure all medicines have a RMP summary.

The final format of the RMP summary and processes for its production and publication are still subject to discussion. Further details will be published on the Agency website and those of national competent authorities (as appropriate) as soon as these are available. However, it is highly likely that the published summary will be based on the elements provided in Part VI of the RMP.

46.20. **How will the PRAC be involved in the review of my RMP? Rev. July 2013**

The oversight of RMPs for products authorised centrally lies with the PRAC. The PRAC is involved in the following procedural and scientific matters regarding RMPs since September 2012:

- Initial MAA
- For Extension application / Extension of indication when including a new or updated RMP
- Updated RMP submitted together with a PSUR if the changes to the RMP are a direct result of the data presented in the PSUR
- For new or updated RMPs submitted as part of a variation

The minutes of the PRAC meeting will be published on the EMA website and contain information about RMPs assessed by the PRAC.
46.21. **Can I submit a version of the RMP after the Opinion to reflect the last minute changes made during the CHMP? New May 13**

A tidied version of the RMP which was agreed at the time of the Opinion should be submitted within 15 days of the Opinion to facilitate publication of the Summary of the RMP and the EPAR. It is important to make it clear that this is not a new version, but a finalised version of what was agreed at the time of the Opinion – therefore, it will keep the version number of the agreed Opinion RMP and have a suffix ‘W’. It should be submitted via Eudralink and should not contain any new data.

**e.g.**:

Version 8.0: RMP agreed at the time of the Opinion but requiring slight amendments due to last minute changes agreed at the Opinion

Version 8.0(W): word version of amended RMP submitted within 15 days of the Opinion

Version 8.0: RMP (including amendments) in closing eCTD sequence.
47. Pharmacovigilance system  

47.1. Requirements regarding the summary of the pharmacovigilance system

Applicants for marketing authorisation are required to provide a summary of their pharmacovigilance system, in accordance with Article 8(3) (ia) of Directive 2001/83/EC, which they will introduce once the authorisation is granted.

The requirement for the summary of the pharmacovigilance replaced the previous requirement for the submission of a detailed description of pharmacovigilance system in the application for marketing authorisation.

The summary of the pharmacovigilance system should be provided in Module 1.8.1 of the application for marketing authorisation and includes the following elements:

- proof that the applicant has at his disposal a qualified person responsible for pharmacovigilance,
- the Member States in which the qualified person resides and carries out his/her tasks,
- the contact details of the qualified person,
- a statement signed by the applicant to the effect that the applicant has the necessary means to fulfil the tasks and responsibilities listed in Title IX of Directive 2001/83/EC,
- a reference to the location where the pharmacovigilance system master file (PSMF) for the medicinal product is kept.

The applicant may combine this information in one single statement using the required statement as per Article 8(3)(ia) of Directive 2001/83/EC regarding the obligation to have the necessary means to fulfil the tasks and responsibilities listed in Title IX. Such statement should be signed by an individual who can act on behalf of the legal entity of the applicant/MAH and by the qualified person for pharmacovigilance (QPPV). The title, role and responsibility of each individual signing the statement should be clearly specified in the document.

The summary of pharmacovigilance system is specific to each marketing authorisation application as per legislation and therefore should be signed by the relevant applicant/MAH.

Applicants are required to include a summary of the applicant pharmacovigilance system at the time of submission of an initial marketing authorisation application (MAA).

The requirement for the summary of the pharmacovigilance system is the same for any marketing authorisation application, independent of the legal basis for the application.

47.2. Requirements regarding the pharmacovigilance system and pharmacovigilance system master file

The MAH has to operate a pharmacovigilance system for the fulfilment of his pharmacovigilance tasks.

The pharmacovigilance system master file (PSMF) is a detailed description of the pharmacovigilance system used by the MAH with respect to one or more authorised medicinal products.

The PSMF is not part of the marketing authorisation (MA) dossier and is maintained independently from the MA. It should be permanently available for inspection and should be provided within 7 days to the Competent Authorities if requested. The PSMF must be located either at the site in the Union where
main pharmacovigilance activities of the marketing authorisation holder are performed or at the site in the Union where the QPPV operates. The QPPV has to both reside and operate in the Union.

Applicants are required, at the time of initial MA application, to have in place a description of the pharmacovigilance system that records the system that will be in place and functioning at the time of granting of the MA and placing of the product on the market. During the evaluation of a MA application the applicant may be requested to provide a copy of the PSM for review.

The PSM has to describe the pharmacovigilance system in place at the current time. Information about elements of the system to be implemented in future may be included, but these should be clearly described as planned rather than established or current.

The pharmacovigilance system will have to be in place and functioning at the time of granting of the MA and placing of the product on the market.

47.3. **Subcontracting pharmacovigilance activities**

The MAH may subcontract certain activities of the pharmacovigilance system to third parties. He will nevertheless retain full responsibility for the completeness and accuracy of the pharmacovigilance system master file (PSMF).

The MAH will have to draw up a list of its existing subcontracts between himself and the third parties, specifying the product(s) and territory(ies) concerned.

When delegating any activities concerning the pharmacovigilance system and its master file, the MAH retains ultimate responsibility for the pharmacovigilance system, submission of information about the PSMF location, maintenance of the PSMF and its provision to competent authorities upon request. Detailed written agreements describing the roles and responsibilities for PSMF content, submissions and management, as well as to govern the conduct of pharmacovigilance in accordance with the legal requirements, should be in place.

For more guidance on the requirements for pharmacovigilance system and PSMF, please refer to the relevant Good Vigilance Modules.

47.4. **Pharmacovigilance system master file number (PSMF)**

Applicants are encouraged to request a PSMF number (MFL EVCODE) in advance of the marketing authorisation application.

If available, the PSMF number (MFL EVCODE) assigned by the extended EudraVigilance Medicinal Product Dictionary (XEVMPD) should be included in the statement in Module 1.8.1. However this information is not part of the compulsory elements as per Article 8(3)(ia) of Directive 2001/3/EC.

For more information on how to obtain a PSMF number, please refer to the Detailed Guidance on electronic submission of information on medicines.

**References**

- Directive 2001/83/EC
- Directive 2010/84/EU

• HMA-EMA Questions and answers on practical transitional measures for the implementation of the pharmacovigilance legislation (EMA/228816/2012 – v.3)

• Guideline on good pharmacovigilance practices - Module I – Pharmacovigilance systems and their quality systems (EMA/541760/2011)

• Guideline on good pharmacovigilance practices - Module II – Pharmacovigilance system master file (EMA/816573/2011)

• EMA Post-Authorisation Guidance regarding the pharmacovigilance system

• Detailed Guidance on electronic submission of information on medicines
48. **What is the CHMP Peer Review?** *Rev. Feb 15*

Peer review is a process by which other members of the CHMP review the (Co) Rapporteurs’ scientific evaluation, as well as the validity of the scientific/regulatory conclusions reached. It applies during the initial phase of the assessment of a new Marketing Authorisation Application (MAA).

Peer review is part of a quality assurance system established at CHMP level. That is the review of the (Co) Rapporteurs’ assessment reports for the purpose of improving the quality of the day 120 List of Questions by those CHMP members that are assigned by the Committee as peer reviewers. It is also the particular task of those members assigned as peer reviewers to judge the quality of the assessment reports from (Co) Rapporteurs especially in relation to potential divergencies in scientific assessment made by (Co) Rapporteurs.

A strengthened peer review system that can improve the consistency of scientific assessments is one of the objectives set out in the EMA Road Map


On appointment of (Co) Rapporteurs during a CHMP meeting, the Committee also appoints Peer Reviewers. The Peer Reviewers are appointed from amongst the members of the CHMP (including co-opted members) or CHMP alternate members and are identified after having put their names forward on a nomination form (nomination form for Rapporteurs). The Committee also decides on the scope of the Peer Review (modules 3, 4, and/or 5) and the number of Peer Reviewers to be assigned to this task.

On Day 112 of the procedure, a Dialogue (e.g. teleconference) is set up between (Co) Rapporteurs, Peer Reviewers and EMA staff to discuss and critically analyse the different objections and concerns raised in the (Co) Rapporteur’s “Overview and draft List of Questions”.

Peer Reviewer’s comments are not made available to applicants. Moreover, it is not intended that applicants directly contact Peer Reviewers or other CHMP members in the context of an ongoing CHMP assessment of a MAA.

**References**

- Notice to Applicants Volume 2A, Chapter 4 Centralised Procedure
49. How can I request a meeting with Rapporteurs to clarify the questions posed by the Committee? **NEW May 2015**

After the receipt of the adopted List of Questions or List of Outstanding Issues and prior to the formal submission of the responses, the applicant can request a clarification meeting with the (Co-) Rapporteurs (from CHMP, PRAC and/or CAT, as relevant) and the EMA (EMA Product Lead and other relevant team members as appropriate). The aim of these meetings is to provide clarifications and guidance to the applicant on the rationale for the Major Objections and/or other issues and to discuss with the Applicants their response strategy and potential need to adjust the response timelines. Such meetings are intended to avoid the submission of inadequate, incomplete or premature responses potentially leading to prolongation of the procedure. It should be emphasised that these meetings are not intended to provide a pre-assessment of the intended responses. These meetings will usually take place via teleconference.

Applicants are advised to refer to “Guidance on meetings with applicants on the responses to questions received from EMA Scientific Committees during the evaluation within the centralised procedure” for further guidance.

**References**

- Guidance on meetings with applicants on the responses to questions received from European Medicines Agency Scientific Committees during the evaluation within the centralised procedure
50. What is an oral explanation and how is it conducted?

NEW May 2015

An oral explanation can be requested either by the applicant or by the relevant EMA committee – the CHMP, CAT (for advanced therapy medicinal products) or, exceptionally, PRAC. Oral explanations are intended to give opportunity to the applicant to explain their position and arguments. They are usually organised when still at Day 180 of the procedure there are major objections concerning the application, which would prevent the Committee from adopting a positive Opinion on the application. It is important that applicants preparing for an oral explanation bear in mind that they are held to only allow clarification of the aspects relating to the outstanding issues.

When the applicant wishes to have the opportunity of an oral explanation, they should present a written request to the relevant committee preferably one month before the anticipated date of the oral explanation and in all cases prior to Day 180. Such request should be sent to the EMA Procedure Manager.

Applicants are advised to refer to “Guidance to applicants / Marketing Authorisation holders on oral explanations at EMA” for practical guidance on preparation for and conduct of oral explanations. Applicants are also reminded that oral explanations are only held in English.

References

- Guidance to applicants /marketing authorisation holders on oral explanations at EMA
51. Where can I find the relevant documents regarding the pharmaceutical legislation? *Rev. March 13*

The **Treaties** on which the European Union and the European Communities are founded can be found on the European Union website: http://eur-lex.europa.eu/en/treaties/index.htm

To exercise the Union’s competences, the institutions may adopt **regulations, directives, decisions, recommendations and opinions**.

Information about the hierarchy of the European Union texts can be found in the Annex I to Chapter 1 of the Notice to Applicants (the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2A, Chapter 1)

The “**Rules governing medicinal products in the European Union**” concerning medicinal products for human use is published on the European Commission website:

  - **Volume 1** – Pharmaceutical legislation, contains most of the relevant Directives, Regulations, Decisions and Communications
  - **Volume 2** – Notice to Applicants (mentioned above)

**Volume 2A** - Procedures for marketing authorisation, is organised as follows:

  - Introduction
  - Chapter 1 – Marketing Authorisation
  - Chapter 2 – Mutual Recognition
  - Chapter 3 – Community Referral
  - Chapter 4 – Centralised Procedure
  - Chapter 5 – Variations
  - Chapter 6 – Community Marketing Authorisation

**Volume 2B** - Presentation and content of the dossier, provides guidance for the compilation of dossiers for applications for marketing authorisation, and is applicable for the centralised procedure and national procedures, including mutual recognition and decentralised procedures.

**Volume 2C** - Regulatory Guidelines, is related to procedural and regulatory requirements e.g. renewal procedures, variation procedures, summary of product characteristics (SPC), package information and classification for the supply, readability of the label and package leaflet requirements.

- **Volume 3** – Scientific guidelines
- **Volume 4** – Good Manufacturing Practices
- **Volume 9** – Pharmacovigilance

With the application of the new pharmacovigilance legislation as from July 2012 Volume 9A is replaced by the good pharmacovigilance practice guidelines (GVP)
released by the European Medicines Agency. However, until the availability of the respective GVP modules Volume 9A remains the reference.

The GVP modules refer to the Commission implementing regulation No 520/2012 of 19 June 2012 on the performance of pharmacovigilance activities. This is a legally binding act published by the European Commission in June 2012 which provides details on the operational aspects for the new legislation: http://eur-lex.europa.eu/LexUriServ/LexUriServ.do?uri=OJ:L:2012:159:0005:0025:EN:PDF Volumes 5, 6, 7 and 8 apply only to veterinary medicinal products

Volume 10 – Clinical trials

The European Commission website offers the possibility to create a CD-Rom with the content of the "Rules governing medicinal products in the European Union" which can be used off-line with an integrated search engine.

The scientific guidelines related to quality, safety and efficacy can be found at the EMA website. It also includes concept papers, draft guidelines and overview of comments received during the consultation on draft versions:


The EMA also publishes on its website procedural and technical guidance and document templates which are intended to provide technical and procedural advice to applicants for marketing authorisations for medicinal products coming within the scope of the centralised procedure, in particular:

- EMA pre-submission guidance for users of the centralised procedure
- EMA Procedural advice for users of the centralised procedure for generic/hybrid applications
- EMA Procedural advice for users of the Centralised Procedure for Similar Biological Medicinal Products applications
- EMA post-authorisation guidance for users of the centralised procedure
- Product information templates

References

- "Procedures for marketing authorisation", The Rules governing Medicinal Products in the European Community, Volume 2A, Notice to Applicants, Chapter 1
- Good pharmacovigilance practice guidelines (GVP)
52. Which European Directorate for the Quality of Medicines and HealthCare (EDQM) activities impact on the centralised procedure? \textit{Rev. Jul 10}

52.1. Introduction

The European Directorate for the Quality of Medicines & HealthCare (EDQM) is a Directorate of the Council of Europe. It was created in 1996.

The mission of the EDQM is to contribute to the basic human right of access to good quality medicines and healthcare, and to promote and protect human and animal health by:

- Establishing and providing official standards for the manufacture and quality control of medicines applicable in all the signatory states of the Convention for the Elaboration of a European Pharmacopoeia.

- Performing the evaluation of applications for Certificates of Suitability of the Monographs of the European Pharmacopoeia (CEPs) and related coordination of related inspections.

- Establishing the list of Standard Terms, which cover pharmaceutical forms, routes of administration and containers used for medicinal products for human and veterinary use.

- Co-ordinating activities performed by Official Medicines Control Laboratories network including annual sampling and testing programme for Centrally Authorised Products (CAPs) within the setting of a network.

- Co-coordinating activities for the elaboration of programmes and policies linking the quality of medicines to the quality and safety of their use, in the fields of pharmaceutical practice and care, risk prevention and management as regards counterfeiting of medicines, and the classification of medicines as regards their supply.

- Publishing and distributing all EDQM publications, including the European Pharmacopoeia.

The EDQM representatives participate as observers to the Agency's Quality Working Party (QWP) and Biologics Working Party (BWP) meetings, the GMP inspection services group meetings as well as HMPC meetings at the European Medicines Agency.

52.2. European Pharmacopoeia and its use for an application

Pharmacopoeias are collections of standardised specifications, so called monographs, which define the quality reference for pharmaceuticals.


The texts of the European Pharmacopoeia cover active substances, excipients, substances or preparations for pharmaceutical use of chemical, animal, human or herbal origin, homoeopathic preparations and homoeopathic stocks, antibiotics, as well as dosage forms and containers. The texts of the European Pharmacopoeia also apply to biologicals, blood and plasma derivatives, vaccines and radio-pharmaceutical preparations.

QWP and BWP are consulted during the preparation and the revision of monographs.
Additionally, chemical and biological reference material of the European Pharmacopoeia (Chemical Reference Substances and Biological Reference Preparations) to be used where relevant as reference standards for the quality control of medicinal products and their constituents are adopted by the European Pharmacopoeia and centrally supplied from the EDQM.

With respect to the quality part (chemical, pharmaceutical and biological) of the dossier, all monographs including general monographs and general chapters of the European Pharmacopoeia are applicable.

When test procedures and methods used for manufacturing and controlling the raw materials and active substances or the starting materials, excipients or finished medicinal products are described in the European Pharmacopoeia, the required description to be included in Module 3 shall be replaced by the appropriate detailed reference to the monograph(s) and general chapter(s).

52.3. **What is the scope of the Certification Procedure of the EDQM?**

The Certification Procedure is intended for substances for which a monograph (general monograph and/or specific monograph) has been adopted by the European Pharmacopoeia Commission. The procedure does not apply for direct gene products (proteins), products obtained from human tissues, vaccines and blood products and preparations.

Under the official procedure described in Resolution AP-CSP (07) 1 (adopted by the Public Health Committee (Partial Agreement), Council of Europe) and Directive 2001/83/EC and 2003/63/EC as amended of the European Union, manufacturers or suppliers of active substances or excipients (organic or inorganic, obtained by synthesis, extraction or fermentation), any product with transmissible spongiform encephalopathy (TSE) risk, or herbal products used in the production or preparation of pharmaceutical products can apply for a certificate of suitability (CEP) concerning:

- The evaluation of the suitability of the monograph for the control of the chemical purity and microbiological quality of their substance according to the corresponding specific monograph; or
- The evaluation of the reduction of Transmissible Spongiform Encephalopathy (TSE) risk, according to the general monograph; or,
- Both of the above; or,
- The evaluation of the suitability of the monograph for the control of herbal drugs and herbal drugs preparations.

A CEP can be used by the manufacturers of pharmaceutical products in their marketing authorisation applications to demonstrate the compliance of the substance used with the monographs of the European Pharmacopoeia as referred in Directive 2001/83/EC, as amended. As a result, the applicants are exempted of providing the concerned data in the relevant parts of Module 3 of the MAA, as deemed to be replaced by the CEP, except for some parts needed for the assessment of the medicinal product. For instance, in case of sterile substances, the applicant has to resubmit the data on the sterilisation of the substance to National Competent Authorities/Agency. Additionally the manufacturer should provide the applicant with the written assurance that the manufacturing process has not been modified since the granting of the certificate of suitability by the EDQM.

In case a new or updated Certificate of Suitability has been issued, the applicant should submit it through the relevant variation procedure.
This procedure is aimed at facilitating and simplifying exchanges between the partners to ensure that the quality of substances is guaranteed and that these substances comply with the European Pharmacopoeia, by issuing a so-called Certificate of Suitability (CEP or CEP for TSE).

CEPs are recognised by all signatory states of the European Pharmacopoeia Convention and by the European Union. There are also other countries which have also chosen to recognise them.

**Note on CEPs for biological substances of non-recombinant origin**

Following EDQM decision to exclude from the scope of the certification procedure the products classified as “other biological substances” by the CMD (h). Applicants are requested to submit full data on the Module 3 for new applications for Marketing Authorisation through the centralised procedure for medicinal products containing these biological substances. Existing certificates of suitability (CEPs) for these substances can be included in the dossiers but should not be used as replacement of the relevant data in the corresponding sections of Module 3.

The reasoning behind this decision is that for biologicals the characterisation and determination of the quality of these products requires not only a combination of physico-chemical and biological testing, but also extensive knowledge over the production process and its control.

The EDQM will therefore not accept any new application for a CEP for these biological substances.

**52.4. List of Standard Terms and its use**

The list of the Standard Terms was drawn up by the European Pharmacopoeia Commission for use in the marketing authorisation application and the product information (SPC, labelling, package leaflet). It has the double purpose of bringing information to the patient/user/prescriber and distinguishing the various presentations of a medicinal product. It should convey essential information on the properties and use of the particular medicinal product presentation.

The Standard Term concerns either the pharmaceutical form, route of administration or container. The pharmaceutical form standard term consists of a combination of the form in which a medicinal product is presented (form of presentation) and the form in which it is administered, including the physical form (form of administration). In special cases (e.g. identical products which may be distinguished only by reference to the container), the information about the immediate container can be included in the pharmaceutical form, e.g. “solution for injection in pre-filled syringes.

Moreover, due to the specificity of a medicinal product the complete characterisation of a pharmaceutical form may be constructed by using a combination of existing Standard Terms, e.g. “powder for solution for injection or infusion”.

The route of administration indicates the part of the body on which, through which, or into which the medicinal product is to be administered.

The container is the packaging immediately in contact with the medicinal product.

When the nature of the medicinal product is such that no existing Standard Term or combination of Standard Terms accurately describes the product presentation, a request for a new Standard Term will have to be made to the EDQM. The need for such a request should be identified by the applicant preferably during the EMA pre-submission meeting. The applicant should submit to the EMA the request for a new standard term, together with appropriate supportive documentation i.e. a detailed description of the pharmaceutical form and proposed new term, together with a justification for the new term including why any of the existing terms are not appropriate and a draft SPC. The request will be reviewed by the Quality Review of Documents and the Quality Working Party groups. The EMA will
subsequently forward the applicant’s request and the common EMA position to the EDQM for final decision.

For more information on Standard Terms please refer to: http://www.edqm.eu/site/page_590.php

References

- EMA website (Inspections section)
- EDQM & HealthCare website
- List of Standard terms
53. When do I have to submit an Environmental Risk Assessment (ERA)? **Rev. Oct 14**

In accordance with Article 8(3) (ca) and (g) of Directive 2001/83/EC, as amended, the evaluation of the potential environmental risks posed by medicinal products should be submitted, their environmental impact should be assessed, and on case-by-case basis, specific arrangements to limit the impact should be considered. In any event this impact should not constitute a criterion for refusal of a marketing authorisation for medicinal products for human use.

The environmental risk assessment (ERA) concerns the risks to the environment arising from the use, storage, and disposal of the medicinal product. Risks arising from the synthesis or manufacture of the product are under the remits of the national competent authorities.

The ERA follows a step-wise, two-phase procedure. The first phase (phase I) estimates the exposure of the environment to the drug substance by calculating the predicted environmental concentration (PEC). The PEC calculation applies to the aquatic compartment (PEC_{SURFACEWATER}). If the PEC_{SURFACEWATER} value is equal or above 0.01 μg/L, then a phase II environmental-fate and effect analysis should be performed.

More details are provided in the guideline on environmental risk assessment of medicinal products for human use and in the related "Questions and Answers on Guideline on the environmental risk assessment of medicinal products for human use" document.

An ERA is required for all new MAAs for a medicinal product through a centralised, mutual recognition, decentralised and national procedure including applications submitted under Article 10 of the mentioned directive.

The ERA, including the relevant study reports, should be provided in module 1.6 of the MAA together with the dated signature of the author, information on the author’s educational, training and occupational experience (curriculum vitae) and a statement of his or her relationship with the applicant.

In the case of medicinal products containing natural substances e.g. vitamins, electrolytes, amino acids, peptides, proteins, carbohydrates, lipids and of vaccines and herbal medicinal products, a justification for not submitting ERA studies should be provided in module 1.6.

In case of an existing marketing authorisation, a re-evaluation of the ERA should be submitted with the application for type II variations or for extension applications.

An ERA is not required for renewals or Type IA/IB variations.

Studies in the context of an ERA are expected to be assessed during the initial marketing authorisation or relevant post-marketing procedures (e.g. extension of indication, extension applications). In the exceptional case that ERA study results are provided stand-alone, they should be submitted as a type IB C.1.z variation as described in the Practical questions and answers to support the implementation of the variations guidelines in the centralised procedure.

**References**

- Directive 2001/83/EC as amended
- Guideline on the Environmental Risk Assessment of the medicinal products for human use
• EudraLex – Volume 2 – Pharmaceutical legislation: Notice to applicants and regulatory guidelines medicinal products for human use

• Questions and answers on the Guideline on the environmental risk assessment of medicinal products for human use

• Practical questions and answers to support the implementation of the variations guidelines in the centralised procedure
54. How are the ATC codes/INN applied within the Centralised Procedure? **New Mar 07**

54.1. **ATC codes**

The Anatomical Therapeutic Chemical (ATC) classification is a system in which medicinal products are divided into different groups according to the organ or system on which they act and their chemical, pharmacological and therapeutic properties. The medicinal products are classified in groups at five different levels.

The Applicant for a Marketing Authorisation should apply for an ATC Code using the application form on the WHO website. For information on data to be submitted together with the application form please refer to the WHO website (www.whocc.no).

Within the Centralised Procedure, the ATC code is used in the application form for a Marketing Authorisation (MAA) and in the Summary of Products Characteristics (SPC). The Applicant should bear in mind that, if an ATC code is not yet assigned to the Medicinal Product, no temporary code should be mentioned in the SPC and “Not yet assigned”, should appear in section 5.1 of the SPC. The proposed/temporary code should however be mentioned in the application form for a MAA, stating its status in brackets. If an ATC code has been assigned, it should be given in section 5.1 of the SPC without any spaces and without brackets (e.g. N02BE01).

When the Applicant receives the final ATC code from the WHO, if this happens before CHMP opinion, the EMA should be informed as soon as possible in writing with the appropriate proof of the change in status from WHO and the SPC should be amended accordingly. If the ATC code is obtained after opinion, the EMA should be informed and the SPC should be amended accordingly either as a Type IA Variation or at the occasion of another variation after the Commission Decision has been obtained. The same procedure applies, in case of a revision of a final ATC code by the WHO for medicinal products already authorised.

54.2. **INN**

An International Non-proprietary Name (INN) identifies a pharmaceutical substance or active pharmaceutical ingredient by a unique name that is globally recognised and is public property. The aim of the INN system has been to provide health professionals with a unique and universally available designated name to identify each pharmaceutical substance. To make INNs universally available they are formally placed by WHO in the public domain, hence their designation as “non-proprietary”.

The names, which are given the status of an INN are selected by the WHO on the advice of experts from the WHO Expert Advisory Panel on the International Pharmacopoeia and Pharmaceutical Preparations. The process of INN selection follows three main steps:

- A request/application is made by the manufacturer or inventor, using an 'INN request form' from WHO website (http://www.who.int)
- After a review of the request a proposed INN is selected and published for comments in WHO Drug Information
- After a time period for objection has lapsed, the name will obtain the status of a recommended INN and is published as such by the WHO if no objection has been raised
If applicants for Marketing Authorisation (MA) wish to apply for an INN, it is strongly recommended to liaise with WHO well in advance of MA submission, in order to obtain a recommended INN for their pharmaceutical substance as soon as possible and preferably no later than the CHMP opinion is obtained. Within the Centralised Procedure, the INN is used throughout the MA dossier. If a recommended INN is not available at submission, the proposed INN can be used in the application form and in the Product Information (PI). When the applicant receives the recommended INN from the WHO, if this happens before CHMP opinion, the EMA should be informed as soon as possible in writing with the appropriate proof of the change in status from WHO and the PI should be amended accordingly. If the INN is obtained after opinion, the EMA should be informed and the PI should be amended accordingly either as a Type IA Variation or at the occasion of another variation after the Commission Decision has been obtained.

For certain biologicals, because of their complexity, general rules for INN are not easily formulated. Some of these substances may have descriptive names assigned by other institutions. These names may not be suitable as INNs. Some nomenclature schemes for groups of biological compounds are provided in the WHO guideline.

For vaccines the INN is not applicable and in these cases either the pharmacopoeial or common name of the antigens should be used.

In the absence of INN, the common name or scientific name of the pharmaceutical substance should be used.

**References**

- WHO Collaborative Centre for Drug Statistic Methodology website
- WHO "Guidelines on the Use of International Non-proprietary Names (INNs) for Pharmaceutical Substances" (WHO/PHARM S/NOM 1570);
- ‘Guideline on Summary of Product Characteristics (October 2005)’ the Rules governing Medicinal Products in the European Community, Notice to Applicants, Volume 2C;
- WHO “International Non-proprietary Names (INN) For Biological and Biotechnological Substances”
55. Can I apply for Design Space or Process Analytical Technology (PAT) in my application? New Mar 07

The ICH Q8 (Pharmaceutical Development) introduces the notion of Design Space, defined as the multidimensional combination and interaction of input variables (e.g. material attributes) and process parameters that have been demonstrated to provide assurance of quality. The Design Space is proposed by the applicant as part of the MAA and thus is subject to assessment.

Additionally the establishment of a robust Design Space is in line with new approaches on quality which focus on building quality into the medicinal product by design (the so-called QbD concept)

PAT is defined as a system for analysing and controlling manufacturing through timely measurements (i.e. during processing) of critical quality and performance attributes of raw and in-process materials and processes with the goal of ensuring final product quality.

PAT is a tool that allows enhanced control of the manufacturing process, can improve process understanding and so facilitates building quality into products and the development of a Design Space. ICH Q9 (Quality Risk Management) provides an approach and a selection of tools which can be used to manage risks associated with these processes.

The main PAT tools are:

- multivariate data acquisition and analysis;
- modern process analysers or process analytical chemistry tools;

The introduction of the PAT system can bring a number of advantages:

- Possibilities to introduce "real time release";
- Reduction of cycle times;
- Improved product quality;
- Possibilities for more efficient and effective control of some changes;

The introduction of PAT system can be applied to new or existing authorised medicinal products.

55.1. When to inform the EMA of the introduction of PAT or Design Space approaches in my application

- Where Design Space concepts or PAT approaches are used, Marketing Authorisation applicants should indicate this in their letter of intent. It is of interest for the Agency and CHMP to be aware of their use so it can be taken into account in the appointment of (Co)-Rapporteurship, as particular expertise from (Co)-Rapporteurs may be needed.
- In addition, when requesting a pre-submission meeting, the applicant should identify it in the relevant question of the pre-submission request form.

55.2. The role of the EMA PAT team

The EMA Process Analytical Technology Team is a forum for dialogue and understanding between Quality and Biologics Working Parties and Ad-Hoc Group of GMP Inspection Services to prepare a harmonised approach in Europe on assessment of applications and inspections of products/systems/facilities for Process Analytical Technology, including quality by design principles and
manufacturing science in the context of PAT. The PAT team may be consulted through QWP or BWP during the assessment of a centralised marketing authorisation application. Applicants using a PAT approach are encouraged to look at the PAT-related guidance and questions and answers document provided on the EMA website. If there are still questions or issues which are not addressed through those documents, applicants could take the opportunity to contact the EMA PAT team at early stage of pharmaceutical development. It should be noted that the PAT team only provides informal and non-binding advice which does not substitute for Scientific Advice/Protocol Assistance.

55.3. Presentation of PAT-related data in the application

When an application for, or variation to, a marketing authorisation is submitted, supporting documentation should be provided in accordance with CTD requirements (Module 3). In addition, the Expert Report provided in Module 2 (Quality Overall Summary) should include a critique highlighting the positive and negative aspects of the Design Space or PAT approach. For more information see: Reflection Paper - Chemical, pharmaceutical and biological information to be included in dossiers when Process Analytical Technology (PAT) is employed. Applicants should note that submission of applications that include Design Space or PAT aspects could result in a specific product related inspection at the manufacturing site.

References

- EMEA website, Inspections section
- ICH (International Conference on Harmonization) Q8, Pharmaceutical Development
- ICH Q9, Quality Risk Management
- The Rules Governing Medicinal Products in the European Union, Volume 2A, Chapter 4 on “Centralised Procedure”
- Reflection Paper: Chemical, pharmaceutical and biological information to be included in dossiers when Process Analytical Technology (PAT) is employed (EMEA/INS/277260/2005)
56. Could my application qualify for a conditional marketing authorisation? *New July 07*

56.1. Criteria and general provisions

For certain categories of medicinal products, in order to meet unmet medical needs of patients and in the interest of public health, it may be necessary to grant marketing authorisations on the basis of less complete data than is normally required. In such cases, it is possible for the CHMP to recommend the granting of a marketing authorisation subject to certain specific obligations to be reviewed annually ("conditional marketing authorisation").

This may apply to medicinal products for human use that fall under Article 3(1) and (2) of Regulation (EC) No 726/2004 and belong to one of the following categories:

- medicinal products which aim at the treatment, the prevention or the medical diagnosis of seriously debilitating diseases or life-threatening diseases;
- medicinal products to be used in emergency situations, in response to public threats duly recognised either by the WHO or by the Community in the framework of Decision (EC) No 2119/98;
- medicinal products designated as orphan medicinal products in accordance with Article 3 of Regulation (EC) No 141/2000.

A conditional marketing authorisation may be granted where the CHMP finds that, although comprehensive clinical data referring to the safety and efficacy of the medicinal product have not been supplied, all the following requirements are met:

- the risk-benefit balance of the medicinal product, as defined in Article 1(28a) of Directive 2001/83/Ec, is positive;
- it is likely that the applicant will be in a position to provide the comprehensive clinical data;
- unmet medical needs will be fulfilled;
- the benefit to public health of the immediate availability on the market of the medicinal product concerned outweighs the risk inherent in the fact that additional data are still required.


The granting of a conditional marketing authorisation should be restricted to situations where only the clinical part of the application dossier is not yet fully complete. Incomplete non-clinical and/or quality data should only be accepted if duly justified and only in the case of a product intended to be used in emergency situations, in response to public health threats.

Conditional marketing authorisations will be valid for one year, on a renewable basis. The holder will be required to complete ongoing studies or to conduct new studies with a view to confirming that the risk-benefit balance is positive. In addition, specific obligations may be imposed in relation to the collection of pharmacovigilance data.

The granting of a conditional marketing authorisation will allow medicines to reach patients with unmet medical needs earlier than might otherwise be the case, and will ensure that additional data on a product are generated, submitted, assessed and acted upon.
56.2. Prior to submission

Applicants for a potential conditional marketing authorisation may request CHMP scientific advice or protocol assistance, as applicable, on whether a specific medicinal product being developed for a specific therapeutic indication falls within one of the categories set out in Article 2 and fulfils the requirement laid down in Article 4(1)(c) (“unmet medical needs will be fulfilled”) of Regulation (EC) No 507/2006. In addition, the intention to request a conditional marketing authorisation and any practical or procedural issues with regards to a potential request for conditional marketing authorisation should be addressed at the pre-submission meeting.

56.3. Timing of the submission and documentation to be supplied

At least seven months before submission, applicants should notify the EMA of their intention to submit an application and include a statement on the intention to request a conditional marketing authorisation (in accordance with Article 14(7) of the Regulation).

The applicant may present a request for a conditional marketing authorisation at the time of the application for marketing authorisation.

If the applicant considers that the grounds for a conditional marketing authorisation apply, the applicant should tick the box 1.5.1 of the application form and include its justification in module 1.5.5. Such justification should show that the medicinal product falls within the scope of the conditional marketing authorisation Regulation (Article 2) and that the requirements for conditional marketing authorisation are fulfilled (Article 4), together with the applicant's proposal for completion of ongoing or new studies, or the collection of pharmacovigilance data. The request may cross-refer to specific parts of the application.

Upon receipt of a valid application containing a request for conditional marketing authorisation, the EMA will inform the Commission.

For further guidance on the criteria for conditional marketing authorisations, justifications to be provided and the procedure to be followed, reference is made to the draft guidance document published on the EMA website (Draft Guideline on the scientific application and the practical arrangements necessary to implement Commission Regulation (EC) No 507/2006 on the conditional marketing authorisation for medicinal products for human use falling within the scope of Regulation (EC) No 726/2004).

56.4. CHMP assessment of a request for conditional marketing authorisation

The Rapporteur, Co-Rapporteur and the other CHMP members will assess the justification/data submitted for a Conditional Marketing Authorisation as part of the overall assessment of the benefit/risk of the application. The assessment of the justification will be reflected in the relevant assessment reports and in the final CHMP assessment report.

A conditional marketing authorisation may be requested by the applicant or proposed by the CHMP. Therefore, during the scientific assessment, after having consulted with the applicant, the CHMP may also propose a conditional marketing authorisation. Normally, the proposal and explanatory reasons will be given to the applicant in the day 120 list of questions, or exceptionally later, in the day 150 joint assessment report and day 180 list of outstanding issues. The reasons for proposing a conditional marketing authorisation will also be detailed in the relevant assessment reports and in the CHMP assessment report.
Upon granting of a conditional marketing authorisation, the specific obligations and the timeframe for their completion will be clearly specified in the conditional marketing authorisation (Annex II.C to the Commission Decision), and will be made publicly available by the Agency as part of the EPAR.

56.5. **Information included in the summary of product characteristics and package leaflet**

In order to provide clear information to patients and healthcare professionals on the conditional nature of the authorisations, the summary of product characteristics and package leaflet will mention that a conditional marketing authorisation has been granted subject to certain specific obligations to be reviewed annually.

**Differences between Conditional Marketing Authorisation and Marketing Authorisation under Exceptional Circumstances**

<table>
<thead>
<tr>
<th>Conditional Marketing Authorisation</th>
<th>Marketing Authorisation under Exceptional Circumstances</th>
</tr>
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<tbody>
<tr>
<td>Demonstrate positive benefit-risk balance, based on scientific data, pending confirmation</td>
<td>Comprehensive data cannot be provided (specific reasons foreseen in the legislation)</td>
</tr>
<tr>
<td>Authorisation valid for one year, on a renewable basis</td>
<td>Reviewed annually to reassess the risk-benefit balance, in an annual re-assessment procedure</td>
</tr>
<tr>
<td>Once the pending studies are provided, it can become a &quot;normal&quot; marketing authorisation</td>
<td>Will normally not lead to the completion of a full dossier and become a &quot;normal&quot; marketing authorisation</td>
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Conditional Marketing Authorisations are distinct from marketing authorisations granted in exceptional circumstances in accordance with Article 14(8) of Regulation (EC) No 726/2004. In the case of the conditional marketing authorisation, an authorisation is granted before all data are available. The authorisation is not intended, however, to remain conditional indefinitely. Rather, once the missing data are provided, it should be possible to replace it with a marketing authorisation which is not conditional, that is to say, which is not subject to specific obligations. In contrast, it will normally never be possible to assemble a full dossier in respect of a marketing authorisation granted in exceptional circumstances.

**References**

- Regulation (EC) No 726/2004
- Notice To Applicants Volume 2A Chapter 4 (Centralised Procedure)
57. How can I get support from the EMA regarding emerging therapies and technologies? Role of the Innovation Task Force (ITF) New July 07

In order to provide support to medicines innovation in EU, the EMA has established an internal multidisciplinary group including scientific, regulatory and legal competences, creating a forum for early dialogue with applicants. ITF members are scientific and legal administrators appointed from different sectors of Human Units, Directorate and Inspection Services. To fulfil its task the ITF may consult as appropriate EMA scientific Committees and Working Parties or individual experts.

The scope of the ITF activities encompasses emerging therapies (i.e. gene therapy, cell therapy and engineered tissues), emerging technologies (i.e. new development strategies, new manufacturing approaches) and borderline therapeutics (i.e. combination of pharmaceuticals and devices) for which there is no established EMA scientific, legal and regulatory experience.

Support available to applicants include:

- General queries relating to Emerging Therapies and Technologies
- Briefing meetings aiming to provide an early guidance and information, in liaison when needed with relevant EMA scientific committees or Working Parties. Additionally briefing meetings complement and reinforce existing formal regulatory procedures e.g. scientific advice
- Requests for regulatory advice on the eligibility to EMA procedures e.g. marketing authorisation, scientific advice, consultation on ancillary medicinal and blood and plasma derivatives in medical devices

For more information on Innovation Task Force and on how to request a briefing meeting or Regulatory Advice refer to the EMA Emerging Therapies and Technologies website. The request forms for Briefing Meetings and Regulatory Advice should be submitted electronically to ITFsecretariat@ema.europa.eu taking into account the dates for submission.

References

- EMA Emerging Therapies and Technologies website
- Mandate of the EMEA Innovation Task Force (ITF)
58. Are there special incentives or assistance for applicants which are Small or Medium-Sized Enterprises (SMEs)? Rev. May 11

Incentives and assistance are available from EMA for SMEs, which focus on reducing financial and administrative entry hurdles for SMEs in pre-marketing authorisation procedures such as scientific advice, the application for marketing authorisation and inspections.

These include:

- Administrative and procedural assistance from the SME Office at the Agency.
- Fee reductions for scientific advice, scientific services and inspections (90% fee reduction).
- Fee exemptions for certain administrative services (excluding parallel distribution).
- Deferral of the fee payable for an application for marketing authorisation or related inspection.
- Conditional fee exemption where scientific advice is followed and a marketing authorisation application is not successful.
- Certification of quality/non-clinical data for advanced therapy medicinal products (ATMPs) intended for human use.
- Assistance with translations of the product information documents submitted in a centralised application for marketing authorisation.
- Waiver of the MedDRA licensing fee when registering with EudraVigilance7.

In determining which companies are eligible for SME incentives, the EMA applies the EU-definition of micro, small and medium-sized enterprises provided in Commission Recommendation 2003/361/EC.

Companies are classified according to their size (micro, small or medium):

- Micro enterprises employ less than 10 persons and have an annual turnover or balance sheet total not exceeding € 2 million;
- Small enterprises have fewer than 50 employees and an annual turnover or balance sheet total of not more than € 10 million;
- Medium enterprises have less than 250 employees and an annual turnover of not more than € 50 million or an annual balance sheet total of not more than € 43 million

and according to their category (autonomous, partner or linked).

Depending on the category in which the enterprise fits, some or all of the headcount and financial data from other partner or linked enterprises may need to be counted when calculating whether the SME criteria are met.

Further information on the definition of an SME is available in “The new SME definition - User guide and model declaration”, published by the European Commission.

A declaration of SME status (form available on EMA website on SMEs) should be submitted to the SME Office prior to requesting financial or administrative assistance from the agency.

7 The MedDRA fee waiver applies to micro and small enterprises only, not to medium-sized companies.
SME Office

The ‘SME office’ has been set up within the agency to address the particular needs of smaller companies. The office aims to facilitate communication with SMEs through dedicated personnel within the agency who will respond to practical or procedural enquiries, monitor applications, and organise workshops and training sessions for SMEs.

Fee Reductions/Deferrals

SME applicants wishing to request a fee reduction and/or deferral should address a letter of intent to the SME Office (see below) of the EMA. It should be noted that fee reductions and deferrals can only be considered once the applicant has been assigned SME status by the EMA and are subject to the SME status remaining valid at the time that their application or request is validated by the Agency. Fee reductions and fee deferrals will not be granted retrospectively. Further information on fee reductions/deferrals is available in the document "Fee reductions/deferrals for micro, small and medium-sized enterprises (SMEs)".

Translation assistance

Because translating product information into all EU languages represents a considerable financial and administrative burden to SMEs entering the EU market, the EMA will provide for translation of product information documents (summary of product characteristics, conditions of the marketing authorisation, label and package leaflet) required for the grant of an EU marketing authorisation. The applicant remains responsible for provision of the Norwegian and Icelandic translations according to the normal timelines and for the maintenance of all translations in the post-authorisation phase.

Due to the timelines required to translate the product information, the Agency will initiate translations through the Centre for Translation (CdT) in Luxembourg prior to CHMP/CVMP opinion (normally around day 180 of the procedure). These translations will then be checked through the national competent authorities in the Member States (see also "QRD product information - Tools used by the EMA to facilitate the streamlining of the European Decision Making process"). To be eligible for translation assistance the applicant’s SME status must be valid at the time the translations are initiated.

Companies wishing to benefit from SME incentives should visit the SME Office section of the EMA website first. This section provides useful information on how to request SME status, and provides a link to useful information sources (e.g. the User Guide for Micro, Small and Medium-sized Enterprises (SMEs) on the administrative and procedural aspects of the provisions, laid down in Regulation (EC) No 726/2004, that are of particular relevance to SMEs).

For further information or requests please contact:

SME Office
Tel.: (44 20) 7418 8575
E-Mail: smeoffice@ema.europa.eu

References

- Commission Regulation (EC) No 2049/2005
- Commission Recommendation 2003/361/EC
- The new SME definition - User guide and model declaration
- Declaration on the qualification of an enterprise as a micro, small or medium-sized enterprise (SME) (EMEA/366649/2005)
• User Guide for Micro, small and Medium-sized Enterprises (SMEs) on the administrative and procedural aspects of the provisions, laid down in Regulation (EC) No 726/2004, that are of particular relevance to SMEs

• Fee reductions/deferrals for micro, small and medium-sized enterprises (SMEs) (EMEA/366526/2005)
Regulation (EC) No 1901/2006 (the 'Paediatric Regulation') lays down obligations, rewards and incentives for the development and placing on the market of medicines for use in children. The Paediatric Regulation places some obligations for the applicant when developing a new medicinal product, in order to ensure that medicines to treat children are subject to ethical research of high quality and are appropriately authorised for use in children, and to improve collection of information on the use of medicines in the various subsets of the paediatric population. The paediatric population is defined as the population between birth and the age of 18 years (meaning up to but not including 18-years).

As set out in Article 7 of the Paediatric Regulation, applications concerning a medicinal product “not authorised in the Community” on 26 July 2008 must include one of the following documents/data in order to be considered ‘valid’:

- The results of all studies performed and details of all information collected in compliance with an agreed Paediatric Investigation Plan (PIP).
  
  This means that the application will have to include the PIP decision but also the results in accordance with the agreed PIP.

- A decision of the EMA on a PIP including the granting of a deferral.
  
  This means that the application will have to include the PIP decision including the deferral granted and if applicable, any completed studies.

- A decision of the EMA granting a product-specific waiver.

- A decision of the EMA granting a class waiver together with the EMA confirmation letter of applicability if requested by the MAH.

Where results of paediatric studies are submitted, applicants should include in the clinical overview a rationale supporting the proposed changes to the Product Information. In particular, if the PIP is completed and the results of all studies are available, the applicant should explicitly discuss why the generated data support or do not support the intended paediatric indication(s) stated in the PIP.

Inclusion of the results of all studies performed in compliance with an agreed Paediatric Investigation Plan requirement in the Product Information is a prerequisite for benefiting from the paediatric reward (Article 36(1) of Regulation (EC) No 1901/2006).

The Global Marketing Authorisation (GMA) concept together with the notion of “same marketing authorisation holder” should be used to determine whether an application concerns a "medicinal product for human use which is authorised or not in the Community". Further information can be found in the Procedural Advice document on "applications for PIPs, Waivers and Modifications" which is available on the EMA website under ‘Special Topics - Medicines for children’.

However, the following types of application are exempted from the application of the above requirements:

- Generic medicinal products (Art 10(1) of Directive 2001/83/EC)
- Hybrid medicinal products (Art 10(3) of Directive 2001/83/EC)
- Similar biological medicinal products (Art 10(4) of Directive 2001/83/EC)
• Medicinal products containing active substance(s) of well-established medicinal use (Art 10a of Directive 2001/83/EC)

Furthermore, when planning submission of their marketing authorisation application, the applicant has to take into account also the need for a "PIP compliance check” to be done.

Such compliance check consists of verifying that the fulfilments of the measures as mentioned in the PIP decision including the timelines for the conduct of the studies or collection of the data are fulfilled. The compliance check procedure is explained in the document “Questions and answers on the procedure of paediatric-investigation-plan compliance verification at the European Medicines Agency”. Applicants are strongly recommended to apply for the compliance check before submission of the marketing authorisation application to not delay the validation phase.

Further details on the format timing and content of PIP or waiver applications as well as on the compliance check can be found in the Commission guideline. In addition, deadlines for submission of PIP or Waiver applications and application templates as well as “Procedural Advice documents respectively regarding applications for PIPs, Waivers and Modifications” and “validation of new MAA, Variation/Extension applications and compliance check with an agreed PIP” are available on the EMA website in section “Special Topics - Medicines for children”.

References

• Regulation (EC) No 1901/2006
• Commission Guideline on “The format and content of applications for agreement or modification of a paediatric investigation plan and request for waivers or deferrals and concerning the operation of the compliance check and on criteria for assessing significant studies”
• Procedural Advice document related to “Paediatric investigation plans (PIPs), waivers and modifications”
• Questions and answers on the procedure of paediatric investigation plan compliance verification at the European Medicines Agency
• EMA website, section “Special Topics - Medicines for children”
60. Can I submit an application for a Paediatric Use Marketing Authorisation (PUMA)? **Rev. Feb 12**

### 60.1. Introduction

According to Article 30 of Regulation (EC) No 1901/2006 ("The Paediatric Regulation"), the paediatric use marketing authorisation (PUMA) is a dedicated marketing authorisation for medicinal products indicated exclusively for use in the paediatric population, or subsets thereof, with, if necessary, an age-appropriate formulation. It has been designed to promote paediatric development of already authorised products which are no longer covered by a supplementary protection certificate (SPC) or a patent qualifying for a SPC.

### 60.2. Eligibility to the centralised procedure

A PUMA application remaining outside the mandatory scope of Article 3(2)(a) of Regulation (EC) No 726/2004 has an ‘automatic access’ to the centralised procedure (Article 31 of the Paediatric Regulation) if the applicant chooses this route of registration.

Before a PUMA application is submitted for the centralised procedure, an eligibility confirmation must be requested by the applicant by submitting a Pre-submission request form (Eligibility) to CPeligibility@ema.europa.eu. For more information on the eligibility request, please refer to the European Medicines Agency pre-submission procedural advice for users of the centralised procedure.

### 60.3. Content of a PUMA application

The same range of supporting documentation should be provided as for other marketing authorisation applications through a combination of new data and/or existing data. Depending on the legal basis of the application, submission of literature and/or cross-reference to the dossier of another medicinal product may be used. In particular, cross-reference to the data contained in the dossier of an authorised medicinal product is possible if the relevant data protection has expired. For further information, please refer to the pre-submission Procedural advice for users of the centralised procedure for generic/hybrid applications.

A PUMA application has to contain the results of all studies performed and details of all information collected in compliance with an agreed Paediatric Investigation Plan (PIP). The corresponding EMA decision as well as the PDCO opinion on compliance or the applicant’s compliance report must be provided in Module 1.10 (please refer to the pre-submission procedural advice for users of the centralised procedure – Q. 48. Do I need to address any paediatric requirements in my application?).

Further details on the submission of a PIP are available on the EMA website in section "Special topics – Medicines for children”.

As per Article 34 of the Paediatric Regulation, applicants are required to detail in a risk-management plan submitted with their PUMA application the measures to ensure the follow-up of efficacy and of possible adverse reactions to the paediatric use of the medicinal product.

### 60.4. Incentives for PUMA

PUMA applications have an ‘automatic access’ to the centralised procedure (Article 31 of the Paediatric Regulation).
PUMA benefits from the 8+2 year period of data and market protection (Article 38 of the Paediatric Regulation).

A medicinal product for which a PUMA has been granted may retain the name of another medicinal product containing the same active substance for which the same holder has been granted an authorisation for use in adults (Article 30(4) of the Paediatric Regulation).

PUMA applications submitted under the centralised procedure benefit from a partial exemption from the payment of fees laid down in the Regulation (EC) No 297/95. This partial exemption applies to the submission of the PUMA application and some of the post-authorisation activities for 1 year as of the date of granting of the PUMA. Please refer to the Regulation (EC) No 297/95 and the Explanatory note on fees payable to the European Medicines Agency.

Further information on PUMA and paediatric requirements related to a PUMA application are available on the EMA website in section “Special topics – Medicines for children”.

References

- Articles 2 and 30 of Regulation (EC) No 1901/2006
- Procedural Advice document related to Paediatric investigation plans (PIPs), waivers and modifications
- Procedural advice for validation of new marketing authorisation application, extension/variation application and compliance check with an agreed PIP
- Commission Guideline on “The format and content of applications for agreement or modification of a paediatric investigation plan and request for waivers or deferrals and concerning the operation of the compliance check and on criteria for assessing significant studies”
- Explanatory note on fees payable to the European Medicines Agency.
- EMA website, section “Special Topics - Medicines for children - PUMA”
61. What is the Community Plasma Master File certification system? *New Mar 09*

The concept of 'Plasma Master File' (PMF) was introduced with the Commission Directive 2003/63/EC in June 2003 amending Directive 2001/83/EC.

The PMF is a compilation of all required scientific data on the quality and safety of human plasma relevant to medicines, medical devices and investigational products which use human plasma in their manufacture. These data cover all aspects of the use of plasma, from collection to plasma pool.

The PMF is a stand-alone document which is separate from the application dossier for a Marketing Authorisation for the medicinal product concerned.

The PMF certification is an optional procedure that follows a similar system to the Marketing Authorisation evaluation procedure (the 'centralised procedure') at the EMA.

Following the satisfactory outcome of an evaluation, the EMA issues a PMF Certificate of compliance with Community legislation, which is valid throughout the European Community.

A Marketing Authorisation (MA) or a Marketing Authorisation Application (MAA) may refer to one or more PMFs or respective certificates. Once the Applicant chooses to use the Community PMF certification system all variations to the corresponding plasma for all the linked MAs will have to be submitted through the same certification system.

The competent authority that will grant or has granted a MA shall take into account the certification, re-certification or variation of the PMF on the concerned medicinal product(s).

For medicinal products that have been evaluated by the EMA through the Centralised Procedure and authorised by the European Commission, the public can find a summary of the quality and safety of the plasma in the product's European Public Assessment Report (EPAR).

For detailed information related to the Plasma Master File certification, please consult the Plasma Master File webpage.
62. What is the Community Vaccine Antigen Master File certification system? *New Mar 09*

The concept of “Vaccine Antigen Master File” (VAMF) was introduced with the Commission Directive 2003/63/EC in June 2003 amending Directive 2001/83/EC.

A VAMF contains all relevant information of biological, pharmaceutical and chemical nature for one given vaccine antigen, which is common to several vaccines from the same marketing authorisation (MA) applicant or marketing authorisation holder (MAH).

The use of the VAMF certification system is optional and the VAMF is a stand-alone part of the marketing authorisation application dossier (MAA) for a vaccine.

The VAMF certification consists of a centralised assessment of the VAMF application dossier submitted by the MA Applicant/MAH, which results in a certificate of compliance to Community legislation, issued by the EMA. This certificate is valid throughout the European Community.

A Marketing Authorisation (MA) or a Marketing Authorisation Application (MAA) may contain one or more VAMF certificates and respective VAMF data. If, when submitting a new MAA, the MA Applicant decides to opt for vaccine antigen master files, the VAMFs must be submitted for all vaccine antigens in the respective MAA.

As a rule, one VAMF should be submitted per vaccine antigen. In the case of a group of antigens aimed at preventing a single infectious disease a VAMF should be submitted for each antigen in the group.

A VAMF application can only be submitted to the EMA for antigens that form part of at least one MA or MAA, which has been, or will be evaluated via a Community procedure (Mutual Recognition (MR), Decentralised Procedure (DCP) or Centralised Procedure (CP)).

Once the Applicant chooses to use the Community VAMF certification system, all variations to the corresponding MAs will have to be submitted through the same certification system.

The competent authority that will grant or has granted a MA shall take into account the certification, re-certification or variation of the VAMF on the concerned medicinal product(s).

For detailed information related to the Vaccine Antigen Master File certification, please consult the Vaccine Antigen Master File webpage.
63. What is Eudravigilance? How will it apply to my Marketing Authorisation? **New Mar 09**

The reporting of suspected serious adverse reactions is defined in the Community legislation. This process involves healthcare professionals, the EMA, national Competent Authorities (NCAs) and MAHs and is applicable to all medicinal products authorised in the EEA. The reporting includes suspected serious adverse reactions occurring both within and outside the EEA.

With effect from 20th November 2005, the electronic reporting of suspected serious adverse reactions, save in exceptional circumstances, has become mandatory.

EudraVigilance is a data processing network and management system, which is used for reporting and evaluating suspected adverse reactions during the development and following the marketing authorisation of medicinal products in the European Economic Area (EEA).

EudraVigilance supports:

- Electronic exchange of suspected adverse reaction reports (referred to as Individual Case Safety Reports) between the European Medicines Agency (EMA), national Competent Authorities, marketing authorisation holders, and sponsors of clinical trials in the EEA.
- Early detection of possible safety signals associated with medicinal products for human use.
- Continuous monitoring and evaluation of potential safety issues in relation to reported adverse reactions.

EudraVigilance is also one of the main pillars of the European Risk Management Strategy and facilitates the process of risk management at several levels including risk detection, risk assessment, risk minimisation and risk communication.

Practical and detailed guidance on the collection, verification and presentation of adverse reaction reports arising from clinical trials on medicinal products for human use can be found via this link: http://ec.europa.eu/health/documents/eudarex/index_en.htm.

A marketing authorisation holder should prepare for the electronic reporting of suspected adverse reactions to the EMA as follows:

- Provide the EMA with a written plan on how the company is going to implement the electronic transmission of ICSRs to the Agency and national Competent Authorities in the EEA. Please address your plan to the attention of Ms Sabine Brosch (sabine.brosch@ema.europa.eu).
- Follow the detailed instructions outlined in “10 Steps to Implementation”, where the procedure for the initiation of the electronic transmission of ICSRs is described.
- Register with EudraVigilance. Please note that a MedDRA license is required for electronic reporting of ICSRs. For further information on the EudraVigilance MedDRA licensing Policy, please refer to MedDRA licensing Policy in this website.
- Provide the required information for the EudraVigilance Medicinal Product Dictionary.

For detailed information related to EudraVigilance, please consult the EudraVigilance webpage http://eudravigilance.ema.europa.eu/highres.htm or contact eudravigilance@ema.europa.eu.
64. What do I have to consider regarding the MA of my centrally authorised medicinal product in Norway, Iceland and Liechtenstein? Rev. Jan 13

Norway, Iceland and Liechtenstein have, through the European Economic Area agreement, adopted the complete Community acquis on medicinal products, and are consequently parties to the centralised procedure. However, legally binding acts from the Community, e.g. Commission Decisions, do not directly confer rights and obligations in Norway and Iceland, but first have to be transposed into legally binding acts in these states, except for Liechtenstein, where they are directly applicable, according to national legislation. According to Decision No. 74/1999 of the EEA Joint Committee, when decisions on approval of medicinal products are taken by the Community, Norway, Iceland and Liechtenstein will take corresponding decisions on the basis of the relevant acts.

The EEA Joint Committee Decision No. 74/1999 on the extension of the Marketing Authorisation Procedures for medicinal products to Norway, Iceland and Liechtenstein entered into force on 1 January 2000.

Specificities for Norway and Iceland

Within the Linguistic Review Process of Product Information in the Centralised Procedure – Human EMEA/5542/02, applicants are required to electronically provide the EMA translations of the agreed product information in all EU languages, including Icelandic and Norwegian, after the adoption of the CHMP EN opinion for review. The Norwegian and Icelandic texts will be checked by the respective Agencies.

Once a Commission Decision is issued, the European Commission publishes the Commission Decision with Annexes in all EU languages on its website. Subsequently, the Norwegian and Icelandic PI texts are published on the EMA’s website.

Norway

The Norwegian authorities will grant a corresponding national authorisation within 30 days following the date of the Commission Decision after receiving final product information in Norwegian from the MAH. Provision of specimens and mock-ups to Norway is not required.

For information regarding the handling of variations in Norway for centralised medicinal products please consult the Norwegian Medicines Agency website:

http://www.legemiddelverket.no/English/regulatory-affairs/variations/Sider/Centralised-procedure---variations.aspx

Please contact:

Norwegian Medicines Agency
P.O. Box 63, Kalbakken
N-0901 Oslo
Norway
Tel.: +47 22 89 77 00
Fax: +47 22 89 77 99
E-mail: pi@noma.no.
**Iceland**

The Icelandic authorities will grant a corresponding national authorisation within 30 days following the date of the Commission Decision. Provision of specimens to Iceland is not required.

For information regarding the handling of variations in Iceland for centralised medicinal products please consult the Icelandic Medicines Agency website: http://www.imca.is/IMCA/News/nr/1120.

At least one month before marketing, the applicant has to provide the Icelandic authorities directly with mock-ups for all product presentations that are intended to be marketed in Iceland. Mock-ups should be sent by e-mail to mockups@ima.is. See also http://www.imca.is/imca/news/nr/1263.

**References**

- Regulation (EC) No 726/2004
- Directive 2001/83/EC
- The Rules governing Medicinal Products in the European Union, Notice to Applicants, Volume 2A, Chapter 1 - Marketing Authorisation, Chapter 4 – Centralised Procedure, Chapter 6 – Procedures for MA , Chapter 7 – General Information
- Decision of the EEA Joint Committee No 74/1999
- The linguistic review process of product information in the centralised procedure – human
- The Revised Checking Process of Mock-Ups and Specimens of outer/immediate labelling and package leaflets of human medicinal products in the Centralised Procedure
65. Is my product subject to batch release by an Official Medicines Control Laboratory and if so, what should I consider? *New May 10*

Live vaccines, immunological medicinal products and medicinal products derived from human blood or plasma may be subject to batch release by a Member State laboratory or Official Medicines Control Laboratory (OMCL).

The OMCL supports the regulatory authorities and the national Inspection Services in ensuring the quality of medicinal products on the market by independent re-testing based on the legal requirements.

The European Medicines Agency and EDQM (European Directorate for the Quality of Medicines and Health Care) on behalf of the OCABR (Official Control Authority Batch Release) Network have been working on a common strategy with the aim of ensuring that the technical expertise of the OMCLs is taken into account in the development and assessment of testing methodologies for vaccines and plasma derived blood products that may be subject to OMCL batch release.

The input of the OMCLs is particularly important for products that include a novel quality control method or where there are known difficulties with a particular assay.

It is therefore strongly recommended for an applicant to enter into early collaboration with the OMCL. This collaboration should ideally begin at least one year before submission of the Marketing Authorisation Application, in order to allow for exchange of information between the OMCL and the Applicant which should be considered in the development of testing methodology.

For this purpose, Applicants are advised to consult the following site on the EDQM webpage for a contact list of OMCLs in the EU carrying out OCABR.

The information on the chosen OMCL by the Applicant will be recorded in the EMA pre-submission meeting and be passed onto the CHMP.

The European Medicines Agency will inform EDQM of any upcoming start of an authorisation procedure with official batch release.

**References**

- Directive 2001/83/EC, Article 114
66. How and when can I withdraw my application? *NEW May 2015*

If the applicant wishes to withdraw their application for marketing authorisation during assessment, it should inform the EMA Procedure Manager by providing a withdrawal letter stating that the applicant withdraws their application, specifying whether in full or partly (e.g. only a certain strength), and indicating reasons for the withdrawal.

The Letter should be signed by the authorised representative of the applicant. Applicants are advised that letters for withdrawal of marketing authorisation applications (in case of a full withdrawal) will be published on the EMA’s website (after redaction of protected personal data).

Applicants can address the withdrawal request to the EMA at any point during the assessment (from validation of the application up until adoption of the final CHMP Opinion).

Of note, the Agency will charge the fee for the application at the start of the procedure, irrespective of its outcome (positive, negative or partial/full withdrawal) and publish information on withdrawn applications.

**References**

- Procedural advice on publication of information on withdrawals of applications related to the marketing authorisation of human medicinal products