

Unique Device Identifier (UDI) Rule Implementation and Compliance Guide

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 - C. Global Unique Device Identification Database (GUDID) Guidance
 - D. European Commission UDI Draft Recommendations
 - E. FDA Guidance on 21 CFR Part 11
 - F. Unique Device Identifier System: Frequently Asked Questions, Vol. 1
 - G. Unique Device Identification System: Small Entity Compliance Guide
 - H. UDI: Direct Marking of Devices – Draft guidance