

MASTER YOUR DATA. STRENGTHEN EUDAMED COMPLIANCE.

Efor & Ackomas: regulatory expertise and digital integration to support EUDAMED compliance.

Turn complex device master data into structured, validated EUDAMED records built for accuracy, traceability and ongoing compliance.



YOUR CHALLENGES



EVOLVING EUDAMED OBLIGATIONS How to determine which EUDAMED requirements and transition provisions apply to your devices and data?



MASTER DATA COMPLEXITY How to clean, consolidate, and format scattered device master data across disparate systems?



SYSTEM & DATA INTEGRATION How to automate data synchronization with EUDAMED modules while satisfying data integrity ?

OUR APPROACH

01 ASSESS

- Regulatory readiness assessment and EUDAMED gap analysis
- Mapping, collection, and organization of required device master data
- Prioritized compliance roadmap and deployment schedule



02 IMPLEMENT

- Master Data constitution and data integrity validation
- Seamless deployment and configuration of the Ackomas software solution
- Automated publication, direct synchronization to EUDAMED, and full CSV execution



03 ASSURE

- Data consistency, completeness and traceability
- Submission testing and validation
- Controlled processes for EUDAMED data updates
- Dedicated technical and regulatory expert hotline
- Targeted training for regulatory, QA, and IT stakeholders

YOUR BENEFITS



CLEAR REGULATORY ALIGNMENT

EUDAMED obligations and applicable transition requirements clearly identified for your device portfolio.



RELIABLE MASTER DATA

Structured, validated device data supporting accurate, consistent and traceable EUDAMED records.



ROBUST EUDAMED INTEGRATION

Controlled data flows and system interfaces supporting reliable submissions and ongoing data integrity.

**READY TO STRENGTHEN YOUR
EUDAMED COMPLIANCE?**

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