



Effectum Medical AG

assists in the development of innovative medical devices and technologies throughout the certification and life cycle management process. Acting as legal manufacturer and by offering our Plug & Play (e)QMS, we speed up your time to market and ensure compliance and excellence in every stage of development.



E F F E C T U M
M E D I C A L

OUTSOURCED LEGAL MANUFACTURER



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OUTSOURCED LEGAL MANUFACTURER

Unlock Your Potential with Our Services

We offer a complete solution for companies to outsource the legal manufacturing, providing a dedicated support team throughout the entire product lifecycle. With products certified under MDD, MDR, and IVDD, partnering with us can help accelerate your time-to-market significantly.

Faster Market Access

- We partner with two leading Notified Bodies, eliminating your search effort
- Take advantage of our QMS certification (ISO 13485/ MDR/IVDR/ QSR)
- „Fast track certification“ provides significant time savings

Mitigate Risks

- QMS audited annually since 2017
- Minimize risks with our approved and trusted supplier network
- Expand your team with experts in every domain of the medical device industry

Focus on your core activities

- Concentrate on new product development and lifecycle management
- Grow your business—we manage the regulatory and compliance challenges
- Maintain full control over your branding and design



Marius Wiederkehr

Head of Legal Manufacturing Services
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FAST TRACK CERTIFICATION



Fast Track Certification:

With our Fast Track Certification, products within the same scope (MDA/MDN Codes for MDR) and similar or lower risk can be marketed immediately after completing the Technical Documentation and issuing the Declaration of Conformity without a technical file review from the notified body.

We hold MDR certificates for several product groups (Class Is and IIa).

Covered Codes (As of 04/2025)

MDN 1202 Non-active non-implantable devices for administration, management and removal of substances including dialysis products (class Is)	MDA 0204 Other active non-implantable devices for monitoring and/or diagnosis (class IIa)	MDA 0313 Active, non-implantable prostheses and devices for rehabilitation (class IIa)
	MDA 0308 Active non-implantable devices for wound and skin care (class IIa)	MDA 0315 Software (class IIa)

QMS setup and certification after 9-15 Months*	ISO/ MDR Audit	Technical File / Product Certification after 6-24 Months*	MDR CE
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Plug & Play (e)QMS (no Legal Manufacturer by Effectum Medical) →

Effectum Medical acts as Legal Manufacturer →

Fast Track Certification
1 - 3 Months* →

*The specified time periods are based on our experience and are non-binding. Liability is excluded. They also do not include the writing of technical documentation.

Contact us today to find out how we can help streamline your product certification process and get your products to market faster!



WHAT OUR CLIENTS SAY



“We have chosen Effectum Medical as Legal Manufacturer because we liked the approach to share risk, simplify processes, reduce time-to-market by increasing efficiency. When challenges arise, the team is very solution-oriented and shows a pragmatic approach. Overall, it is a very satisfying collaboration for us, and we have not been disappointed.”

Emanuela Pufe,
Leiterin, mediQ (PDAG)



„Effectum Medical guided us through the regulatory landscape, enabling us to work efficiently under a Quality Management System (QMS) despite initial inexperience. Their eQMS significantly improved accuracy, reduced errors, and saved time. Most importantly, we accelerated time-to-market by bypassing the need for our own Notified Body relationship and internal resources. Our successful launch, even without reimbursement, proves that patients are willing to pay out-of-pocket.”

Frederic Fappereau,
Head of Product, Biospectral

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SUPPORT OPTIONS



Outsourced Legal Manufacturer:

Streamline market access and compliance with our trusted legal manufacturer services. Need speed? Ask about our fast-track option!



Plug & Play (e)QMS:

Accelerate your launch with a ready-made EN ISO 13485, MDR/IVDR, and FDA-compliant quality management system.



Plug-and-Play eQMS Bootcamp:

Flexible training to empower self-management of your QMS.



Rent an Expert:

Add quality management and regulatory expertise to your team as needed.



Regulatory Opinions:

Access expert insights for global compliance and market strategy.



AI/ML Device Certification:

Certification services for AI and machine learning-based medical devices.

For more information visit:
www.effectummedical.com



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