



Regulatory Fundamentals Training for Medical Device Consultants under § 83 MPDG

Solid expertise for your professional practice – concise, clear and hands-on.

 Available in German and English



3.5 Hours

Total training duration



8 Modules

Structured and sequential



Online, at your own pace

Location-independent, anytime



Handouts

For every module, for reference



Final Exam

15 questions · 75% pass mark



Certificate

Upon passing the exam

The training provides a structured introduction to the regulatory fundamentals for medical device consultants under § 83 MPDG. Across eight sequential modules, you build the required expertise in the legal framework, due-diligence obligations and documentation – location-independent and at your own pace. Each module comes with a handout for reference in your daily work.

PARTICIPATION FEE

€279 net

per person · plus statutory VAT

REGISTRATION

Register by email to info@qm-medizintechnik.de. You will then receive a personal access link (valid for 1 week).

 info@qm-medizintechnik.de



If you pass the exam, you receive a certificate. If you do not pass, you receive a certificate of participation.