



FOR MEDICAL DEVICE, IVD, AND COMBINATION PRODUCT MANUFACTURERS

# TO NAVIGATE THE PATH TO REGULATORY COMPLIANCE, YOU NEED A TRUSTED PARTNER. THAT'S NSF.

**Trust NSF. We're a proven partner, helping manufacturers as they bring new and innovative** medical devices, IVDs, and combination products to market. With our expert advice and complete range of services meeting international regulatory compliance and standards is faster and more efficient.

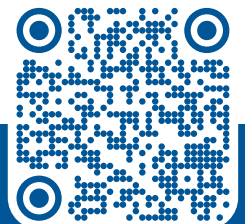


**NSF experts have "BEEN THERE, DONE THAT" and help prepare and train your team through:**

- > Remediation
- > Consultation
- > Mock inspections
- > Gap analyses
- > Acquisition diligence support
- > And much more

## Complete training, consulting, and auditing services:

- > US FDA, EU MDR/IVDR Regulatory & Compliance Experts
- > Mock Inspections & Gap Assessments to ISO 13485, ISO 14971, US and EU Regs, MDSAP and more
- > Remediation Services
- > On-site and Remote Project Management
- > Acquisition Diligence Support
- > Corporate and Supplier Audits
- > Regulatory Strategy
- > Quality Management Systems
- > Risk Management
- > Technical Documentation
- > Clinical Evaluation & Clinical Investigation Support
- > Trustee Service



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