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

Get started today by emailing: **medicaldevices@nsf.org**
Visit: **www.nsf.org/medical-devices**