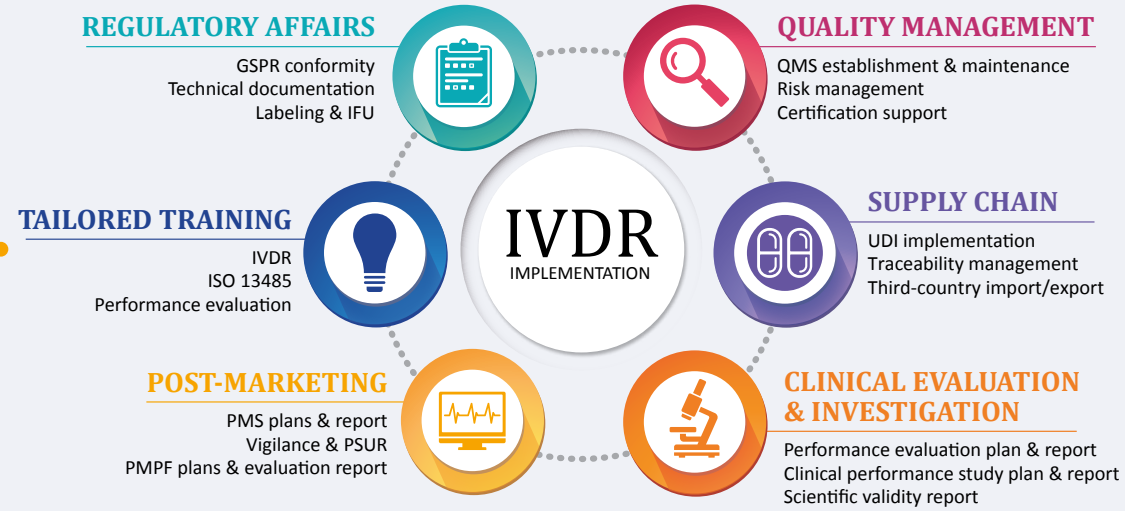
Targeted IVDR implementation for your *in vitro* diagnostics

Experts at SFL know what Notified Bodies and Regulators want.

Combining Notified Body and industry insights with long-term experience in implementing regulatory requirements, the SFL team can support you with all aspects of IVDR implementation for *in vitro* diagnostic medical devices. Our expertise covers a broad range of *in vitro* diagnostics, including companion diagnostics, genetic tests, and near-patient testing devices.

SFL's experienced cross-functional teams, including PhDs and medical doctors, perform gap analyses and create, review and update technical documentation, clinical performance evaluation as well as quality and risk management systems according to the IVDR requirements.

Our in-house cross-functional expertise includes Regulatory Affairs, Quality Management, Supply Chain, Clinical Performance Evaluation, Post-Marketing, and Tailored Training.



Take advantage of SFL's extensive experiences to get ready for your conformity assessment under the IVDR. We would be happy to set up a 30-minutes call or meeting free of charge to discuss your project and how we could support you.

Please contact us at office@sfl-services.com

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Solutions For Life sciences