

Key stages where we partner with companies



Specialised in Medical Device, IVD & Software as a Medical Device (SaMD)

Our Impact

125 200+ 350
Approvals Customers Projects

Experts

We understand the cross functional subject matters better than the conventional consulting

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Registered Office

No. 213, GF, Mallappa Layout, Babusapalya, Bengaluru, Karnataka, India - 560 043.

Scan to book a





free consultation

Our Services

Regulatory Consulting

- CDSCO Licenses (Manufacturing, Import, Test, Wholesale)
- USFDA approvals (510k, IDE, Denovo)
- CE Certification (EU MDR & IVDR)
- Regulatory Strategy & Gap analysis
- Predicate analysis/Substantial Equivalence
- Technical Documentation (DMF, PMF, CER, Risk Management etc.)

Startup Support

- Quality & Regulatory Staff augmentation
- QMS Audit assistance
- Bench testing guidance (stability, sterility, packaging etc.)
- Manufacturing facility design
- Grant application assistance (BIRAC, PRIP, ELEVATE, SAMRIDH etc.)
- Investors connect

Quality Consulting

- ISO 13485, 27001, 9001
- ISO 15189 & 17025
- MDSAP & QMSR
- DSIR Certification
- · BIS Certification
- NABH, NABL & CAP Accreditation

Pilot Clinical Study

- Protocol Development
- IEC approval & CTRI registration
- Feasibility Study for Device, IVD and SaMD
- Health Technology Assessment
- IBSC & GEAC approvals
- IVD kit's PQ evaluation

Licenses & Registrations

- GeM Registration
- SOFTEX Registration
- EPR Registration with CPCB (e-waste, battery and plastic)
- · Legal Metrology License
- WPC ETA (Wireless Equipment Type Approval)
- Factory License

Prototyping

- Wearable & IoMT (Internet of Medical Things)
- Point of care (POC) & Remote patient monitoring
- Digital diagnostics & Digital therapeutics
- Pre & full compliance testing services (EMI/EMC) assistance
- Cybersecurity testing
- ESDM (Electronics System Design & Manufacturing)

Fast-track your innovation in Medical Devices, Diagnostics & Digital Health

Expertise Across Global Compliance Frameworks





















