CURRICULUM VITAE

Dr. Vidya Sagar

Founder, CEO & Managing Director – Salvation Global Pvt. Ltd.

EXECUTIVE PROFILE

Visionary MedTech leader, scientist, and entrepreneur with over 20 years of global experience in medical devices, regulatory affairs, quality systems, clinical trials, artificial intelligence, and product innovation. Founder & CEO of Salvation Global Pvt. Ltd., driving transformation in medical device development, regulatory intelligence, and global compliance ecosystems. Holds 6 international patents (4 granted), 65+ publications, and extensive experience in CE, FDA, ANVISA, CDSCO, IMDR, and MDSAP regulatory pathways.

CORE COMPETENCIES

- Global Regulatory Affairs (EU MDR, FDA, ANVISA, TGA, PMDA)
- Quality Systems & Risk Management (ISO 13485, ISO 14971, MDSAP)
- Medical Device Product Development & Clinical Trials
- AI/ML & Digital Transformation in MedTech
- Intellectual Property & Patent Development
- Leadership, Strategy & Business Development
- Academic Mentorship & Scientific Research

PROFESSIONAL EXPERIENCE

- Founder, CEO & Managing Director – Salvation Global Pvt. Ltd. (2022 – Present)
- Head – QA/RA – Relisys Medical Devices, Hyderabad (2011 – 2020)
- Chief Scientist – Relisys Medical Devices, Hyderabad (2005 – 2011)
- CSIR Project Fellow – Relisys Medical Devices (2003 – 2005)

EDUCATIONAL QUALIFICATIONS

- Ph.D. – Medical Devices, Kakatiya University (2001–2006)
- M. Pharm (Pharmacology) – Kakatiya University (1999–2001)
- B. Pharmacy – Kakatiya University (1994–1998)
- PG – AI/ML/Data Science – University of Texas, Austin (2021–2022)
- MBA (Artificial Intelligence) – NIBM (2021–2022

PATENTS (6 TOTAL, 4 GRANTED)

- FORMULATION & COATING FOR SIROLIMUS STABILITY – Patent No. 2878/CHE/2011
- ELECTRO-POLISHING TOOL FOR IMPLANT DEVICES – Patent No. 63/CHE/2012
- EVEROLIMUS COATING FOR MEDICAL DEVICES – Patent No. 62/CHE/2012
- HEMOSTATIC FORMULATION PROCESS – PCT/IB2021/051222

SCIENTIFIC CONTRIBUTIONS

- 65 international publications and 25 national publications.
- Research areas: cardiology, pharmacology, biomaterials, metabolism studies, ML in MedTech.

CERTIFICATIONS

- ISO 13485 – Lead Auditor (BSI)
- ISO 14971 – Risk Management (GLC)
- MDSAP Auditor Training – DQS
- PG Diploma – Clinical Data Management
- Applied Biostatistics – ISI

CURRICULUM VITAE

Mr. Udayabhaskhar Kilari

Quality Assurance & Regulatory Compliance Expert

Professional Summary

Quality Assurance (QA) and Regulatory professional with 10+ years of experience in ISO 15189, ISO 13485, and ISO/IEC 17025-based systems across diagnostics, medical devices, and healthcare. Skilled in internal audits, risk management, CAPA, documentation control, and accreditation processes for laboratories and MedTech organizations.

Core Skills

- ISO 15189, ISO 13485, ISO/IEC 17025 Auditing
- Clinical Laboratory QA/QC Systems
- Internal Audits, Risk Management & CAPA
- SOP Development & Regulatory Documentation
- Conformity Assessment for Diagnostics & Medical Devices
- Training & Capacity Building

Professional Experience

- Scientist D – KIHT, AMTZ (May 2023 – Present)
- Assistant Director – NABL (Feb 2023 – May 2023).
- Executive Officer – NABL (Sep 2016 – Aug 2020)
- Technical Support Specialist – Ortho Clinical Diagnostics (May 2016 – Sep 2016)
- Clinical Biochemistry Incharge – Manipal Super Speciality Hospital (May 2013 – May 2016)
- Tutor – Krishna Teja Pharmacy College (Sep 2012 – Apr 2013)

Education

- Master of Science in Medical Biochemistry – Sri Venkateswara University of Medical Sciences.
- ISO 15189 Auditor Course – Siddhartha Medical College (2016)

Key Achievements

- Led quality systems implementation aligned to ISO 13485, facilitating clinical validation readiness for MedTech startups.
- Guided ISO 15189 accreditation for diagnostic labs that supported early-phase trials and disease surveillance programs.
- Trained over 150 lab professionals and auditors in regulatory-compliant QA practices relevant to research and diagnostics.

CURRICULUM VITAE

Ms. Revathi Korupolu

Quality Management & Regulatory Affairs Professional

Professional Summary

Experienced Quality Assurance and Regulatory Affairs professional with strong expertise in ISO 13485, ISO 9001, and ISO 17021-1-based certification systems. Skilled in medical device QMS audits, documentation control, CAPA, risk management, SOP development, and regulatory compliance. Demonstrated background in diagnostics, CROs, and medical device industries.

Core Skills

- ISO 13485:2016 & ISO 9001:2015 Lead Auditor
- QMS & MDQMS Auditing
- Regulatory Documentation & Technical File Management
- Risk Management & CAPA
- Internal Audit Planning & Implementation
- SOP Development & Quality Documentation
- Regulatory Compliance for Medical Devices

Professional Experience

- Quality Manager – KIHT Certification Services (KCS), AMTZ (Jun 2022 – Present)
- Executive – QARA, Trivitron Healthcare Pvt. Ltd. (Nov 2021 – May 2022)
- Assistant – QA, DNA Xperts Pvt. Ltd. (Dec 2020 – Aug 2021)
- Team Lead – Longdom Group (Dec 2017 – May 2020)

Education

- Master of Pharmacy (Regulatory Affairs & Pharmaceutical Management) – Andhra University (2017)
- Bachelor of Pharmacy – Andhra University (2015)

Certifications

- ISO 13485:2016 Lead Auditor – BSCIC
- ISO 9001:2015 Lead Auditor – Intertek
- ISO/IEC 17021-1 & 17021-3 Accreditation Training – APAC
- Biomedical Quality Assurance Certification – IBSC
- Training on Indian Medical Device Rules (IMDR 2017)

Publications

- Review on Coronary Bioresorbable Vascular Scaffold System – JOCCT (2017)
- Review on European Medical Device Regulations – PharmaTutor (2018)
- Pre-Clinical Requirements for Drug-Eluting Stents – IJRASET (2023)
