



Revolutionizing Pharmaceutical & Lifescience Solutions



Global Pharma Regulatory,
Quality Services & Software
Company



Leadership/Scientific Team



Dr. Ramesh Pennamareddy
Founder & Managing Director



Mrs. Swarna Priya Pennamareddy
Founder & Chief Executive Officer



Dr. Muralidhara Babu Gavini
Director & Former USFDA GMP Inspector



Dr. Khemraj Hirani
Chief Regulatory Officer - Masuu USA



Mr. Shiv Kumar Somani
Director - Operations & Technical



Mr. Concepción "Coki" Cruz
Former USFDA GMP Inspector



Mr. Ariel Cruz Figueroa
Former USFDA GMP Inspector



Mr. Akhilesh Khale
Former TGA GMP Inspector



Mr. Harvey Jaramillo
Former INVIMA Regulatory Technician



Ms. Sara Ruggieri
Former AIFA PV Specialist



Dr. Walid Smadi
Technical Head MENA Region



Mr. Ankamma Rao Thotkura
EU & UK QP (Qualified Person)



Mr. R V Ravindranadh
Director - International Marketing



Mr. B.H.K. Satyadev
Sr. Vice President - Audits & Training

Industries



Formulation



R&D Labs



API/Raw Material



Medical Device 510(K)



CMO/CDMO/CRO



Analytical Labs



Packaging & Excipients



OTC/Cosmetics

Dossiers



DMF/CEP/ASMF



ANDA



NDA/505(b)(2)



IND/MAA/BLA



PMA/510(K)



ANADA/NADA



ANDS/NDS



Drug Listing

Our Services



Global Dossier Filing



Medical Device
510(k) Filing



Quality, GMP &
GxP Audits



Quality System
Development



Due Diligence/
Acquisition/Merger



eCTD Publishing &
Submission



Labeling & Artwork



Medical Writing & PV



Global Language
Translation



NextGen eCTD
Software



Toxicology Reports
(PDE/OEL/OEB)



US Agent Services

Our Expertise & Matrix

Offering Comprehensive Services and Software for Pharmaceuticals, Biopharmaceuticals, Life Sciences, Cosmetics, Nutraceuticals, and Medical Devices.



39+ Countries Serving

50k+ Projects Delivered



100+ Global Partners

8+ Office Locations



150+ Regulatory Affairs SMEs

50+ Quality/cGMP SMEs



80+ Pharma Software Developers

450+ International Clients



About Us & Deliverables

Masuu Global Solutions: Your Trusted Partner in Pharma Regulatory, Quality & PV Services, and Software Solutions.



700+ Global Dossier Submissions

5+ In-house Software
(eCTD, SPL, PDF Plugin, eLN and eQDR)



400+ Software Setup & Installation

250+ Medical Writing/PV



100+ US Agent & FDA Registration

500+ Quality, cGMP & GxP Audits



10,000+ Global Labeling & Artwork

45k+ eCTD Publishing & Submission



Our International Clientele



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