







CRDO & CDMO



**Custom Synthesis** 



Certified Reference Standards



Nitrosamines Standards



Stable isotopes and Metabolites



Analytical Services

## **About Us**

K. M. Pharma Solution (P) Ltd, is an integrated CRDO (Contract Research and Development Organization) and GMP CDMO (Contract Research and Development Manufacturing Organization).

We are a STATE-FDA Audited & Accepted Facility & bagged with ISO 17034, ISO 17025 & ISO 9001 Certification. We are specialized in chemistry based innovation driven company involved in synthesis of nitrosamines impurities, pharmacopeial and non-pharmacopeial impurity standards, reference standards, drug metabolites, glucuronides, process degradants, excipient related impurities, unknown impurity isolation and characterisation and stable isotope products. Also is an integrated contract research, development and manufacturing organisation providing scientific services - from early discovery to commercial supply.

We are Supplying all the Pharmacopeial Reference Standards (i.e. USP, EP, BP, IP, JP). Apart from this we offer Pesticide Standards, Phyto-chemical Standards, Food & Environmental Standards, etc. K. M. Pharma is having more than 6,000 In-house Impurities ready to ship available.

We are providing all the Analytical Services. (i.e. Method Development, Method Validation & Transfer, Stability Studies, Contract Analysis, etc)

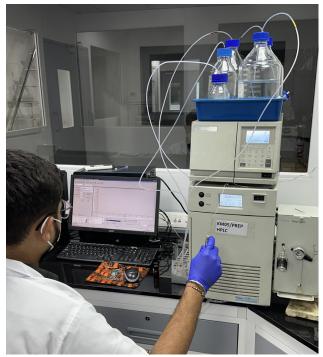
**WHO GMP Employees Founded R&D Centre Manufacturing Sites** 2012 100+ 01 01 Certified **International** Commercial Indian Reference **APIs** Customer **Customer Standards** 10,000+ 100+ 25+ 15+

















## **Accreditations**



## Clientele







































