



A phase-appropriate manufacturing philosophy in our facilities, which are compliant with the regulatory guidelines is followed {present day technological exclusive current good manufacturing practices (cGMP) and manufacturing capabilities}. A demonstrated capabilities and experience to manufacture batches of small molecule drug substances, advanced intermediates as well as regulatory starting materials for pre-clinical, clinical, and commercial applications at appropriate facilities, and are further expanding our capabilities to cater to growing demand from global innovators in both pharma (Human Health & Animal Health) as well as non-pharma industries.

A dedicated state-of-the-art cGMP manufacturing facilities support the small molecules contract manufacturing business. Since inception, we supply clinical (early & late phase) as well as commercial drug substances (NCEs), advanced intermediates, regulatory starting materials and select generic APIs.



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The Doctor/Lawyer Team

Revive Medico Legal, has a team of lawyers who are also doctors. This enables us to better litigate cases that involve pharmaceuticals and medical issues, especially complex medical malpractice cases.

Revive team's holistic approach and in-depth knowledge of the industry with a core experienced team allows delivering an integrated service offering to life sciences and healthcare companies around the globe.

Revive would be happy to assist you regardless of what you produce or sell —medicines, medical devices, cosmetics or household chemicals.

A collaborative approach brings the customers our consulting capabilities in compliance, tax, customs, and anti-monopoly regulation along with the access to the resources of a global out reach.

In our Medical Law practice, Revive Medico Legal Services provides advice on legal compliance issues to medical professionals, hospital management, healthcare networks and nursing home facilities, defend medical practitioners and institutions from cases of medical malpractice, as well as pursuing claims of malpractice for patients.

Revive Medico Legal services is well renowned for its wisdom of medical legal experience. Our team has extensive experience in medical legal cases commerce with both patients and medical sources. Our team of legal experts, physician advisory panel, paralegals, trained support staff and demonstrative experience gives us the competitive benefit in medical legal cases, including, Medical Legal Counseling, Contracting & Negotiation Services, Empower Teams & Increase Productivity, Contract Management Services and Solutions, Contract Management Services, Consumer Laws.

We at Revive Medico Legal are focused to provide unified Legal & Conciliatory solutions for Medical Fraternity, Medical Regulatory & Industry Oriented, Companies engaged in manufacturing, clinical trials, supply of medicines, medical devices and extend special legal services relating to the usage of biological materials, utilisation of medical waste, and circulation of cosmetics and household chemicals.

Pharmaceutical industry is rapidly growing across the globe.

We at REVIVE help the pharmaceutical sponsors with robust technology scope in the industry with harmonizing international standards.

Discovery and Development:

Research and development testing of molecular compounds and new technologies followed by the development of a new drug in the analytical and chemical laboratory.

Pre-clinical Phase:

Pre-clinical research is mainly conducted on microorganisms and animals for answering basic questions about drug safety.

Clinical Research:

Drug testing on people, which is further divided into Phase I, II, III & IV of clinical trials.

REVIVE's Global Regulatory Affairs Services enable Life Sciences, Consumer Pharma and Bio-Med companies to understand and efficiently manage the demanding complexity of the diverse Regulatory requirements in order to successfully launch their products in new markets and maximize their market value.

Regulatory offerings will be successfully in completing the audits from Indian & International regulatory agencies, sponsor audits from Indian and International clients and study submissions will also be made to other countries.

All the services are executed in strict compliance with regulatory and ICH GCP requirements.

(CSR):

Corporate Social Responsibilities: Our mission is to improve the lives of people and patients through scientific research, improving the quality of life and environment for the communities around us and build social and economic equity, providing economic stability for women entrepreneurs, provide access to basic amenities in collaboration with the state governments and Non-Governmental Organizations (NGOs).

Most of the programs are undertaken for a minimum of three years to provide sufficient time for an intervention to become sustainable.

