Vanank Regpharma Solution

Vanank Regpharma Solution has emerged as a reputed exporter and supplier of a varied range of products. The assortment of products that we offer to the clients includes Megazole Powder, Closantel Base Powder, Closantel Sodium Dihydrate, Cyromazine Premix, Fenbendazole Powdera and many more. We Deals in Services like Pharmaceutical Contract Manufacturing Services and CTD and ECTD Dossier Preparation Services. Backed by a reputed group of vendors, the company brings the finest quality products as per the specific needs of the clients. What makes us stand out among others working in similar domains is the fact that we never compromise on the quality of the products that we offer? Additionally, the clients can avail our products at highly competitive prices.

company profile

Vanank Regpharma Solution is a Haryana; India based company and aims to be global leader in the area of operations. Since inception, the company is running under the supervision of professionals and scaling new heights of success.

Quality Satisfaction

We are a quality-conscious firm and try hard to maintain in our products and services. Also, we have adopted strict measures for ensuring high quality standards. Apart from this, we sternly check the entire lot prior to final dispatch for the attainment of complete client satisfaction.

Industries We Serve

We have satisfied clients in India and overseas with our dependable and cost effective services. We provide our services to the following industries :

- Pharmaceutical Formulations
- Pharmaceutical APIs / Bulk drugs
- Herbal Manufacturers
- Biotech Industries

Why Us?

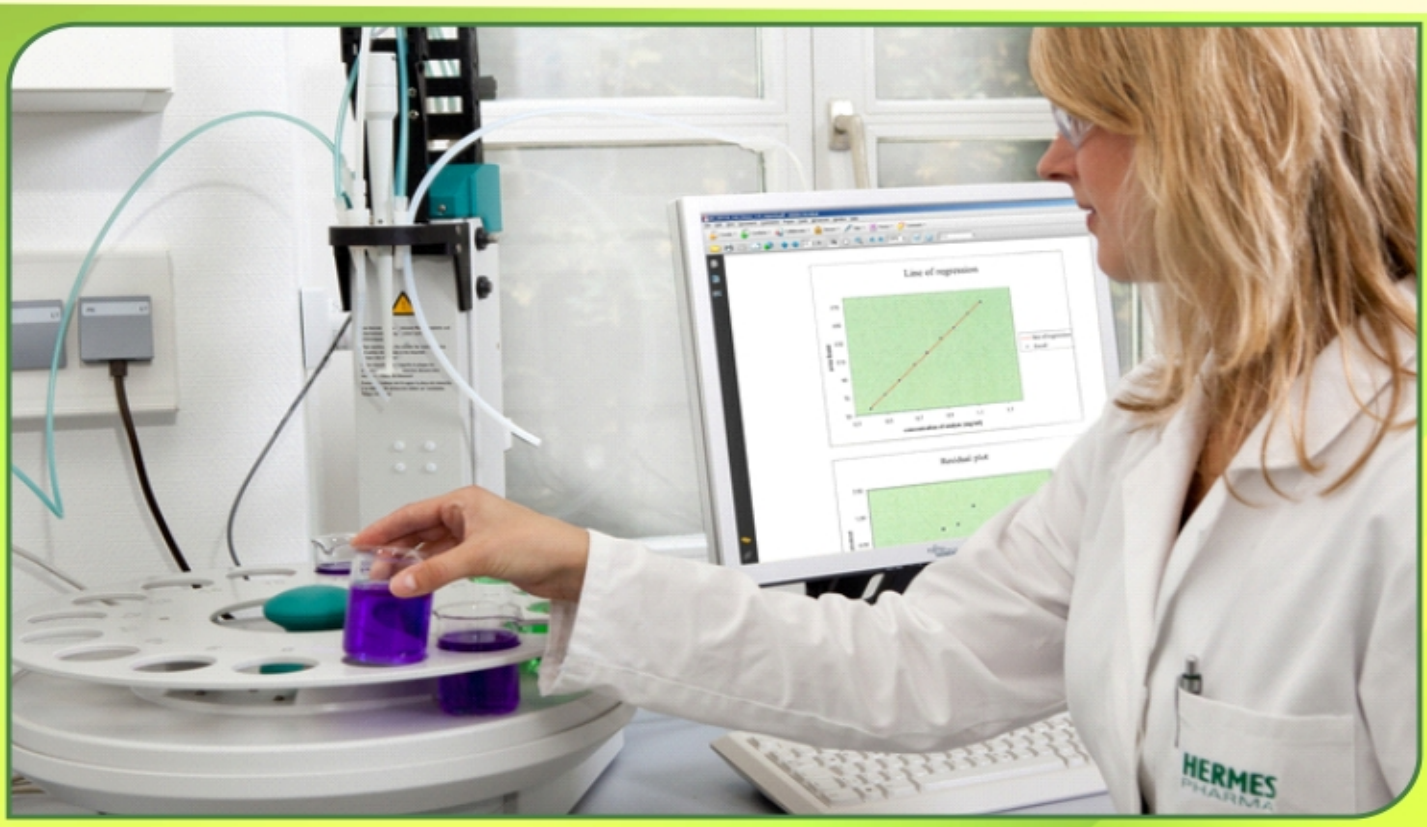
Points that distinguish us from others are :

- Technical Expertise
- Complete Database
- Quality services
- On-time delivery
- Complete Dossier Solution

our products

Analytical methods validation

The robustness of an analytical procedure is a measure of its capacity to remain unaffected by small, but deliberate variations in method parameters and provides an indication of its reliability during normal usage

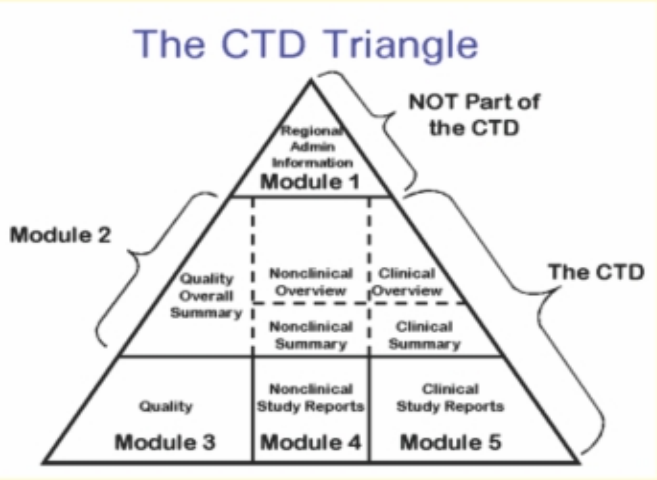


BA/BE study

assess equivalence, including: i comparative bioavailability (bioequivalence) studies, in which the active. drug substance or one or more metabolites is measured in an accessible. biological fluid such as plasma, blood or urine. ii comparative pharmacodynamic studies in humans.

CTD And ACTD Dossier For CIS And Asean And Gulf Region

The Common Technical Document (CTD) is a set of specification for application dossier for the registration of Medicines and designed to be used across Europe, Japan and the United States. It is an internationally agreed format for the preparation of applications regarding new drugs intended to be submitted to regional regulatory authorities in participating countries. It was developed by the European Medicines Agency (EMA, Europe), the Food and Drug Administration (FDA, US) and the Ministry of Health, Labour and Welfare (Japan). The CTD is maintained by the International Conference on Harmonisation of Technical Requirements for Registration of Pharmaceuticals for Human Use(ICH)



Clinical Trial Report

In medicine, a clinical study report (CSR) on a clinical trial is a very long and detailed document giving much detail about the methods and results of a trial. A CSR is a scientific document addressing efficacy and safety, not a sales or marketing tool; its content is similar to that of a peer-reviewed academic paper.[1] Results of trials are usually reported in a briefer academic journal paper, but methodological flaws are often glossed over in the briefer paper.[2]

The International Conference on Harmonisation of Technical Requirements for Registration of Pharmaceuticals for Human Use (ICH) is a body bringing together the regulatory authorities and pharmaceutical industry of Europe, Japan and the US to discuss scientific and technical aspects of drug registration;[3] in 1995 it produced a tripartite harmonised ICH guideline on the format and content of a study report to be acceptable in all three ICH regions.[4] Recommended prerequisites and content for producing a report conformant to ICH guidelines have been outlined by SE Caldwell.

our products

Contract Manufacturing of Finished Product for Export And Domestic Purpose

Vanank Regpharma Solution have tieup with the many WHO-GMP certified plant , who are making the tablet, capsule, injection, liquid syrup, in india and overseas.

Vanank Regpharma Solution understands the importance of Research & Development in the present scenario. When we are committed to give the best Quality Products at most economic rates, the usage of R&D in developing newer combinations, rather rational combinations in a single dosage form, becomes utmost important. For example, for the treatment of a single disease, when patient has to take three to four tablets, he feels disgusted & there are quite good chances that he may forget to take one or another tablet. In Today's era of Jet Age, when everything is so fast, patients as well as the doctors equally find themselves helpless. We at Vanank Regpharma Solution swung into action to provide them whatever is their need. This is not a child's play. This requires lot of R&D, as we have to take care or rather ensure that the bio availability of individual drug should not be altered or affected. Similarly their dissolution rate, dispersibility rate (D.T.) etc. should not be changed. In it's product basket, Vanank Regpharma Solution has so many such products to offer.

The main aim of R&D at Vanank Regpharma Solution is to simplify the sophistication / complications of latest technology so that it can be used for the welfare of masses.

Second but not the last , R& D ensure that each dose/ particle of drug in Tablet or Capsule or every drop of liquid oral should give the claimed benefit to suffering patient & this benefit should be there till last particle or drop. This will instill the faith in patients that still there are medicines which treat them fully.

Tablet

Tablet Manufacturing and Packaging Just about any size, any shape, any coating available for your custom formulation.

our products

Syrup

Syrup, Dry Syrup, Suspension, Drops manufacturing with various packing available.

Capsules

Available as antibiotics, pain killers, vitamin supplements in a variety of sizes and packaging

Ointment

We manufacturing include balms, creams, gels, oils, lotions, patches, ointments and other products that you apply to your skin.

CTD and ECTD Dossier Preparation Services

Deeply rooted in Kurukshetra (Haryana), we are an organization that renders excellent CTD and ECTD dossier preparation services to the clients at competitive charges. ECTD means an electronic common technical document that is required by the pharmaceutical industry for transferring regulatory information. Thus, CTD & ECTD is an essential documentation that requires the right assistance. We are a group of committed professionals who strive to provide proficient solutions to our valuable clients.

Vanank Regpharma Solution

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