

Oasis group of companies

The Group companies

Analytical services

- Oasis Test House Limited, Jaipur
- Oasis Test House, Ahmedabad

Pharma production contract services

- Jaipur Pharmaceutical Works, Jaipur
- Cris Pharma (India) Limited, Dehradun

Software services

- Oasis Infotech, Jaipur

Vision

To provide prompt, consistent, reliable, cost effective quality services to pharma sector, worldwide, maintaining confidentiality and integrity.

Milestones

Jaipur Pharmaceutical Works, Jaipur started in 1970

Oasis Test House Limited, Jaipur established in 1981

Oasis Infotech started in 1987

Development of LIMS and Pharmaster in 1987

Development of LIMS, 21 CFR Part 11 compliant version

Development of CrisCon, Pharma ERP for compliance with cGMP / WHO GMP

Pharma production contract services

- Jaipur Pharmaceutical Works, Jaipur
- Cris Pharma (India) Limited, Dehradun

Jaipur Pharmaceutical Works, a professionally managed organisation firmly believing in ethical promotion of high quality therapeutic formulations, has been in operation for over 35 years and consistently making efforts to prosper and grow.

The company has state-of-the-art, modern (GMP, ISO 9001) certified production facility at its unit situated at Jaipur, Rajasthan, India. It manufactures different dosage forms, such as tablets (general and beta lactam), liquid orals and capsules (general and beta lactam) and has spread its wings not only in manufacturing products for many reputed pharma companies, but also in marketing its products.

The company employs strict measures at each stage of production to maintain quality of products, in conformance with the requisite standards, using pharma specific ERP (enterprise resource planning) software package.

The company specializes in outsource contract manufacturing for many pharma companies including MNCs. With growing product offering and clientele, the group has expanded manufacturing facility and operations with its new plant at Dehradun, Cris Pharma (India) Limited. It is also managed by a dedicated team of professionals under the guidance of its CEO. It is looking forward to touching new heights by selling its products in sixteen states, marketing newer and latest formulations across the country successfully.

Quality Policy

To provide consistently quality products and services as per the need of the society, by having commitment and involvement of the employees, to do things right first time and every time, and continual compliance and improvement in the quality management system.

Our commitment

Diligently practice the QMS and thus to serve customers with great and prompt responsiveness.

Enhance customer satisfaction by meeting their requirements, expectations, besides complying with relevant statutory and legal obligations.

Establish, implement and review the quality policy and its objectives, with a view to ensure their continuous suitability through improvements as necessary.

Analytical services

- Oasis Test House Limited, Jaipur
- Oasis Test House, Ahmedabad

Oasis Test House Limited, a reputed government approved public testing lab (ISO 9001 certified), is another group company and is one of the leading test laboratories in India. It has its branch at Ahmedabad also.

The test labs have excellent, modern, updated facilities at its units at Jaipur, Rajasthan and Ahmedabad, Gujarat and provide analytical services, outsourcing, inspection and certification services to over 800 clients in India as their preferred partner for quality, professionalism, performance and solutions.

Use of sophisticated and modern instruments makes the testing facilities comparable to best anywhere. The labs are being managed by dedicated teams of qualified, experienced and dynamic professionals. Many of the professionals have been working with the labs for more than last 15-20 years. This enables the labs to provide quality services to the pharmaceutical industry.

The labs serve on-line with eoasislims.com, an innovative software to provide remote access to the laboratory data from anywhere in the world.

Software services

- Oasis Infotech, Jaipur

Oasis Infotech

This group company has been instrumental in developing software to computerise the whole gamut of manufacturing and analytical activities. It has been serving the pharma industry for over 19 years with its software packages; has been successful in undertaking varied assignments of software development, implementation and consulting services to pharma industry, bulk drug, consumer health care units, QA/QC laboratories and contract research organisations, including many multinationals.

Product line

OasisLIMS 1.0 for Windows / LIMS for DOS

OasisLIMS 1.0 for Windows (21 CFR Part 11 compliant)

OasisLIMS 1.0 for Windows for Commercial testing laboratories

OasisLIMS 1.0 for Windows for CRO (Contract research organisations)

CrisCon 1.0 for Windows (ERP Solution, for compliance with New Schedule M)

OasisLIMS: Laboratory Information Management System (LIMS)

OasisLIMS, an innovative software solution, encompasses unleashing power and excellent graphical user interface (GUI), such that the user feels at home with windowing environment. It is a configurable client / server LIMS solution comprising of various features including, sample registration and tracking, specifications and limits, SOPs / STPs maintenance, protocols and worksheets, bar coded labels, test maintenance and automatic calculations, reporting and certification, automatic sample approval / rejection, MIS / DSS tool and many add-on modules. The working of add-on modules in conjunction with Core LIMS brings numerous advantages in implementing the latest regulatory / quality assurance requirements.

CrisCon: Enterprise Resource Planning and Management (ERP)

CrisCon is an ERP (Enterprise Resource Planning) and business management software solution, specifically designed for pharmaceutical manufacturing units. It integrates all area of business such as, sales, purchase, production, inventory management, quality assurance and control, accounts and invoices, human resource. It helps production managers and analysts to easily implement cGMP including new Schedule M guidelines, WHO GMP. **CrisCon** helps its users to maintain entire documents / records correctly, completely and timely, as per the requirements of new Schedule M. The user company is thus, always in a state of compliance with the regulatory requirements.

Valued clients

Johnson & Johnson, Fulford, Wyeth, Elder, Glaxo Smithkline, Sunways, Encube, Kilitch, Biodeal, Troikaa, Astron Research, Ivax, Brown & Burk, Micro, Rusan, Indoco Remedies, Neon, Inga, LiTaka, Ipca, RDPL, Akums, Helios, Marck, Alpa, HiGlance, BHPL, PRK Pharamanlyst, Cosmos (Nairobi), Smilax, Biogenetics, Cris, Indoco Healthcare, Next Wave, Lagray (Ghana), Astron Research (U.K.) and many more ...

About CEO

The group was founded by Mr. Vinod Kalani, a pharmacy graduate from BITS, Pilani, with a vision to establish the group as a global pharmaceutical sector service providing company. In fact, it is the spirit of entrepreneurship that has shaped the group to become what it is today and he is well known for his passion for exploring new ideas and innovation.

Board and management

The top management has many decades of collective experience in the pharmaceutical sector. It is committed to provide quality services to its customers, organisation and the employees. It plays active role in targeting the objectives, planning, business growth, allocation of resources, technology upgradation and strategy implementation.

The executive team

The executive team comprises of professionals who possess vast experience across several technology platforms.

The Executive team comprising the technocrats and officials of various discipline handle the day-to-day operations. They help the Board of Directors in implementing the goals and objectives by following the corporate strategy. Their technical guidance has helped the Organisation to grow to its heights.

Services offered by Oasis group of companies

Analytical research and development

- Analysis of bulk drugs, pharmaceutical formulations, cosmetics, food, water (chemical, microbiological, instrumental analysis)
- Test procedure development, stability indicating methods
- 2nd testing of production batches on randomized basis
- Stability protocol preparation, analysis and monitoring
- Stability testing as per ICH guidelines
- Dissolution profiling
- Analytical method validation
- Co-ordinating conduct of clinical / BE studies at Regulatory approved clinical test laboratory

Formulation research and development

- Formulation development
- Trial batches
- Trouble shooting

QA documentation

- Documentation as per new Schedule M
- Site Master File development
- MFR / BMR / BPR development, SOPs
- Process Validation
- Validation master schedule
- Calibration
- Self-inspection and quality audit
- Drug master file / dossier development

Training

- Training to analytical chemists
- Pharma ERP
- Validation

Software services

- Tailored software development including 21 CFR Part 11 compliance
- Multi-location interconnectivity of operations on WAN using various options
- Computerization of pharma production, QA and QC activities
- Up-gradation of existing software
- Software validation