

Committed for better healthcare through Global Partnership

LEADING MANUFACTURER & SUPPLIER OF QUALITY PHARMACEUTICALS

YOUR TRUSTED PARTNER FOR QUALITY PHARMACEUTICALS

GSS Pharma is an emerging pharmaceutical company in the export market. Our head office is based in Bangalore, India, and we have offices in the United Kingdom. We are committed to providing our customers with high-quality pharmaceutical products that meet international standards.

Our manufacturing facilities are state-of-the-art and certified to the highest standards, including WHO-GMP and ISO 9001:2015. We have a comprehensive product portfolio, including tablets, capsules, eye and ear drops, injectables, creams and ointments, liquids and syrups, and nutraceuticals.

We are a team of passionate, dedicated professionals committed to providing our customers with the best possible products and services. We are always looking for ways to improve, and we are committed to continuous innovation.

Our Vision

To be a leading global pharmaceutical company that provides high-quality products and services that meet the needs of our customers.

Our Mission

Provide customers with innovative, high-quality pharmaceutical products that meet international standards.

Our Values

- + Quality: We are committed to providing our customers with high-quality products and services.
- + Innovation: We are always looking for ways to improve our products and services.
- + Customer Focus: We are committed to meeting the needs of our customers.
- + Teamwork: We believe that teamwork is essential to our success.
- + Integrity: We are committed to conducting our business honestly and with integrity.

Why GSS Pharma

- + Quality: GSS Pharma is committed to providing the high-quality products and services. Our manufacturing facilities are state-of-the-art and are certified to the highest international standards, including WHO-GMP, NAFDAC (Nigeria), and FDA (Ghana).
- + **Range:** GSS Pharma has a wide range of products to choose from, covering a variety of therapeutic segments. This includes tablets, capsules, eye and ear drops, injectables, creams and ointments, oral liquids, and dry powder.
- + Affordability: GSS Pharma is committed to providing high-quality products at affordable prices. We offer competitive pricing on all of our products, and we are constantly working to find new ways to reduce our costs so that we can pass the savings on to our customers.
- + **Customer service:** GSS Pharma is committed to providing excellent customer service. We have a team of experienced and knowledgeable staff who are available to answer your questions and help you find the right product for your needs. We also offer a various customer support services, including online chat, phone support, and email support.
- + **Trust:** GSS Pharma is a trusted brand. We have been in business for over 20 years, and we have a long history of providing our customers with high-quality products and services.

Manufacturing Facilities

At GSS Pharma, we take immense pride in our state-of-the-art manufacturing facility, which serves as the backbone of our operations. With a commitment to excellence and a focus on innovation, our facility stands as a testament to our dedication to delivering the highest quality pharmaceutical products.

Cutting-Edge Infrastructure

Our manufacturing facility is equipped with cutting-edge technology and modern infrastructure to ensure the production of pharmaceuticals that adhere to global quality standards. We have implemented stringent quality control measures and follow Good Manufacturing Practices (GMP) to guarantee the integrity and safety of our products.

Industry-Leading Expertise

Our team consists of highly skilled professionals who possess a deep understanding of the pharmaceutical industry. With years of experience and expertise, they ensure that every step of the manufacturing process is conducted with precision and attention to detail. From research and development to production and packaging, our experts work diligently to deliver products of exceptional quality.

Quality Control

Quality control is at the heart of our manufacturing facilities. We have established a comprehensive quality assurance program that encompasses rigorous testing, robust quality control processes, and adherence to regulatory guidelines. From raw material inspection to in-process testing and final product analysis, we leave no stone unturned to ensure that our products meet the highest standards of quality and purity.

State-of-the-Art Testing Facilities

To support our quality control efforts, we have invested in advanced testing equipment and laboratories. Our testing facilities are equipped with cutting-edge instruments and are staffed by experienced analysts who perform a wide range of tests to ensure the efficacy, safety, and stability of our products. We continuously upgrade our testing capabilities to stay ahead of evolving industry requirements.

Continuous Improvement

We believe in the continuous improvement of our manufacturing processes. Through ongoing monitoring, data analysis, and feedback mechanisms, we identify areas for enhancement and implement corrective actions to drive efficiency and quality. Our commitment to continuous improvement is a testament to our dedication to delivering high-quality pharmaceuticals.

Sustainability and Environmental Responsibility

We are dedicated to environmental sustainability and reducing our ecological footprint. Our manufacturing facility incorporates eco-friendly practices, minimizing waste generation and optimizing resource utilization. We continuously explore innovative solutions to ensure that our operations align with our commitment to the environment and the well-being of future generations.

Trust GSS Pharma

With our cutting-edge infrastructure, industry-leading expertise, and an unwavering commitment to quality control, GSS Pharma stands as a trusted partner in pharmaceutical manufacturing. We take pride in delivering products that improve the lives of patients worldwide, and we invite you to join us on this journey towards excellence.

GSS PHARMA SERVICES

PHARMACOVOGILANCE

- + Complete ICSR Processing: Streamlined and efficient processing of Individual Case Safety Reports (ICSRs) to ensure timely identification, collection, evaluation, and reporting of adverse events.
- + Literature Screening: Thorough and systematic screening of scientific literature to identify relevant safety information and ensure comprehensive monitoring and analysis of potential risks.
- + Risk Management Plan: Development and implementation of comprehensive plans to proactively identify, assess, and mitigate risks associated with pharmaceutical products, ensuring patient safety and regulatory compliance.
- + Aggregate reporting: Robust generation of high-quality aggregate safety reports, including Periodic Safety Update Reports (PSURs) and Development Safety Update Reports (DSURs), to provide a comprehensive overview of product safety profiles.
- + Maintenance Of Eudravigilance Portal: Expert management and maintenance of the Eudravigilance database and reporting system, ensuring accurate and up-to-date data entry and compliance with regulatory requirements.
- SmPC review: Thorough review and analysis of Summary of Product Characteristics (SmPC) to ensure accurate and comprehensive information regarding product safety, efficacy, and administration, enabling informed healthcare decision-making.
- + Addendum To Clinical Overview (ADCO): Expert preparation of comprehensive addendums to clinical overviews, providing updated safety and efficacy information based on emerging data and regulatory guidelines.
- Medical Information Call Centre: Responsive and knowledgeable call center services, providing accurate and timely medical information to healthcare professionals and patients, ensuring appropriate use and understanding of pharmaceutical products.
- + **Signal Management:** Vigilant monitoring and analysis of safety signals, including the identification, evaluation, and communication of emerging safety concerns, promoting continuous improvement of product safety profiles.

REGULATORY AFFAIRS

- + **Regulatory Strategy Development:** Expert guidance and formulation of strategic plans for navigating complex regulatory landscapes, ensuring successful product development, approval, and commercialization.
- + Regulatory Submissions and Filings: Timely and accurate preparation, submission, and management of regulatory documentation, including Investigational New Drug (IND) applications, New Drug Applications (NDAs), and Marketing Authorization Applications (MAAs).
- + **Product Registration and Licensing:** Efficient management and facilitation of product registration and licensing processes, ensuring compliance with regulatory requirements and smooth market access.
- Quality Assurance and Compliance: Rigorous implementation of quality assurance systems and compliance measures to meet regulatory standards, ensuring product safety, efficacy, and adherence to Good Manufacturing Practices (GMP).
- + **Regulatory Intelligence and Updates:** Continuous monitoring and analysis of regulatory guidelines, policies, and updates to maintain compliance, anticipate changes, and adapt regulatory strategies accordingly.
- Interactions with Regulatory Authorities: Effective communication and collaboration with regulatory authorities, including pre-submission meetings, regulatory consultations, and regulatory agency interactions, to address inquiries, clarify requirements, and expedite the approval process.
- Food Supplement Regulatory Services: Specialized expertise in regulatory requirements and compliance for food supplements, including ingredient assessment, labeling, claims substantiation, and notification or registration processes.
- + Cross-functional Collaboration and Project Management: Seamless coordination and collaboration with crossfunctional teams, including R&D, manufacturing, marketing, and legal departments, to ensure alignment of regulatory strategies and efficient project management throughout the product lifecycle.

GSS PHARMA SERVICES

ARTWORK & LABELLING

- Artwork Design and Development: Creative and visually appealing design of pharmaceutical artwork, including
 packaging labels, leaflets, and product materials, ensuring compliance with regulatory requirements and brand
 guidelines.
- Regulatory Compliance Review: Thorough review and evaluation of artwork and labeling materials to ensure compliance with regulatory standards and guidelines, including accurate and complete information, proper formatting, and adherence to legal requirements.
- + Label Design and Layout: Expert design and layout of product labels, incorporating essential information such as product name, strength, dosage instructions, warnings, and regulatory markings, while maintaining clear and user-friendly presentation.
- Branded Packaging Solutions: Tailored packaging solutions that reflect brand identity and meet regulatory requirements, including selection of appropriate materials, colors, fonts, and design elements, ensuring product differentiation and compliance.
- + Patient Information Leaflets (PIL) Design: Clear and concise design of patient information leaflets, presenting important safety information, dosage instructions, and precautions in a user-friendly and easily understandable format, enhancing patient compliance and safety.
- + **Barcoding and Serialization:** Implementation of barcode and serialization systems to enable accurate product identification, tracking, and supply chain management, ensuring regulatory compliance and safeguarding against counterfeiting.
- + **Product Identification and Tracking:** Development and implementation of systems for product identification, tracking, and traceability throughout the supply chain, promoting regulatory compliance, product integrity, and patient safety.
- + Artwork Management and Version Control: Efficient management and control of artwork files, including version control, storage, and retrieval, ensuring accurate and up-to-date artwork materials for regulatory compliance and production purposes.
- + **Regulatory Guideline Adherence:** Strict adherence to regulatory guidelines and requirements throughout the artwork design, development, and approval process, ensuring compliance, consistency, and adherence to legal and regulatory standards.





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