Bharat Parenterals Limited is a specialized pharmaceutical manufacturing organization that has built a reputation for excellence based on its manufacturing & operational expertise, formulation innovation and high quality products. Our state-of-the-art manufacturing & formulation R&D infrastructure developed over an area of 3,00,000 sq. ft. offers finished dose formulations for emerging & semi-regulated global markets.

Facility Highlights

- 3 Independent & dedicated production blocks: General, Beta Lactam & Cephalosporin
- Separate air handling units for each section for air conditioning & dehumidification as per product requirements
- Our Integrated Formulation & Development Department has been recognized by the Department of Scientific & Industrial Research (DSIR)
- More than 30% manufacturing activities are powered by renewable energy sources such as solar & wind energy

Future Expansion

- Infrastructure for future expansion & development of other specially products such as lines for
 - Lyophilized
- Pre-filled syringes
- Emulsion formulations

Regulatorily approved Manufacturing & Formulation R&D infrastructure ISO 10002-2018 DSIR WHO GMP AEO-TI CERTIFIED TWO STAR EXPORT HOUSE







Manufacturing Excellence in wide range of Formulations

Three dedicated production units: General unit, Beta Lactam unit & Cephalosporin unit ensures zero cross-contamination. Our large batch size production capacity and 3+ decades of operational excellence enables us to cater to even unprecedented business demands.







Global Presence

BPL enjoys long-standing presence and commercial relationships in high-growth geographies. We are the trusted partner of 60+ clients exporting to over 40 countries.





Formulation Development & Analytical Development Strengths

Our integrated F&D Laboratory was recognized by the Department of Scientific and Industrial Research (DSIR), Govt. of India in the year 2016. With an emphasis on NDDS, our team of scientists are engaged in various areas such as method development, validation, formulation of novel dosage forms & improving existing formulations. Each year we invest in formulation research with an expertise in development of:

- Modified Release Tablets
- Extended-Release Dosage Forms
- Gastro-Retentive Dosage Forms
- Pulsatile Dosage Forms
- Lyophilized Injectables

Regulatory Affairs

Our dedicated team gives strategic and technical assistance to the company, from medicine development to effective product marketing by keeping track of a dynamic global legislative landscape.

- Number of CTD Dossier: 200
- Product Filings: 450 +
- Markets: Semi-regulated & ROW regions
- Focus Therapies : Anaesthesia & Pain management, Cardiovascular & CNS, Anti-malaria & Antibiotics

Quality Control & Quality Assurance

Well established Quality Systems with essential IP/QC (in-process quality control) process. Our in-house integrated quality control & testing laboratory is compliant with GLP standards, & equipped with high quality testing instruments that produce consistently reliable results.





Core Capabilities & Competitive Advantage

- Contract development & GMP manufacturing expertise for wide range of formulations
- Full range of integrated services including formulation development, analytical development & commercial manufacturing
- With 450 + product filings, our Regulatory Affairs teams are well versed with complex compliance requirements of various countries

BPL Group of Companies





Marketing BPL certified quality pharmaceuticals as branded generics in highly competitive Indian domestic landscape through highly skilled sales professionals





India's first-of-it's kind facility for development & manufacturing of complex/specialty drugs for US / Europe markets in niche areas of oncology, Alzheimer's and pain



Varenyam Biolifesciences Limited

Poised to leverage complex manufacturing capabilities & market access to further expand presence in high margin, emerging & regulated markets

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