neuland

Your CDMO Partner for Small Molecules and Peptides

Founded in 1984, Neuland Labs is a globally recognised Contract Development and Manufacturing Organization (CDMO), offering comprehensive customised NCE and peptide development services. Our team comprises of highly skilled scientists, engineers, and technicians who utilise their expertise and advanced technologies to address the complex and evolving requirements of our clients. We collaborate with pharmaceutical and biotechnology companies to support all aspects of their chemistry needs, ranging from early development stages to commercial manufacturing, including process development, optimisation, analytical testing, and regulatory compliance.

Neuland Labs possesses the capabilities and flexibility to manage both single-step processes and complete synthesis, handling projects from lab to commercial supply. We provide quantities suitable for clinical trials as well as full commercial-scale production of new chemical entities (NCEs), key starting materials (KSMs), active pharmaceutical ingredients (APIs), and their intermediates, ensuring efficient tech transfer timelines.

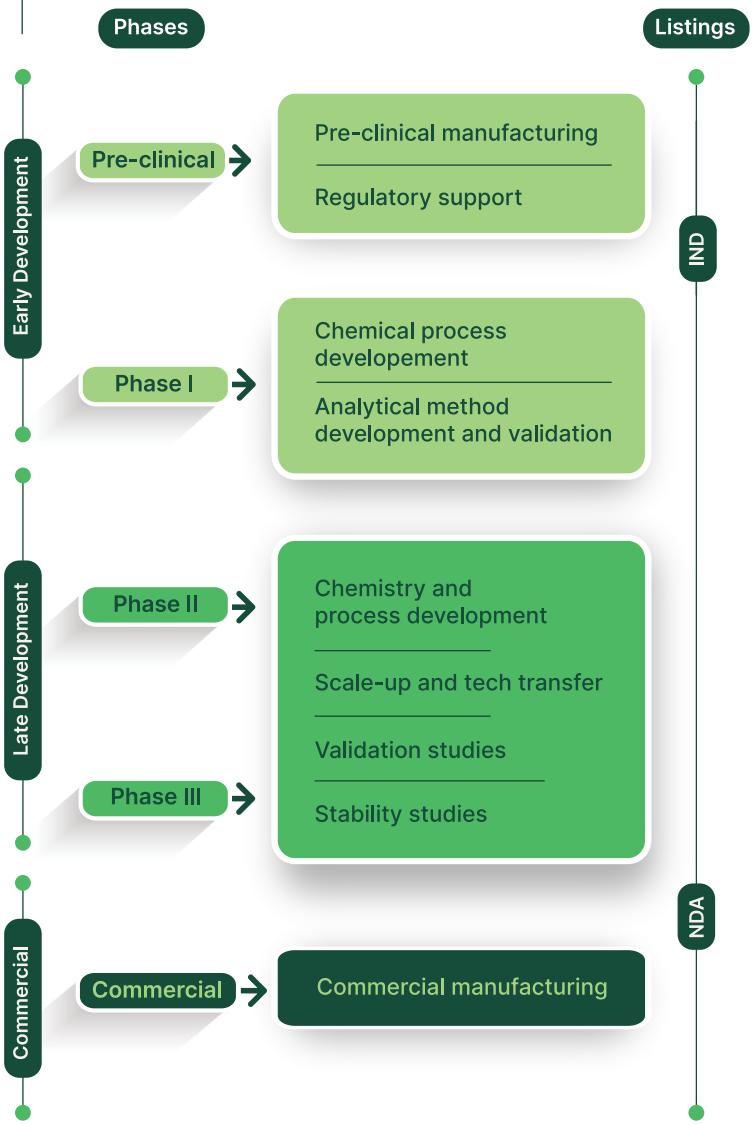
Our CDMO Services

- Custom NCE design and development
- Process optimization and validation
- Custom cGMP manufacturing - clinical production to commercial
- Analytical development and validation
- Process safety and engineering
- IND & NDA filings support

Enabling Services

- Quality Control & Assurance
- Project Management
- Supply Chain Management
- Regulatory CMC Services
- Intellectual Property

We provide phase-appropriate solutions across the entire drug development cycle



Experience the Neuland Difference

Proven Track Record
Over 4 decades of experience in small molecule API development

Customer-Centric Culture
Delivering innovative solutions with agility and a collaborative approach in project execution

Scalable Infrastructure
3 cGMP manufacturing facilities with 1174 kL reactor volume

Compliance & Integrity
Transparent documentation and full traceability

Clinical & Commercial Readiness
150+ NCEs projects successfully executed

18+ commercial APIs / Intermediates programs with global innovator companies

Quality Excellence
Strong history of 17+ successful USDFA audits

SILVER | Top 15% ecovadis Sustainability Rating MAR 2025

S&P Global Ratings

BRITISH SAFETY COUNCIL Five Star Occupational Health and Safety Audit 2024

UN Global Compact United Nations Global Compact

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