

Evonik Rexim - Your CDMO

Downstream capabilities and manufacturing expertise for high-quality purified life science products



1

Purification

2

Crystallization

3

European GMP expertise

Tailored CDMO Solutions for pharmaceutical and food products

Evonik Rexim is a European CDMO specializing in downstream processing of fermentation- and synthesis-derived small molecules. With decades of expertise in amino acids, peptides, polysaccharides, and specialty ingredients, we help customers transform intermediates into high-purity, market-ready products while accelerating development and commercialization.

OUR CAPABILITIES

- Processing of fermentation- and synthesis-derived molecules
- Chemical modification, including acetylation and hydrolysis
- Ion exchange chromatography and industrial-scale purification
- Ultrafiltration for bacterial and endotoxin removal
- Crystallization and centrifugation
- Evaporation, drying and spray drying
- Milling
- GMP-compliant packaging and supply

YOUR BENEFITS

- High-purity finished products
- Accelerated path to commercialization
- Reduced investment in internal downstream processing capabilities
- Flexible scale-up from development to commercial supply
- Reliable European partner with extensive industry expertise
- Decades of experience in amino acids, peptides, polysaccharides, and specialty ingredients

**WANT TO LEARN MORE
ABOUT OUR
CDMO CAPABILITIES?**

CONTACT OUR CDMO REPRESENTATIVE
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CHECK OUR WEBPAGE ►



Certifications

- GMP compliance (ICH Q7)
- ISO certification 9001, 14001, 50001
- FSSC 22000
- USP, EP, JP monograph compliant
- BSE/TSE Certificates
- High-purity chemical grade; controlled microbial/endotoxin levels
- Free from key contaminants and hazardous residues*
- Kosher and halal certifications
- Statement on allergens, residual solvents and Bovine Spongiform
- Non-animal origin

Registrations and Regulatory Affairs Support

- International Regulatory Agencies (FDA, PMDA and ANSM example compliance)
REACH registration
- Expertise in global registration processes and regional regulatory requirements
- Proactive implementation of new regulations
- Creation and maintenance of DMF and CEP
- Regulatory affairs support and consultancy

HACCP: Hazard Analysis and Critical Control Point Concept; BSE: Encephalopathy/TSE: Transmissible Bovine Encephalopathy available; USP: United State Pharmacopoeia, EP European Pharmacopoeia, JP: Japanese Pharmacopoeia; DMF: Drug Master Files , CEP: Certificates of Sustainability

*Free from chlorinated solvents, preservatives, antioxidants, mycotoxins, aflatoxins, pesticides, hormones, dioxins, and cytotoxic compounds

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