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PEPTIDE GENERIC API

Generic API	CEP/DMF	Application	Site
Atosiban	DMF	Reproductive Medicine	СН
Bivalirudin	DMF	Cardiovascular Disease	СН
Buserelin	DMF	Oncology, Reproductive Medicine	СН
Desmopressin Acetate	CEP, DMF	Diabetes Insipidus	USA
Exenatide Acetate	DMF	Diabetes Mellitus	USA
Glucagon	DMF	Diabetes Mellitus	СН
Gonadorelin Acetate	CEP, DMF	Oncology, Reproductive Medicine	СН
Goserelin Acetate	CEP, DMF	Oncology, Reproductive Medicine	СН
Icatibant Acetate	DMF	Hereditary Angioedema	СН
Lanreotide	DMF	Oncology	USA
Leuprolide Acetate	CEP, DMF	Oncology	СН
Octreotide Acetate	CEP, DMF	Oncology	СН
Somatostatin	CEP	Gastritis	СН
Teriparatide Acetate / pTH (1-34) (human)	DMF	Osteoporosis	USA
Tetracosactide	DMF	Oncology, Diagnostics	СН
Triptorelin Acetate	DMF	Oncology, Reproductive Medicine	СН
Triptorelin Pamoate	DMF	Oncology, Reproductive Medicine	СН
(Arg ⁸)-Vasopressin	DMF	Diabetes Insipidus	USA
Liraglutide*	DMF	Diabetes Mellitus	USA/CH
Semaglutide*		Diabetes Mellitus	СН

SMALL MOLECULE API

CEP/DMF	Application	Site
DMF	Ophthalmology	СН
DMF	Ophthalmology	СН
DMF	Ophthalmology	СН
CEP, DMF	Epilepsy and Parkinson's Disease	СН
CEP, DMF	Sedatives and Anesthetics	СН
CEP, DMF	Sedatives and Anesthetics	СН
	DMF DMF DMF CEP, DMF CEP, DMF	DMFOphthalmologyDMFOphthalmologyDMFOphthalmologyDMFEpilepsy and Parkinson's DiseaseCEP, DMFEpilepsy and Anesthetics

PIPELINE

Application
Oncology
Short Bowel Syndrome
Short Bowel Syndrome
Diabetes Mellitus
IBS, Chronic constipation
Obesity

Please note: Some products may be restricted in certain countries.

* Bachem provides this product solely for uses within the scope of any statute or law providing for an immunity, exemption, or exception to patent infringement ("Exempted Uses"), including but not limited to 35 U.S.C. S 271(e)(1) in the United States, the Bolar type exemption in Europe, and any corresponding exception to patent infringement in any other country. It is the sole responsibility of the purchaser or user of this product, and the purchaser or user of this product agrees to engage only in such Exempted Uses, and to comply with all applicable intellectual property laws and/or regulations. The purchaser of this product agrees to indemnify Bachem against all claims in connection with the performance of the respective commercial agreement (e.g. supply agreement) and possible infringements of intellectual property rights.

FULL RANGE

of technical and regulatory support

- We enable clients to meet the requirements of the authorities by providing comprehensive technical and regulatory information for filing.
- Our service includes the compilation of technical and regulatory documents and we offer support for regulatory requests.
- Through our expertise we enable clients to be the first to file their generic drugs in a highly competitive market.



EXPERIENCE

- Bachem is the leading independent supplier of complex active pharmaceutical ingredients (APIs) for the human and veterinary pharmaceutical market.
- We have over 50 years of experience and a strong track record in the production and filing of peptide and small molecule drug substances.
- Building on our heritage, we pioneer innovations to deliver the best quality for every API need at competitive pricing.

QUALITY

- Bachem has facilities in Switzerland and in the US, approved by international regulatory authorities.
- We produce active ingredients from milligrams of research material for preclinical work up to hundreds of kilograms of API.
- Inspections by national authorities and more than 50 customer audits per year confirm our high quality standard and our compliance with Good Manufacturing Practice (cGMP) regulations.

PARTNERSHIP

- With our commitment to supply products of the highest quality, we have established long-term contracts with important industrial partners.
- Our partners count on Bachem's dedicated, well-trained and responsible project managers, realizing supply solutions tailored to their needs.
- The tight execution of projects and our customer-friendly service set us apart.

RELIABILITY

- We follow sound economic and ecological principles and comply with relevant international regulations.
- Our customers rely on the constant quality of our generic APIs for the manufacturing of their drug products.
- Your trust in Bachem is built on our reliability and our willingness to go the extra mile for you.

- Over 50 years of experience
- Strong track record in the successful production and filing of APIs
- Tight execution of projects and customer-friendly service
- In compliance with sound economic and ecological principles

BACHEM

GLOBAL BUSINESS

Bachem facilities are located in Switzerland, the UK, and in the US. All cGMP manufacturing sites are inspected by the US-FDA and national authorities.







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