With Chiracon to new entities and APIs

www.chiracon.de

From structure to drug substance

Chiracon is the independent and reliable service partner for the pharmaceutical and biotech industry. Our certified laboratories and production facilities are located at the Biotechnology Park in Luckenwalde. Since its foundation, Chiracon has successfully completed over 330 projects, including over 30 projects for active pharmaceutical ingredients at various stages of development. We are currently producing for numerous clinical projects.

| CONTACT

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chiracon

SERVICES

Chiracon has been your API manufacturer in Brandenburg for over 25 years

www.chiracon.de

APIs

API in GMP quality for industry and pharmacies

Active pharmaceutical ingredients (APIs) are manufactured in compliance with applicable standards and regulations (GMP, AMG, AMWHV, etc.). By manufacturing APIs in Germany, our customers receive a high quality promise and security of supply for their product.

The results of the entire process, from research and development to scaling and production, are monitored, controlled, documented and continuously improved by the quality control and quality assurance departments to ensure consistently high quality.

CONTRACT RESEARCH

The development of a modern manufacturing process for your API

For over 25 years, Chiracon has been offering know-how in the development, optimisation and scaling of synthesis of active pharmaceutical ingredients as well as in the manufacture of customised products. We contribute our expertise to numerous collaborative projects, including those funded by third parties.

SCALE-UP

We optimise and scale up your synthesis process

We offer you the scaling of your production processes, usually up to 100 kg of product per year. Larger delivery quantities are also possible by arrangement. Our scaling projects are of course also carried out under ISO-certified conditions. Your process can also be GMP-certified and documented accordingly, if required.

INTERMEDIATES

High-quality raw materials for pharmaceutical and technical applications

One of Chiracon's core competences is the synthesis development and production of enantiomerically pure compounds and intermediates, including for pharmaceutical applications. Both classical chemical and enzymatic synthesis methods are utilised. Our synthesis projects are carried out under ISO-certified conditions. If required, the processes can be GMP-certified and documented accordingly.

STABILITY STUDIES

Stability studies according to ICH guidelines

Chiracon offers stability studies in accordance with ICH guidelines (incl. IVb, Brazil) to accompany drug synthesis development or as a service for your products. On request, we carry out the analytical characterisation of decomposition products and, if required, the synthesis of the decomposition products.

Our stability studies are of course also carried out under ISO-certified conditions. If required, the procedures can be GMP-certified and documented.

SAMPLES FOR CLINICAL TRIALS

GMP production of clinical trial samples

From preclinical testing to authorisation and then later commercial production: make Chiracon your partner and let us work together to bring your project to market maturity. Regardless of whether you already have a synthesis process or want us to carry out the process development for you, Chiracon is your reliable partner all the way.

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