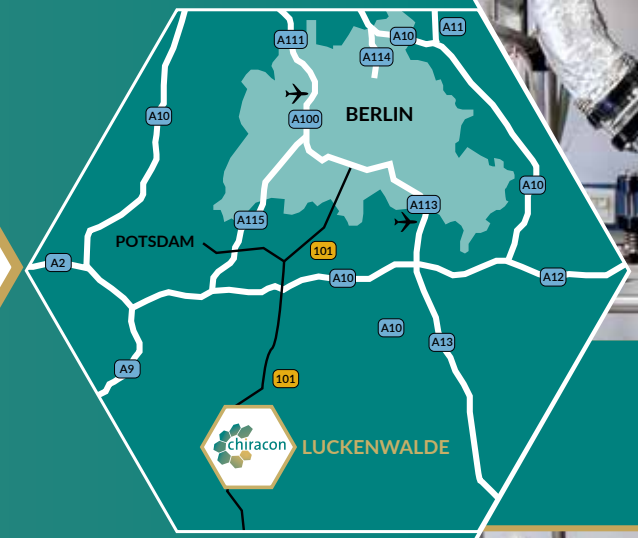
Active Pharmaceutical Ingredients (APIs) for the Pharmaceutical Industry

From preclinical testing to approval, and later to commercial production: Make Chiracon your chemical project partner, and together we can bring your project of choice to market. No matter whether you already have a method of synthesis or you would like us to develop the complete synthesis, contact us for a fast and detailed offer.

With Chiracon your project is in good hands. We have 20 years of experience specializing in the synthesis of small molecules. Chiracon provides all the necessary analytical and documentary services concerning the regulatory aspects of drug development. We have experience with all types of syntheses and APIs for all applications for oral and dermal administration. Our versatile production facilities and our highly qualified staff make us your partner of choice for all requirements of the project process. Approach us for your individual offer meeting your specific requirements in your needed quality, and all this at favorable pricing.

We cooperate directly with drug formulation, processing, and toxicological working industry partners to bring your project directly to the clinical phase without any time lag.

Chiracon is an API manufacturer that has been inspected in accordance with – Art. 111 (1) of Directive 2001/83/EC transposed in the following national legislation: Sect. 64 para 1 Arzneimittelgesetz (German Drug Law) (GMP certificate).



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Wirkstoffe • Klinische Prüfmuster • Chirale Intermediate • Auftragssynthese

chiracon

From Structure to Drug Substance

Active Pharmaceutical Ingredients

Active pharmaceutical ingredients according to GMP guidelines for the pharmaceutical industry and for formulations for pharmacies and hospitals

APIs for Clinical Trials

Synthesis of APIs for clinical trials according to GMP guidelines including drafting the respective documentation (IMPD)

Analytical Support

Chiracon offers a whole range of analytical services for pharmaceutical research and production. Through our expertise in drug development and manufacturing we are able to provide analytical services that meet your needs and requirements.

Thanks to our pharmaceutical analysis specialists who have many years of experience in developing, optimizing and validating methods in line with ICH guidelines we can assure fast, flexible and accurate solutions through a broad range of services:

- *method development, validation and transfer*
- *analytical chemistry for raw materials, bulk and finished products*
- *forced degradation, stability testing and storage (according to ICH guidelines and customer specific needs)*

As a GMP-certified laboratory, we conduct all the relevant tests in line with international pharmacopoeia. The analytical focus of Chiracon is on product purity using state of the art equipment including HPLC, GC, HS-GC, IR, NMR spectroscopy, isolation and synthesis of degradation products in drug substances and products.

Chiracon gladly integrates your methods or develops methods geared to your specific needs. We conclude technical and confidentiality agreements with our customers to ensure the necessary legal framework is in place.

Chiral Intermediates

Selected chiral intermediates for synthesis and industrial applications

Custom Synthesis

Development of synthesis and scale-up for research projects or preclinical studies