tbd PHARMATECH

Your CDMO based in the EU

End-to-end service

TBD's skilled in-house team handles every aspect of your API development and manufacturing needs, from route scouting, in-house QC and regulatory support to commercial production.

We deliver

a tailored range of fine chemical development and manufacturing solutions to pharmaceutical, biotech and chemical companies for ~20 years.

A reliable partner

Long cooperation with aknowledged European pharmaceutical companies. Over 80% of new projects come from existing partners.

CONTRACT DEVELOPMENT

- **API DEVELOPMENT**
  Generic and innovative molecules
- **EARLY PHASE CLINICAL API DEVELOPMENT**
  Producing clinical batches for innovative APIs

Focus on quality

Our operations follow stringent quality management principles to uphold standard. TBD has been GMP-compliant since 2008 and certified to ISO 9001:2015

CERTIFIED ISO 9001

GOOD MANUFACTURING PRACTICE GMP

CONTRACT MANUFACTURING

Originate from completed in-house development or through customer-led tech transfer.

Outstanding track record

< 30 APIs developed for human and veterinary pharma for Phase I/II and generics under several categories.

>10 ASMFs submitted in EU, Switzerland, UK, Canada, Australia etc.

OTHER SERVICES

- **Conjugation chemistry**
  To improve drug attributes or create specialized treatments like ADC conjugates.
- **Custom synthesis and analytics of specialty compounds**
  Tailored solutions for generating reference materials, excipients, intermediates and impurities.
- **Regulatory support**
  Prepare and submit key regulatory documents like DMF, ASMF, CMC, to expedite approvals from authorities.

> *TBD offers a unique level of expertise in API synthesis combined with flexibility, open communication as well as speed in all activities.*
>
> Tiefenbacher Group

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