



Integrated CDMO Services for Smarter Drug Development

HWI group is an independent, family-owned service provider to the pharmaceutical and biotech industries. We support pharmaceutical and biotech companies with our end-to-end service concept throughout the entire drug development process.

Combining formulation development, analytical expertise, manufacturing capabilities as well as regulatory support, we provide tailored CDMO services from early-stage development to clinical trial material production.

CMC Development

- Development of new drug products with focus on complex small and large molecules e.g. poorly soluble APIs
- Optimisation and reformulation of existing products
- Development of herbal medicinal products

Dosage Forms and Packaging

- **Liquid dosage forms:** Solutions, micro- and nanosuspensions, emulsions and lyophilisates for oral, nasal and parenteral use
- **Semi-solid dosage forms:** Creams, oleogels, hydrogels, emulsions and self-emulsifying systems (SEDDS, SMEDDs) for topical applications
- **Solid dosage forms:** Tablets, coated tablets, mini tablets, hard capsules, pellets, granules and powders for oral use
- **Primary packaging of liquid and semi-solid dosage forms:** Vials, syringes, cartridges, bottles, tubes
- **Primary packaging of solid dosage forms:** Bottles, blister packs, containers, sticks, pouches

Manufacture of CTM and Small Commercial Batches

- Solid, semi-solid and liquid dosage forms
- Micro- and nanosuspensions, nano emulsions
- Nasal sprays
- Handling of APIs up to OEB 5 and narcotic drugs
- Tailor-made primary and secondary packaging incl. cold chain
- Customised labelling, serialisation and aggregation
- Patient kit labelling and assembly
- Worldwide distribution to clinical trial sites under controlled conditions
- Comparator sourcing
- IMPD Writing
- Import/export and customs clearance

Integrated Services

- Analytical development and validation
- Release and stability testing
- QP batch release and regulatory support
- Nanotechnology-based formulation approaches to enhance bioavailability



Sterile Fill & Finish of Liquid and Lyophilised Products

Sterile Powder Development and Filling

- Powder formulation development and the transfer from laboratory to production scale
- Filling of powders for terminal sterilisation
- Manufacture of technical batches, stability batches, and batches for clinical trials and commercial use with batch sizes of up to approx. 10,000 units.

New Service from 2029: Steril Fill & Finish

- Sterile Fill & Finish of Liquid and Lyophilised Products
- Manufacture of sterile finished products to medium sized batches of up to 30,000 items for large and small molecules
- Filling in vials, syringes, and cartridges.
- **The new facility is projected to start operations in 2029**



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As an end-to-end service provider, HWI complements its portfolio with laboratory services, pharmacovigilance and QP services as well as regulatory support, all from a single source.

Whether you are looking for short-term expertise or a long-term strategic partner, HWI delivers tailored solutions to meet your specific requirements.

Get in touch with us to find out how we can support your project.

HWI group

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