Navigating you through drug development

We are Ardena, a fully integrated Contract Development & Manufacturing Organization (CDMO) with a core focus on bringing molecules to the clinics.

We operate from six state-of-the-art, purpose-built facilities across Western and Northern Europe.

We service a broad and global customer base including prominent blue-chip companies. We have an outstanding quality and regulatory track-record, verified by full GMP and GLP compliance.

Get in touch with our deligates

Matteo Seregni
Business Development Director
Solid State Research

Dave Seghers
Business Development Director
Drug Development & Manufacturing

Hilbert Branderhorst
Business Development Director
API & Nanomedicine

Seven centers of excellence to support your multidisciplinary project

Drug Substance Development & Manufacturing
- Route scouting
- DS process dev
- GMP manufacturing
- Analytical method dev
- Stability testing

Drug Product Development & Manufacturing
- Preformulation
- Formulation & process development
- Solubility enhancement
- Analytical method development and validation
- GMP manufacture
- Stability studies

Solid State Research
- Polymorph screening
- Salt screening
- Co-crystal screening
- Crystallisation process dev
- IP support

API/ Nanomedicines
- (Pre)formulation dev
- DP process dev
- GMP manufacturing
- Analytical method dev
- Stabilty testing

Clinical Supply
- GMP Packaging & labeling
- Patient kits
- Clinical logistics (DP+biosamples)

Bioanalysis
- Discovery support
- Bioanalytical method dev
- Preclinical bioanalysis
- Clinical bioanalysis
- Biomarkers

Regulatory Dossier Development
- Data management
- Scientific report writing
- Quality document writing
- Regulatory dossier writing
- Dossier submission and maintenance