

# Taking pride in putting science at the heart of serving customers to **bring better, safer products to market faster**

## Who We Are

Solvias is a world leading CRO/CDMO headquartered in Switzerland. **Our highly trained scientists deliver unrivaled expertise, service, and value to customers in the pharmaceutical and biotech industry.**

Drawing on a proven 20+-year track record of scientific excellence, we provide cGMP-compliant contract analytical services to help you bring safe and effective therapies to patients faster. Partner with us to gain access to a wide range of capabilities for characterization of biologics, ATMPs and small molecule drugs and method development, release testing and stability services. We also offer one of the largest ligand portfolios for catalytic transformations and a suite of related custom API synthesis and catalysis technology services.

Our experts use a phase-appropriate analytical testing approach to help you navigate the regulatory environment. With ready access to our subject matter experts and professional project management **we are invested in your success.**



**800+ FTEs**  
**175+ PhD-level scientists**



**GMP, GLP and ISO9001 certified**



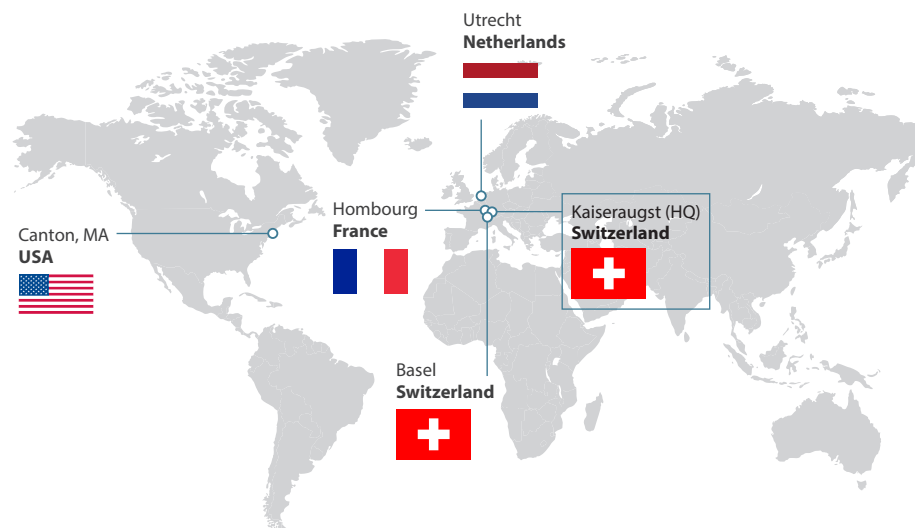
**700+ current customers**



**Successful FDA inspection (2023)**

## High Quality Compliance

- 22.5K m<sup>2</sup> of lab capacity
- 100+ successful customer audits per year
- Five sites in global biopharma hubs



## From Our Customers

“

Whatever they do, is **fully compliant with regulations**. I can trust them on quality control, which is very important in the analytical testing space.

”

“

They're not a new player, they had time to develop **an approach to analytical testing that makes them unique**. They have also worked with big pharma companies before, which makes the collaboration easier.”

”

“

There is no doubt that they are **very highly technologically capable**, both in terms of their methodology and in their staff.

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# Analytical Solutions that Span the Entire Product Lifecycle

	Preclinical	Phase I	Phase II	Phase III	Commercial
<b>Synthesis</b>	[Dark Blue Bar]				
Custom Synthesis	[Light Blue Bar]				
API Synthesis		[Light Blue Bar]			
Process R&D		[Light Blue Bar]			
Ligand Supply	[Light Blue Bar]				
<b>Solid State</b>	[Dark Blue Bar]				[Light Blue Bar]
Salt/Co-Crystal Selection	[Light Blue Bar]				
Polymorph Selection		[Light Blue Bar]			
Crystallization Development			[Light Blue Bar]		
<b>Analytical</b>	[Dark Blue Bar]				
Characterization	[Light Blue Bar]				
Method Development		[Light Blue Bar]			
Method Validation		[Light Blue Bar]			
Functional Analysis	[Light Blue Bar]				
<b>Quality Control</b>		[Dark Blue Bar]			
<b>Stability Studies</b>	[Dark Blue Bar]				
Forced Degradation	[Light Blue Bar]				
ICH Stability		[Light Blue Bar]			
Post Approval Studies				[Light Blue Bar]	
<b>Troubleshooting</b>	[Dark Blue Bar]				

Small molecules

Biopharmaceuticals

## World-Class Facilities

	 Kaiseraugst, Switzerland (HQ)	 Hombourg, France	 Basel, Switzerland	 Canton, MA United States	 Utrecht, Netherlands
<b>Size</b>	~12,540 m <sup>2</sup>	~3200 m <sup>2</sup>	~2790 m <sup>2</sup>	~2090 m <sup>2</sup>	~230 m <sup>2</sup>
<b>Employees</b>	~489	~214	~45	~47	~38
<b>Quality Accreditation</b>	GMP, ISO 9001	GMP, ISO 9001, ISO 17025	GMP, ISO 9001	GMP, GLP	ISO 17025
<b>Activities</b>	Comprehensive GMP analytical services Biopharmaceuticals and small molecule Drug substance and drug product	Microbiology Molecular biology Cellular biology Virology	Ligand and specialty catalyst production Related synthesis and catalysis services	GMP analytical services E&L programs Small molecule API manufacturing services	Large molecule ATMP testing Biologics research and development

### Contact us to speak with an expert

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