

# SAFC®

Pharma & Biopharma Raw  
Material Solutions

# MERCK



# Complex Small Molecule Manufacturing

## Schaffhausen Facility Overview

As a development, launch, and commercial supply site for APIs and lipids as well as activated PEGs, our cGMP facility in Schaffhausen (Switzerland) offers production capacities for custom synthesis from grams to tons. We are the world's leading supplier of reduced folates and have more than 50 years of experience in pre-clinical to commercial supply. Our tailored approach enables our customers to move their products to market quickly and efficiently.

## Chemistry Development Services

Specialized in lipids, PEGs, reduced folates and other APIs, our comprehensive chemistry development services include:

- Route finding
- Chemical feasibility studies
- Process characterization
- Process optimization and full development including QbD and DOE
- Scale-up for kilo lab and commercial scale up to 60 L
- Process Safety evaluation
- Impurity characterization and synthesis

## Analytical Services

We offer comprehensive analytical support for all our cGMP operations from raw material quality control (QC) to final product release.

- Thorough chemical analytical testing
- Identification of impurities and by-products
- Microbio lab for bioburden and endotoxin testing
- Development of analytical and microbiological methods

- Method validation according to ICH Q7
- Stability studies according to ICH Q7 with internal storage capacity
- Direct support by our technical teams
- SAP QM as integrated LIMS with automatic data transfer

### General applicable methods

- UPLC (UV/RI/ELSD/CAD/MSD)
- GC/GC-MS/MALDI-TOF
- DSC/TG
- IR/UV/Optical rotation
- Wet chemistry (Titrations)
- CE/Ion Chromatography
- TOC
- Microbiological testing

### Special methods

- XRPD – differentiation of polymorphic forms
- NMR – structure elucidation
- SEC – Mw distribution

## Manufacturing

Schaffhausen provides APIs and key intermediates (drug delivery compounds) out of state-of-the-art cGMP facilities for use in clinical trials and commercial product applications.

### Equipment Overview

QTY	Equipment	Capacity	Temp. Range
11	Glass line reactors	250 – 630 L (nominal vol.)	-40°C – 200°C
3	Hastelloy® reactors	250 – 630 L	-20°C – 160°C
2	Hastelloy® centrifuges	800 mm (diameter)	-25°C – 100°C
1	Inverted bag centrifuge	450 mm	-10°C – 100°C
1	Hastelloy® double cone dryer	750 L (300 L product)	-30°C – 200°C
1	Titanium filter dryer	280 L	-20°C – 135°C
2	VA Shelf dryers	1150 – 2050 L	25°C – 100°C
2	VA centrifuges	800 mm (diameter)	-25°C – 100°C
1	VA filter dryer	250 L	-10°C – 100°C
2	Mobile vacuum contact dryers	160 L (nominal value)	25°C – 80°C

### Clean Rooms

- 5 ISO-8 clean rooms for dispensing active materials

### Other Features

- Endotoxin controlled purified water system

To place an order or receive technical assistance in Europe, please call Customer Service:  
 France: 0825 045 645 Spain: 901 516 645 Option 1  
 Germany: 069 86798021 Switzerland: 0848 645 645  
 Italy: 848 845 645 United Kingdom: 0870 900 4645

For other countries across Europe, please call: +44 (0) 115 943 0840  
 Or visit: [MerckMillipore.com/offices](http://MerckMillipore.com/offices)  
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## Technologies

- Aromatic chemistry
- Protected amino acids
- Esterification
- Alkylation
- Heterocyclic chemistry
- Pressure reactions
- Oxidation
- Reduction
- Chiral synthesis
- Chiral resolution
- Hydrogenation

## Compliance, Regulatory & EHS

Schaffhausen complies with the quality systems and standards of Merck KGaA, Darmstadt, Germany and has been inspected regularly by various regulatory authorities including the FDA (USA) and our principal regulator, the Swissmedic (Switzerland). A dedicated team of experts ensures full regulatory support/documentation:

- Global expertise in API registration
- Intellectual Property (IP) and Freedom-to-Operate (FTO) evaluation
- Proven track record of successful DMF submissions globally
  - Preparation of regulatory filings (NDAs)
  - 150+ customer audits over the past 10 years
- Excellent FDA inspection track record (no Form 483 issued)
- Quality systems cGMP/ICH Q7
- Compliant with US-FDA generic drug user fee act (GDUFA)
- Annual Kosher inspections
- Annual Halal inspections
- License for handling controlled substances

## Dedicated Experts

### Research & Development

Our team of dedicated scientists and highly skilled experts support the development of innovative APIs, activated PEGs, PEG-hydrogels and specialty lipids suitable for parenteral use. Our position as a large European manufacturer allows us to offer a full spectrum of services from reaction screening to commercial supply.

### Project Management

From evaluation to execution, our dedicated project managers are coordinating multidisciplinary teams, international site activities and timelines throughout the lifecycle of your program.

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