



BACCINEX

your  fill & finish expert

CMO Pharmaceutical manufacturer

*Aseptic filling of sterile liquid and
lyophilized dosage forms*

Location



Rue de la source, 3
CH-2822 COURROUX
info@baccinex.com



[Web link: Baccinex – Your Swiss Fill and Finish Expert](#)



History

1999
Foundation of Baccinex SA

2003-2004
Installation, qualification and validation activities

2004
Certification by Swissmedic

2021
Start of the project of building
of a new warehouse

Q4 2024
Revamping of the filling line

Mission

Production of vials for

- › Clinical batches → for worldwide clients
- › Commercial batches → for Europe + US
- › Technical batches
 - for preclinical studies
 - for stability studies
 - for technological transfer

Mission

One stop fill&finish partner :

› For clinical manufacturing



› For commercial manufacturing



Mission

Client-oriented approach and specific strengths related

- › Utmost flexibility
- › Strong project management system for a proactive follow-up of the projects
- › Technical and GMP regulatory support related to the clinical project
- › Extensive expertise in management of limited quantity of valuable API
- › Standard APS available
- › No minimum batch size

Strategy

Pure CDMO = Multi product facility managed by :

- › Risk assessment to determine the feasibility of the project at Baccinex
- › No handling of products requiring dedicated facilities (β -lactam antibiotics...), nor alive microorganisms

Strategy

Multi product facility managed by :

- › Standard APS (covering most common worst cases) for automatic filling of vials
- › Standard visual inspection qualification for operators
- › Standard autoclave loads qualification (e.g.: mixed load)

Standard validated processes to be checked and accepted by clients during audit. If not in line with client regulatory understanding, specific process validation can be performed

Key figures

- › 55% of clinical batches / 45% of commercial batches
- › 40% of lyophilized products/60% of liquid products
- › 20 years of GMP experience
- › ~45 different manufacturing projects managed each year

Development services

- › Linked to scale-up and industrial transfer
 - › Filter/material compatibility
 - › Filter retention
- › Collaboration with companies specialized in formulation and lyophilization development

Manufacturing services

- › Formulation
- › Aseptic compounding
- › Aseptic filling of vials
- › Lyophilization
- › Integrity testing
- › Visual inspection
- › GMP Storage (15-25°C; 2-8°C; -15/-25°C; <-70°C)

Batch sizes

Dosage form	Liquid	Freeze-dried
VIALS		
Type	Maximum batch size	
DIN 2R	10'000	11'500
DIN 6R	6'200	6'100
DIN 10R	4'900	4'900
DIN 15R	4'900	4'900
DIN 20R	3'200	3'200
DIN 30R	2'500	2'500

Manufacturing services

> Labelling/secondary packaging :

Activity	Labelling	Secondary packaging
Clinical batches	Customized/ randomized labelling	Customized secondary packaging such as : <ul style="list-style-type: none">- Preparation of kits- Packaging for blinded clinical studies
Commercial batches	Automatic labelling	<ul style="list-style-type: none">- Secondary packaging of finished products (folding boxes, leaflets)- Serialization- Tertiary packaging

> Shipment

Collaboration with a qualified carrier specialized in temperature monitored shipment of pharmaceutical products

Quality Control services

Laboratory

Activities

Physico-chemic lab

HPLC-UV (assay, purity),
Spectrophotometry UV-Vis
Spectrophotometry IR
Sub-visible particle counting

pH

Residual water content (Karl Fischer volumetric or coulometric)

Viscosity

Osmolality

TOC

Potentiometry

Microbiological lab

Bioburden and specific germs contamination

Endotoxins (turbidimetric or chromogenic technics)

Sterility testing under isolator

Incubation and reading of APS at 20-25°C & 30-35°C during 7 days

Environment control in classified areas

Quality Control services

- › Validation of non compendial analytical methods / Suitability testing of microbiological methods :
 - › Bioburden
 - › Sterility
 - › Endotoxins
 - › HPLC etc.
- › **Stability study** : Stability storage according to ICH conditions
 - › 40°C/75%RH
 - › 30°C/65%RH
 - › 25°C/60%RH
 - › 5°C +/-3°C
 - › -20°C +/-5°C
 - › Analysis according to stability program

Quality Assurance

- › Inspection by Swissmedic (Swiss regulatory authorities every 2 years)
- › GMP confirmation of the batches :
 - › GMP confirmation established by Baccinex' QP
 - › Partnering with an European regulatory specialized company who performs EU release of batches
- › Around 20 external clients' audits / year
- › Quality system in place to ensure GMP compliance for any of performed activities

Project Management

Personalized and constant support through complete project lifecycle

- › Initial project assessment with client
- › Proactive dedicated Project Managers as interface between client and internal teams
- › Dynamic project team including the appropriate internal experts
- › Specific project management tools for identifying and establishing detailed format client needs
- › Recognized flexibility and strong anticipation skills on the overall project requirements
- › Adaptability to project and client constraints

Conclusion: Baccinex engagement



Contacts



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