The worlds leading CDMO for Microbiome therapeuthics and next generation probiotics.

Developing and producing Bacteria as a drug for over 70 years





commitment agility responsibility expertise







PROCESS DEVELOPMENT

Microbiotic candidates

Aerobic & anaerobic

Consortia & multi or single-strains

Commensal and delivery GMO

Post-biotic

R&D Microbiology Platform

Risk group assessment (Bioinformatics)
Strain Isolation & characterization: Screening,
Strain-specific ID (Molecular tools), Strain characterization
Analytical methods transfer development & validation

R&D Process Platform

Manufacturability screening and process development Fermenters: 0.5L, 2L, 5L, 20L, 150L
Centrifugation & TFF
Freeze-driers: $2 \times 0.5m^2 - 1.5m^2 - 4m^2$

Dosage form: Capsule, tablet, sticks, oil-suspension, cream

Powder & Product characterization

Comparability tests

commitment agility responsibility expertise

120 M€ invested to maintain our position as the world leading CDMO for microbiome therapeutics and next generation probiotics

GMP DRUG PRODUCTION

DRUG SUBSTANCE

Scale-up & manufacturing

GMP MCB and WCB manufacturing / according to LBP Guidelines and storage at -80°c & -20°c

DS1: 300L and 3 500L - Centrifugation & TFF + 25m² freeze-dryer

DS2: 2 000L - Concentration + 6m² freeze-dryer

DS3: 4 x 800L - 2x 30m² freeze-dryer

DS4: 150L - Centrifugation + 1m² freeze-dryer

DS5: 2 000L - Centrifugation + 12m² freeze-dryer

DS6:5000L-Centrifugation+30m² freeze-dryer

Commercial or clinical batches released by our QP (PharmD)

Powder placebo production for Capsule & Tablet & Sachet

DRUG PRODUCT

Formulation for Capsules / Tablets / Sachets / Oils & Creams

Blending from 10 to 400 kg

Dosage forms: Capsules, Tablets, Powder

Enteric Banding and Coating

Packaging: Blisters, Vials, Sachets, Tubes, Creams & Oils.







KEY FIGURES



- + 766 DS batches released in 2023
- + 318 DP batches released in 2023
- + 75 Tonnes of commercial drug substance 160 000 QC analyses done in 2023

52000m² centre of excellence + applied R&D laboratory in Boston, US.

30+ clinical lines including 3x phase III projects in Asia, Europe & the US for Microbiome therapeutics and next generation probiotics

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Main facility, R&D and GMP production - Aurillac France R&D and Pilot Plant - Boston USA

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