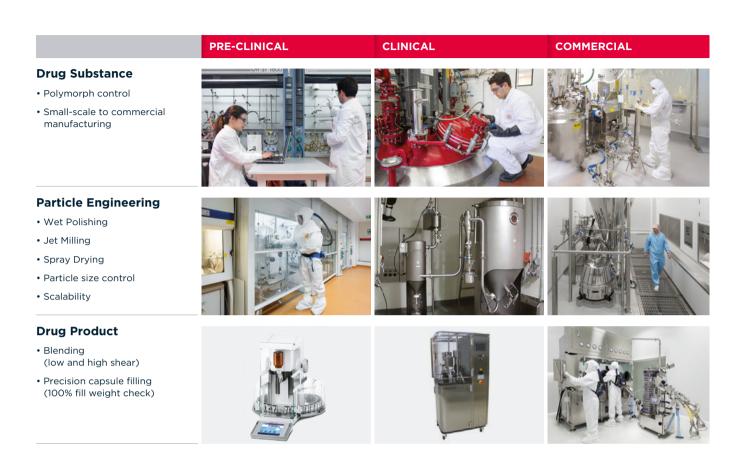


Everything for Inhalation



In it for life

Integrated Inhalation Development Services



Particle Engineering Technologies

Unprecedented particle size control

- Median particle size control down to 0.1 μm
- Spans down to 0.6
- At any scale

Advanced Engineered Particles for low and high dosages

- Microencapsulation
- Nanocoating

API alone

Control over interfacial properties

STRENGTHS

Jet Milling

- Simple
- Solvent free

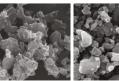
Wet Polishing

- Reduced amorphization
- PSD control
- Reproducibility
- Scalability

Spray Drying

- PSD control
- Reproducibility
- Scalability • Thermo-labile
- compounds

JET MILLED API





WP FP 2µm



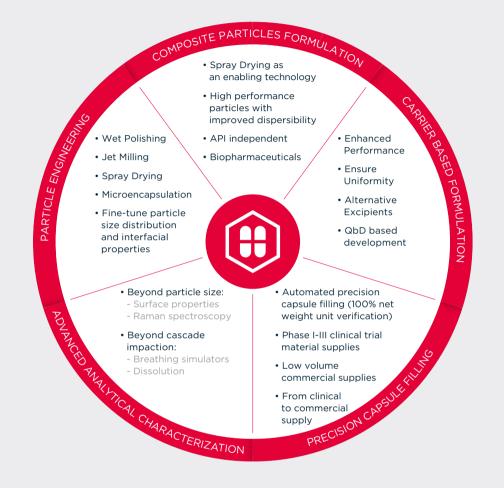
JM Fluticasone Propionate (FP) 2µm

JM Mometasone Furoate (MFA) 2µm

WET POLISHED API

WP MFA 2µm

Everything for Inhalation



Inhalation drug product development at Hovione has a strong focus on formulation for Dry Powder Inhalers (DPI), particularly for capsule-based and reservoir-based DPIs.

From formulation composition optimization and precision capsule filling with 100% fill weight check, to analytical characterization and scale-up to a commercial scale, Hovione can take your product from the early proof of concept stages up to a fully developed and scaled process. Together with our API and Particle Engineering capabilities, this means that your product can be fully developed and produced at Hovione, from API synthesis, to size reduction and final drug product.

Carrier-based Formulations

Blending

Most carrier-based formulations are lactose-based and they can carry different inhalation grades of lactose, typically a combination of coarse and fine lactose, and a range of API percentages. The lactose grades for your specific product will be carefully chosen to provide the best possible aerodynamic performance and stability through the life cycle of your product. With our low-shear and high-shear blenders (Turbula and Diosna), we are equipped to perform blend production from very small batches of only a couple grams up to a fully commercial scale on the kilogram range. Our cGMP blending equipment (Diosna P1-6 and Diosna PVAC 10-60) will allow production of blends from 150 g up to 15 kg (per batch).

Carrier-free Formulations

Capsule filling

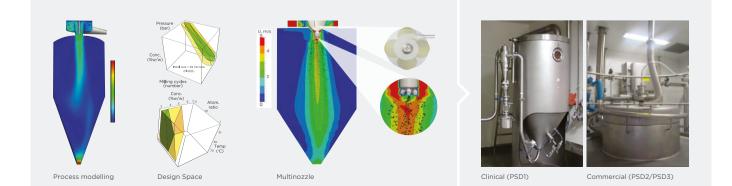
Also on the capsule filling side of the process, we offer a range of different pieces of equipment that can satisfy your demand. From a lab-scale weight-based Quantos capsule filler, that is semi-automatic and will fill up to 100 capsules per hour, to a capacitance-based (MultiNett system) pilot scale FlexaLab (MG2) that will fill up to 3 000 capsules per hour with 100% fill weight verification, Hovione can guarantee production of batches of up to 10 000 capsules, both in R&D and cGMP environments, so your development and clinical supply can be fully covered. With a recent investment on a Tekna (MG2) capsule filler, we now have the capability of going up to 70 000 capsules per hour and assure your low volume commercial supply of a capsule-based formulation for DPI.

API-alone formulations

For the kind of applications where no excipients can be used, for example proteins which are denatured when in contact with lactose and other sugars, you may need to develop an API-alone formulation. In this case, due to the necessary small size of the API particles for pulmonary delivery, the formulation can become too cohesive. Therefore, the properties of the powder need to be optimized in order to provide an appropriate aerodynamic performance. With Hovione's extensive expertise in Particle Engineering technologies, we are capable of achieving the best possible performance for your product. Spray drying, wet or dry milling, among others, can be applied to fine-tune the particle interactions and optimize the efficiency of delivery.

Composite particle formulations

For the platform of composite particle formulations, the optimization of the formulation in terms of composition and ratio of excipients and the particle engineering steps are both carried out simultaneously using spray drying technology. This allows the application of Hovione's strong expertise in spray drying directly to your formulation, which goes hand in hand with the application of QbD principles. From internal predictive tools using CFD and statistical modelling, to scale-up methodologies specific for inhalation particles, including patented multinozzle and taylor-made cyclone approaches, excellent control over the process as well as optimization of the final product performance can be guaranteed.



State-of-the-Art Analytical Characterization for Dry Powder Inhalation

Particle/Physical Characterization

- PSD by Laser Diffraction (Malvern and Sympatec)
- Computerized microscope imaging for particle characterization
- DVS
- Ultra-picnometer
- XRPD
- DSC
- Thermogravimetric analysis
- Scanning electron microscopy
- Surface area by BET & Raman spectroscopy

Formulation Characterization

- Fast Screening, Andersen Cascade and Next Generation Impactors
- Dosage Unit Sampling Apparatus (DUSA)
- Dissolution testing
- Breath simulators (closer to in vivo deposition)

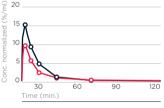
Biopharmaceutical Characterization

Microbiology

Beyond Cascade Impaction Evaluation

The use of anatomical throat models and breathing simulators allows formulation development under conditions mimicking different target patient populations.

Dissolution testing is an integral part of the development of inhalation formulations.





-O-Budesonide FPD -O-Budesonide ED

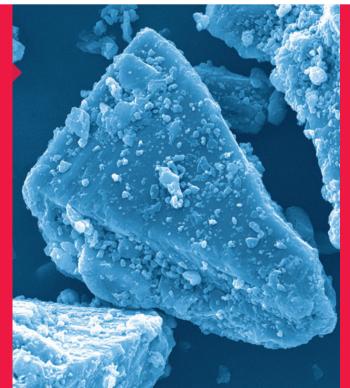
Breathing simulator

Portfolio of Off Patent APIs

Expertise in Customized APIs for Inhalation

Hovione's inhalation APIs are designed at the particle level to bring unique performance to your formulation.

- Fluticasone
- Salmeterol
- Umeclidinium
- GlycopyrrolateMometasone
- Vilanterol



Hovione 🌐

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