

DRY POWDER INHALER | SIEVED, MILLED, FINE MILLED LACTOSE

INHALAC

Technical brochure

General information

MEGGLE's sieved, milled and fine milled alpha-lactose monohydrate for dry powder inhalation: InhaLac®

MEGGLE's dry powder inhaler business has a long-standing and trustworthy partnership with leading formulators for dry powder inhalation (DPI). With years of experience and expertise, MEGGLE guarantees the production of high-quality lactose for pharmaceutical applications. MEGGLE is a partner to its global customers, providing them not only with high-quality products but also formulation services and ideas.

MEGGLE is known to collaborate with various pharmaceutical companies covering the entire pipeline of DPI development and commercialization. MEGGLE is committed to supporting its customers based on their needs. Thanks to its own globally distributed laboratories, MEGGLE is able to generate valuable application data rapidly.

The delivery of active pharmaceutical ingredients (APIs) via the lung is becoming increasingly important as more patients worldwide suffer from chronic respiratory diseases [1]. Additionally, the use of the respiratory tract for systemic application is gaining more attention than in the past.

MEGGLE benefits:

- Customization and tailored solutions
- Leading manufacturer of DPI lactose
- Comprehensive and constantly expanding portfolio
- Broad spectrum of tightly controlled particle sizes
- DPI grades available for all types of devices
- Individualized service with quick response time
- Expertise and support in the development of DPI formulations
- Global application laboratories to optimize new and existing formulations
- Collaboration with leading industrial partners and scientific institutes
- Lactose for parenteral or ophthalmic application available



Product description

DPIs are widely used in pulmonary drug delivery. This is because of their advantages like ease of use, small size, portability, and lack of breath-actuation coordination requirement [2]. They are also environmentally friendly as they are propellant-free and comparatively stable as solid-particle formulations [3].

Typically, this dosage form comprises a device, one or more APIs, and excipients. The main function of the excipient lactose is to act as carrier. Carriers are used to help deposit the active ingredient in the lung and may have a secondary role in diluting the active to ensure that dosages can be properly metered. Other excipients, like lactose fines or force control agents, may be used to further modify drug detachment.

Lactose has a long tradition in inhalative dosage forms and is proven to be safe. Therefore, lactose is the excipient of choice in pulmonary drug delivery. MEGGLE's InhaLac® product family is highly specialized and produced using a dedicated and well-documented process.

MEGGLE offers a wide range of sieved, milled, and fine milled inhalation grades of lactose monohydrate – the InhaLac® types. All of them have excellent physicochemical characteristics and conform to the highest compendial requirements. The broad MEGGLE portfolio allows to meet any formulator's expectations.

APPLICATION

InhaLac®

- InhaLac® is suitable for use in pulmonary and nasal drug delivery.

BENEFITS

InhaLac®

- Highly controlled powder characteristics
- Low microbial burden including endotoxines
- Customized grades

Regulatory & quality information

MEGGLE's InhaLac® grades comply with the current USP-NF, Ph. Eur., JP and ChP monographs. In order to meet the special requirements for pulmonary drug delivery, additional and in some cases even stricter specification limits are in place for all InhaLac® grades. For many InhaLac® grades a drug master file (DMF) is available for drug product submission review and approval in China. Specifications and regulatory documents can be downloaded from www.meggle-excipients.com.

MEGGLE's pharma-dedicated production facility in Wasserburg, Germany is certified according to DIN EN ISO 9001 and has implemented GMP according to the Joint IPEC-PQG (Good Manufacturing Practices Guide for Pharmaceutical Excipients) and USP-NF General Chapter <1078> GOOD MANUFACTURING PRACTICES FOR BULK PHARMACEUTICAL EXCIPIENTS. MEGGLE has been an EXCiPACT™-certified excipient

manufacturer and supplier since 2014. All InhaLac® products are manufactured on product lines exclusively dedicated to inhalation lactose. Additionally, MEGGLE is a member of IPEC (International Pharmaceutical Excipients Council).

MEGGLE invests considerably in the sustainability of raw material sourcing, production standards, and efficiency and is actively engaged in environmental protection. In order to guarantee the quality of the products, commitment and adherence to established pharmaceutical standards is the highest priority.



Particle Size Distribution (PSD) & Morphology (SEM)

InhaLac® grades are produced through crystallization, followed by sieving or milling, using a dedicated production line. The optimized and standardized production process consistently ensures the highest quality production, made in Germany.

Lactose is usually used in a DPI formulation as a coarse carrier to help deposit the active ingredient in the lung and may have a secondary role in diluting the active to ensure that dosages can be properly metered. Moreover, finely milled lactose is often used to improve the performance (detachment of API). Formulation requirements vary depending on the API's target concentration and characteristics, like particle size and shape hydrophilicity/lipophilicity, as well as on the device. Additionally, downstream processes like blending and filling are also important to consider.

The different requirements have to be considered during development to ensure a high and repeatable delivery of the API to the lungs as well as a robust production process. Carrier properties are hereby significantly determining the product performance. Therefore, MEGGLE's InhaLac® portfolio offers a variety of both sieved (tomahawk-shaped particles) and milled (irregular shaped) grades with distinct and highly controlled particle sizes as carriers, as well as fine milled grades for DPI performance fine tuning.



Figure 1: Specified PSD of InhaLac® grades by laser diffraction.

DPI Selection Guide

InhaLac® is suitable for use in pulmonary and nasal drug delivery.

MEGGLE offers for each application the right lactose grade, but there is no one fits all solution. The **figure 2** gives some examples for selection of suitable DPI grade lactose. Please contact the MEGGLE experts for further help to choose your InhaLac®.

The many different available InhaLac® grades offer you a high flexibility (“tool-box”) to achieve the desired product performance.

Sieved						Milled							
Coarse sieved lactose			Sieved lactose			Milled lactose					Fine milled lactose		
InhaLac® 70	InhaLac® 120	InhaLac® 160	InhaLac® 230	InhaLac® 240	InhaLac® 251	InhaLac® 180	InhaLac® 140	InhaLac® 145	InhaLac® 150	InhaLac® 300	InhaLac® 400	InhaLac® 500	
250 µm Median particle size						The difference in particle morphology between sieved and milled should be also considered for selection.							
11%						40% Compressibility/Carr's Index							
0%						75% Amount of intrinsic fines < 5 µm							
InhaLac® 70	InhaLac® 120	InhaLac® 160	InhaLac® 230	InhaLac® 240	InhaLac® 251	InhaLac® 180	InhaLac® 140	InhaLac® 145	InhaLac® 150	InhaLac® 300	InhaLac® 400	InhaLac® 500	
E.g. ideal for metered dose/reservoir based devices, which typically require good to fair flowability (CI 11–20%); low amount of intrinsic fines						E.g. typically used for capsule & blister based formulations, for which some cohesivity is required; upper limit depends on the dosing principle and filling volume: Dosator principle and drum dosing are both suitable for micro-dosing of a variety of different powders; with drum dosing even very cohesive powders (CI > 38%) can be filled precisely. Dosing disc system might have some limitations regarding micro-dosing precision and powder cohesivity (typically CI 20–31%)						E.g. typically used to fine tune DPI performance (typically addition of 2–20%) or for soft pellet formulations	

Figure 2: Filling and selection of device is essential for DPI development, from most common filling principals and devices beneficial powder characteristics are given.

Key factors influencing DPI performance

Achieving optimal performance in Dry Powder Inhalation (DPI) formulations depends on the precise interaction of several critical elements as schematically shown below.

Each of these factors must be carefully optimized and harmonized to deliver a DPI product that is both effective and reliable. MEGGLE's expertise in DPI formulation support helps customers navigate these complexities to achieve high-performance solutions.

API (Active Pharmaceutical Ingredient):

The physical and chemical properties of the API, such as particle size, shape, and surface energy, play a major role in determining aerosolization efficiency and deposition behavior.

Carrier:

The choice of carrier material, typically lactose, impacts the uniformity, flowability, and overall stability of the powder blend. Carrier properties such as surface morphology and particle size are crucial for ensuring consistent drug delivery.

Fines:

The presence and proportion of fine particles (<5 µm) can significantly affect drug detachment from the carrier during inhalation, influencing the respirable dose and therapeutic effectiveness.

Blending:

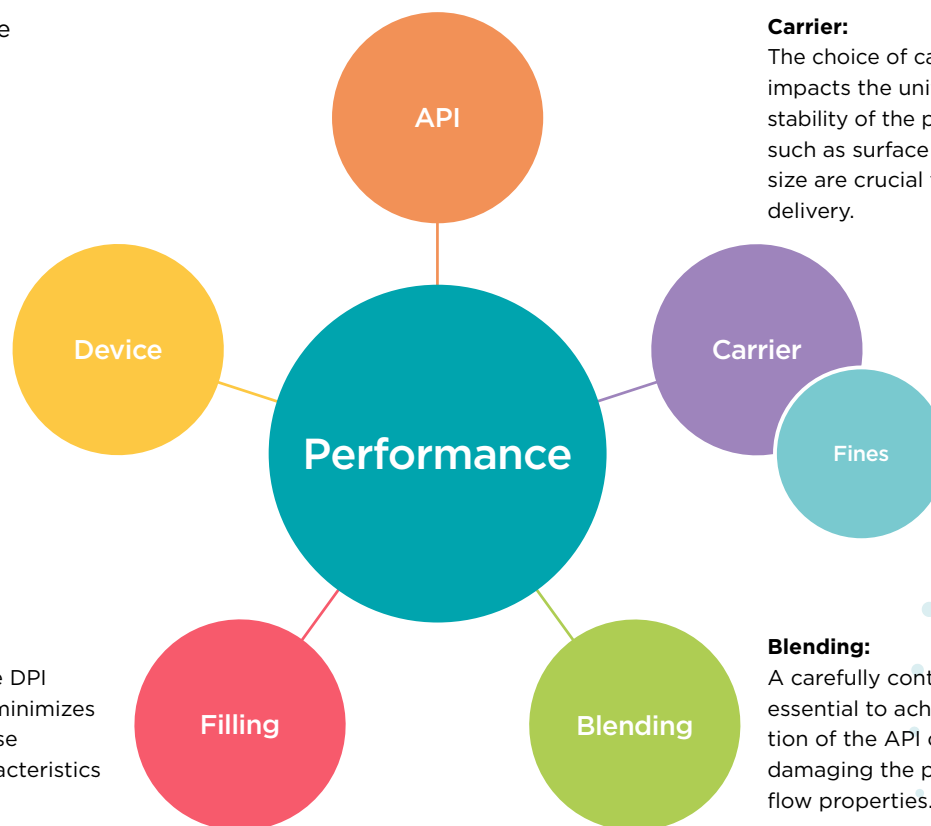
A carefully controlled blending process is essential to achieve homogeneous distribution of the API on the carrier surface without damaging the particles or altering powder flow properties.

Filling:

Accurate and consistent filling of the DPI device ensures correct dosage and minimizes variability. Filling challenges may arise depending on the powder flow characteristics and dosing system used.

Device:

The inhalation device design, including its resistance, dispersion mechanism, and airflow patterns, must be compatible with the formulation to maximize delivery efficiency and patient usability.



Sieved lactose for inhalation – Properties

Sieved inhalation lactose grades consist of single or agglomerated crystals, mostly typically referred to as tomahawk-shaped. Coarser material exhibits a higher share of agglomerate particles.

Sieved InhaLac® grades with good flowability – most suitable for reservoir formulations

InhaLac® 70, the coarsest, sieved product, has a typical median particle size of approximately 215 µm, is virtually free of fines (particles < 5 µm). It shows a narrow PSD and is best suited to cyclone-based inhalation devices. InhaLac® 70 has additionally an agglomerated structure and poses certain pockets and cavities, which makes it very suitable for higher drug loading applications.

InhaLac® 120 (typical median particle size: ~130 µm) and InhaLac® 160 (median particle size: ~110 µm) are also a coarse sieved grade, but finer compared to InhaLac® 70. They have good to excellent flowability (Carr's Index ≤15%) and are well suitable for application focusing e.g. on reservoir devices.

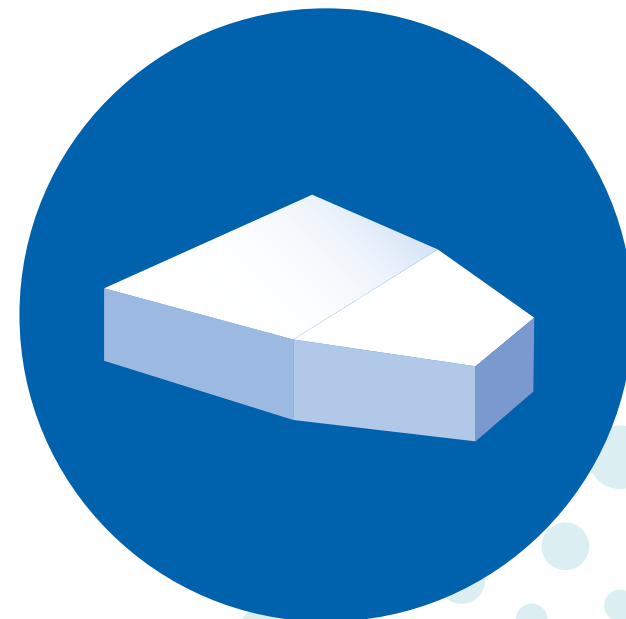
Sieved InhaLac® grades – ideal for capsules or blisters

InhaLac® 230 (median particle size: ~100 µm) and a fines content between 3 – 5%.

InhaLac® 240 is a sieved grade with a fine median particle size (~65 µm), but still fair flowability characteristics (Carr's Index ≤20%).

InhaLac® 251, the finest sieved lactose grade, has a median particle size of approximately 50 µm. The product is characterized by a higher fines content. The compressibility/flowability is more comparable to milled grades.

Stay tuned for regularly published application studies and white papers: www.meggle-excipients.com



Sieved lactose for inhalation – Particle Size Distribution (PSD)

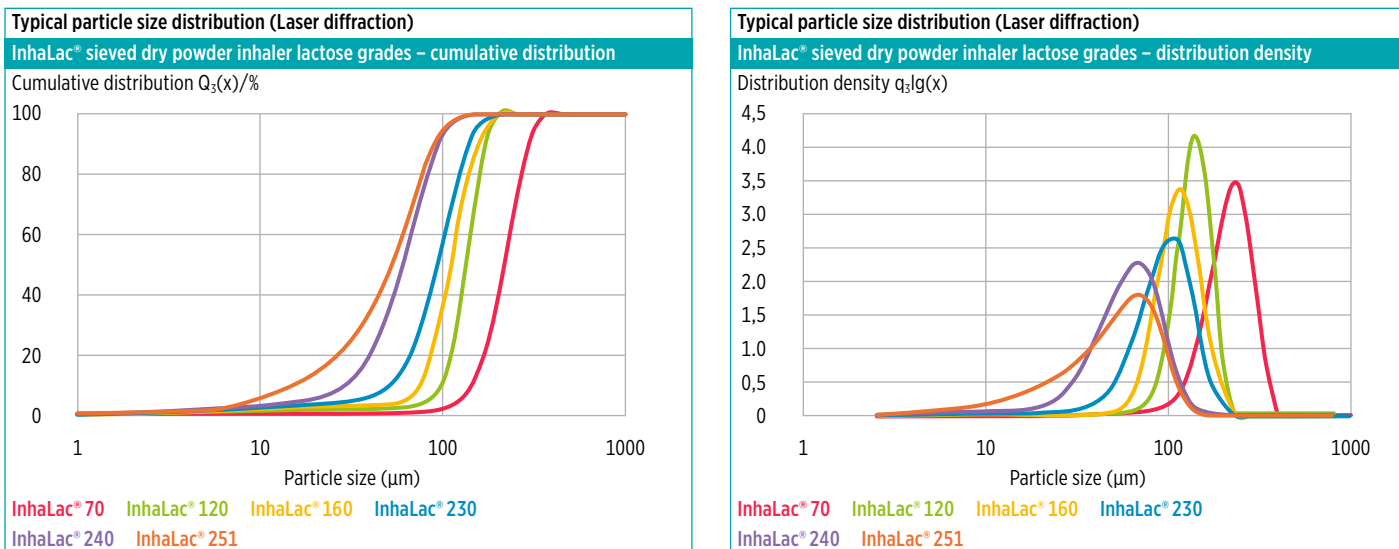


Figure 3 – 4: Typical cumulative particle size and density distribution of MEGGLE's sieved inhalation lactose grades InhaLac® 70, InhaLac® 120, InhaLac® 160, InhaLac® 230, InhaLac® 240 and InhaLac® 251. Analyzed by Sympatec®/Helos & Rodos particle size analyzer.

Sieved InhaLac® grades		InhaLac® 70	InhaLac® 120	InhaLac® 160	InhaLac® 230	InhaLac® 240	InhaLac® 251
Lactose type		typical	typical	typical	typical	typical	typical
Particle size distribution	x_{10}	140 μm	97 μm	75 μm	50 μm	30 μm	15 μm
	x_{50}	215 μm	137 μm	111 μm	96 μm	63 μm	52 μm
Laser diffraction	x_{90}	295 μm	178 μm	152 μm	144 μm	101 μm	92 μm
	Span $[(x_{90}-x_{10})/x_{50}]$	0.7	0.6	0.7	1.0	1.1	1.5
	% fines < 5 μm	0	1	2	2	3	3

Figure 5: Typical PSD values of MEGGLE's sieved inhalation lactose grades by laser diffraction.

Milled lactose for inhalation – Properties

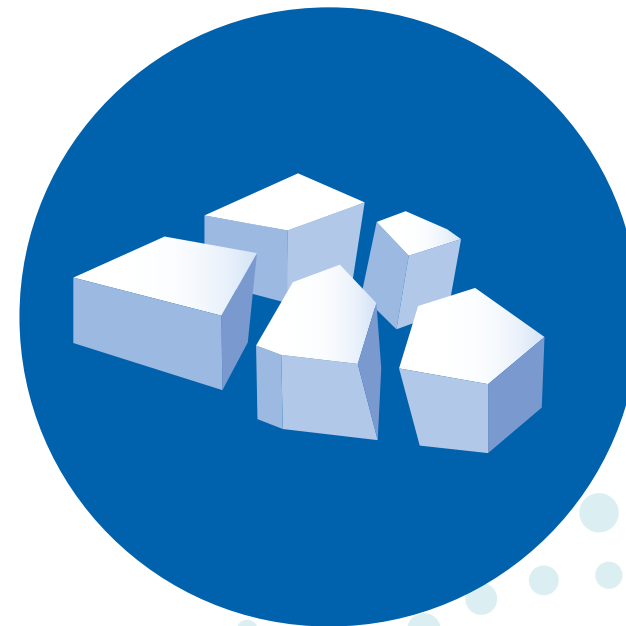
Milled InhaLac® for superior performance during blister and capsule filling

In contrast to the sieved grades, milled and fine milled grades consist of lactose particles that are finer, more irregular and sharp-edged due to the manufacturing by milling. Milled lactose typically has a broader particle size distribution and a smaller mean compared to sieved grades. Therefore, cohesiveness is typically higher (Carr's Index $\geq 31\%$). milled grades are carriers for capsule and blister based formulations with intrinsic amount of fines.

MEGGLE's current portfolio contains four milled carrier grades, for which the PSD characteristics are highly controlled. InhaLac®145 is a middle sized carrier with a mean particle size of 35 μm and approximately 11% of particles below 5 μm . In comparison InhaLac®140 shows a mean of 50 μm and InhaLac®150 has with 24 μm the smallest mean particle size. The possibility to choose between the variety of milled grades allows the formulator to evaluate the design space.

The portfolio is completed by the blockbuster product InhaLac®180. It features the frequently requested particle sizes of $x_{10}=5-25\text{ }\mu\text{m}$, $x_{50}=50-100\text{ }\mu\text{m}$ and $x_{90}=120-160\text{ }\mu\text{m}$. InhaLac®180 could be described as a broad lactose suitable for many applications. It contains approximately 6% of intrinsic fine particles below 5 μm .

Check out MEGGLE's latest application studies for blockbuster formulations using milled InhaLac® grades:
www.meggle-excipients.com

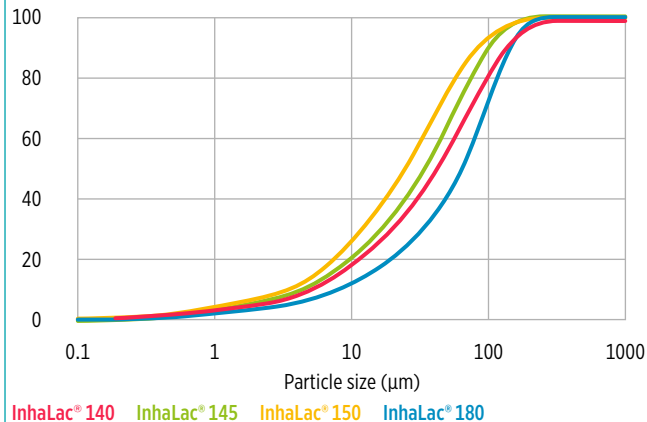


Milled lactose for inhalation – Particle Size Distribution (PSD)

Typical particle size distribution (Laser diffraction)

InhaLac® milled dry powder inhaler lactose grades – cumulative distribution

Cumulative distribution $Q_3(x)/\%$



Typical particle size distribution (Laser diffraction)

InhaLac® milled dry powder inhaler lactose grades – distribution density

Distribution density $q_3lg(x)$

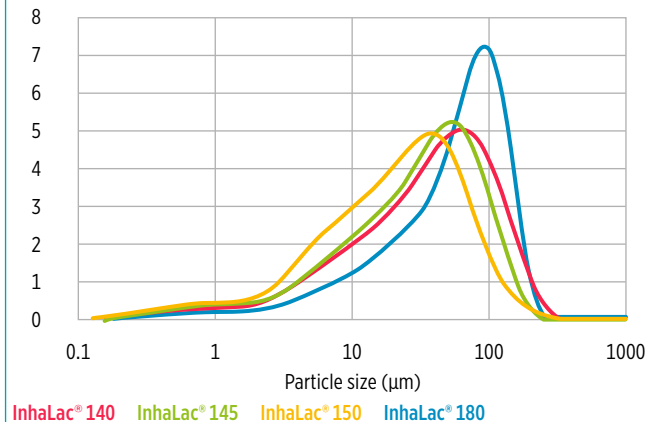


Figure 6 – 7: Typical cumulative PSD and distribution density of MEGGLE's milled inhalation lactose grades, InhaLac® 140, InhaLac® 145, InhaLac® 150, and InhaLac® 180. Analyzed by Malvern Mastersizer 3000 laser diffraction system.

Milled InhaLac® grades

	Lactose type	InhaLac® 140	InhaLac® 145	InhaLac® 150	InhaLac® 180
		typical	typical	typical	typical
Particle size distribution	x_{10}	5 μm	4 μm	4 μm	9 μm
	x_{50}	45 μm	34 μm	25 μm	65 μm
Laser diffraction	x_{90}	135 μm	102 μm	80 μm	143 μm
	Span $[(x_{90}-x_{10})/x_{50}]$	2.9	2.9	3.0	2.1
	% fines < 5 μm	10	11	14	6

Figure 8: Typical PSD values of MEGGLE's milled inhalation lactose grades by laser diffraction.

Fine milled lactose for inhalation – Properties

A well-balanced proportion of finely milled or micronized lactose in the DPI blend is often important to achieve the required performance of the DPI formulation. This is because fines can advance the deposition via various mechanisms like enhanced aerolidsation/dispergation. However, different quantities and qualities of lactose fines can not only influence the deposition performance of the API, but they may also impact the flowability and further downstream processing of the material.

MEGGLE offers three grades of fines with different characteristics and particle size distribution. This allows the formulator to choose according to requirements and to evaluate the design space. For each formulation, a compromise must be found between the titration of active sites, the adjustment of the performance parameters and the flow behavior of the powder bulk.

InhaLac® 300 – coarsest fine quality

InhaLac® 300 contains 90 % of particles below 35 – 50 µm (Sympatec) and 40 – 56 µm (Malvern). The reason to add this grade is to modulate the cohesiveness of the powder rather than to adjust the deposition of the API. Cohesiveness is particularly important during encapsulation.

InhaLac® 400 – “the multitalent”

InhaLac® 500 – “the micronized”

MEGGLE’s finest grades are used to tune DPI performance in tertiary blends. The addition of fines usually improves aerosolisation performance by enhancing the API detachment or co-agglomeration. InhaLac® 400 and InhaLac® 500 are both fine lactose grades produced under different milling conditions to achieve the desired particle size distribution (PSD). InhaLac® 500 is finer, with 90 % of particles smaller than ≤ 10 µm. InhaLac® 400 is slightly coarser with a typical median particle size of $x_{50} = 8$ µm.

Typical concentrations of fines ranges from a few percent up to 20% for InhaLac® 400 and up to 10% for InhaLac® 500. If the fine content is too high, flowability and further processing may be impaired.

InhaLac® 500 is also an excellent choice for soft pellets. Soft pellets typically consist onyl of API and fines.

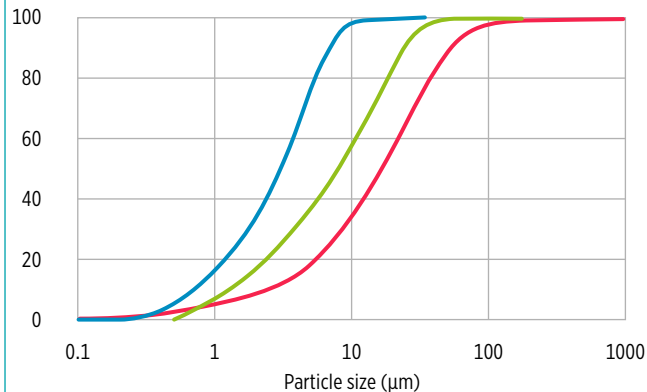


Fine milled lactose for inhalation – Particle Size Distribution (PSD)

Typical particle size distribution (Laser diffraction)

InhaLac® fine milled dry powder inhaler lactose grades – cumulative distribution

Cumulative distribution $Q_3(x)/\%$

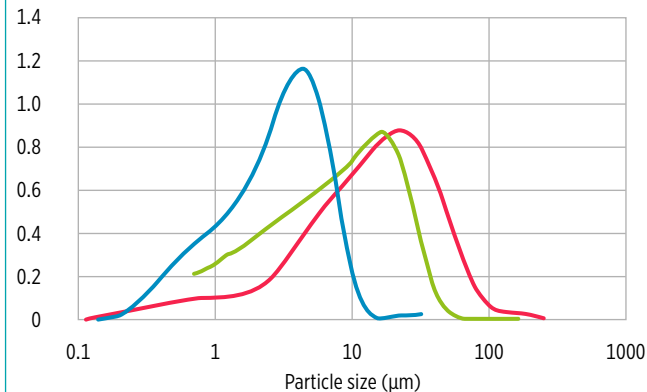


InhaLac® 300 InhaLac® 400 InhaLac® 500

Typical particle size distribution (Laser diffraction)

InhaLac® fine milled dry powder inhaler lactose grades – distribution density

Distribution density $q_3 \lg(x)$



InhaLac® 300 InhaLac® 400 InhaLac® 500

Figure 9 – 10: Typical cumulative PSD and distribution density of MEGGLE's fine milled inhalation lactose grades, InhaLac® 300, InhaLac® 400 and InhaLac® 500. Analyzed by Sympatec®/Helos & Rodos particle size analyzer.

Fine milled/micronized InhaLac® grades

Lactose type		InhaLac® 300	InhaLac® 400	InhaLac® 500
		typical	typical	typical
Particle size distribution	x_{10}	2.5 μm	1.2 μm	–
	x_{50}	16.0 μm	8.0 μm	3.1 μm
Laser diffraction	x_{90}	50.0 μm	25.0 μm	9.9 μm
	Span $[(x_{90} - x_{10})/x_{50}]$	3.0	3.0	2.5
	% fines < 5 μm	18	38	75

Figure 11: Typical PSD values of MEGGLE's fine milled inhalation lactose grades by laser diffraction.

Functional related characteristics

MEGGLE ensures highest quality for critical properties. MEGGLE monitors FRC (functional related characteristics) and supports customers with their expertise. MEGGLE's sophisticated knowledge management allows fast and easy answering of customer questions.

Typical powder technological values

Figure 12 provides additional information on the other functional characteristics of the inhalation lactose grades.

Typical powder technological values					
InhaLac®					
	BET surface (m²/g)	Density bulk (g/ml)	Density tapped (g/ml)	Hausner ratio	Carr's Index (%)
Sieved					
InhaLac® 70	0.2 ¹	0.66	0.77	1.17	14
InhaLac® 120	0.3 ¹	0.73	0.83	1.14	12
InhaLac® 160	0.3 ¹	0.71	0.83	1.17	14
InhaLac® 230	0.3 ¹	0.73	0.87	1.19	16
InhaLac® 240	0.3 ¹	0.71	0.87	1.23	18
InhaLac® 251	0.4 ¹	0.62	0.87	1.40	29
Milled					
InhaLac® 140	0.9 ¹	0.58	0.90	1.55	35
InhaLac® 145	1.3 ¹	0.54	0.87	1.61	38
InhaLac® 150	1.3 ¹	0.51	0.84	1.65	39
InhaLac® 180	0.4 ¹	0.63	0.91	1.44	31
Fine milled					
InhaLac® 300	1.5 ¹	0.44	0.72	1.67	39
InhaLac® 400	2.2 ²	0.35	0.58	1.66	40
InhaLac® 500	5.3 ²	0.21	0.32	1.52	34

Figure 12: Typical technological powder values of MEGGLE's inhalation lactose grades (Quantachrome Autosorb-3, Krypton adsorption/Nitrogen adsorption²).

Microbiology

All of MEGGLE's InhaLac® grades have stricter or additional microbial limits compared to the current monographs of the Pharmacopoeia. This guarantees the highest safety in the use of InhaLac® grades in DPI formulations. All microbiological parameters listed in **figure 13** are part of the product specification. MEGGLE has a validated production process with respect to bacterial endotoxins.

Microbiology	
InhaLac®	
Parameters	Specified
Total aerobic microbial count (TAMC)	NMT 10 cfu/g
Total combined yeasts and molds count (TYMC)	NMT 10 cfu/g
Bile tolerant gramnegative bacteria	absence/10 g
Escherichia coli	absence/10 g
Pseudomonas aeruginosa	absence/10 g
Staphylococcus aureus	absence/10 g
Salmonella spp.	absence/10 g
Burkholderia cepacia	absence/10 g
Bacterial endotoxins	< 5 EU/g

Figure 13: Specified microbiological parameters of MEGGLE's inhalation lactose grades.

Batch-to-batch consistency

Batch-to-batch consistency for all lactose products can be attributed to MEGGLE's long history and experience in lactose manufacturing, and broad technical expertise. Constant in-process and final product testing ensures consistency and quality.

Packaging and shelf life

Packaging material complies with Regulation (EC) No. 1935/2004 and 21 CFR 174, 175, 176, 177 and 178. Stability tests were performed according to ICH guidelines and an ongoing stability program is in place. **Figure 14** provides information on packaging size, material, and shelf life.

Packaging and shelf life			
Sieved			
	Size	Material	Shelf life
InhaLac® 70	25 kg	Carton box with an aluminium laminated inliner	24 Months
InhaLac® 120			
InhaLac® 160			
InhaLac® 230		Carton box with aluminium laminated and PE-EVOH-PE inliner	
InhaLac® 240			
InhaLac® 251			
Milled			
InhaLac® 140	25 kg	Carton box with aluminium laminated and PE-EVOH-PE inliner	24 Months
InhaLac® 145	20 kg		
InhaLac® 150			
InhaLac® 180	25 kg		
Fine milled			
InhaLac® 300	15 kg	Carton box with aluminium laminated and PE-EVOH-PE inliner	24 Months
InhaLac® 400		Carton box with aluminium laminated inliner	
InhaLac® 500	6 kg	Carton box with aluminium laminated and PE-EVOH-PE inliner	18 Months

Figure 14: Packaging and shelf life of MEGGLE's inhalation lactose grades.

Customization according to your needs

The performance of inhalers depends on the particle size and flow characteristics of the selected lactose. Precise control of these factors is crucial for efficacy and dosing accuracy.

MEGGLE offers tailor-made inhalative lactose solutions, using long-standing expertise and flexible production to deliver customized lactose with specific particle size distributions and high quality. Understanding your requirements is essential for a successful outcome.

Collaboration with MEGGLE's Inhalation Experts ensures a structured development process, guided by a dedicated project manager.

The result: a well-characterized product that fully meets your individual needs.

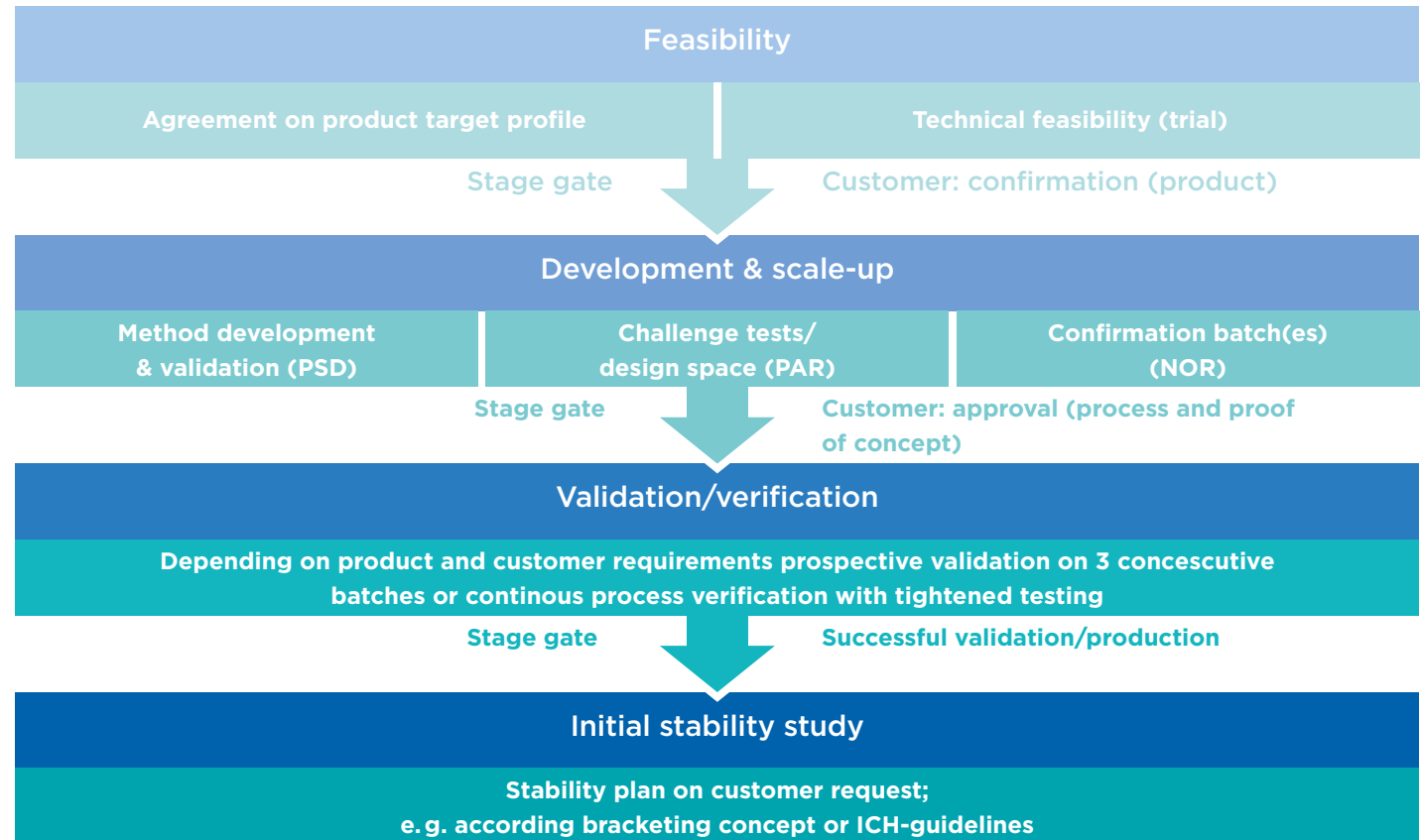


Figure 15: Stage-Gate-Model for the development process of customized products.

Experts in dry powder inhalation – Technical support

Global service labs, the Innovation & Formulation Campus Munich and partnering with leading research institutions in the field to accelerate your development

With a long history in manufacturing and distribution of excipients for the pharmaceutical industry MEGGLE has a lot of expertise to share. The MEGGLE R&D team works in close collaboration with research institutes and universities all over the world. This provides customers with additional technical and analytical data and support. The capabilities of the MEGGLE experts as well as the product portfolio are continuously being expanded.

MEGGLE can utilize an application lab to cope with any challenge during the development of your DPI formulation.

Why is blending so important and how MEGGLE understands your need

A major challenge is to reach content uniformity for the typically low dose of the API (<5% API). In this respect, the current trend towards combining several APIs in one formulation increases the precautions that formulators have to take to ensure blend uniformity. For example, a modern DPI formulation is very often a multi-component system containing 2 or 3 APIs, lactose carriers and lactose fines and additional Force Control Agents such as magnesium stearate. There are several ways to blend these components and a multitude of influencing factors (type of blender, sequence and method of addition of different components, mixing energy), all of which can affect homogeneity and DPI performance.

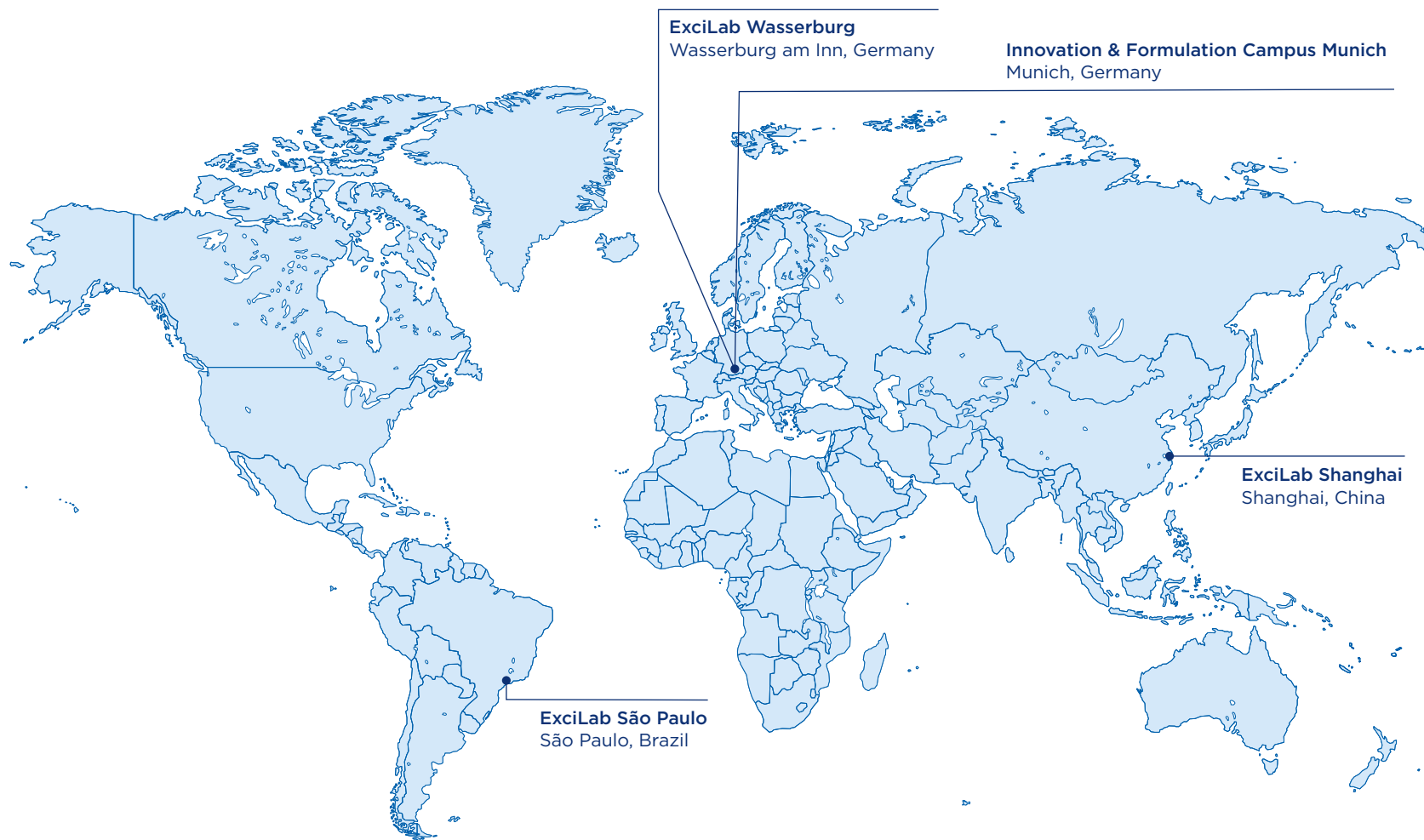
Therefore, a profound understanding of blends and blending is important. In the fully equipped Innovation & Formulation Campus Munich, the customer's process can be mimicked and helped to elucidate the optimal blending. Reversed engineering and typically analysis like homogeneity, particle size distribution and morphology are suitable as well. The customer is supported in every aspect of formulation development.

SERVICE & SUPPORT

Includes

- Manufacturing of blends
- Advanced powder and blend characterization
- DPI testing (NGI, DUSA)
- Application studies for selected formulations

Experts in dry powder inhalation – Locations



Literature

- [1] Bousquet, J., Khaltaev, N. (2007). Global surveillance, prevention and control of chronic respiratory diseases: a comprehensive approach WHO Library Cataloguing-in-Publication Data: ISBN 978 92 4 156346 8 (NLM classification: WF 140), World Health Organization.
- [2] Labris, N.R., Dolovich, M. (2003). Pulmonary drug delivery. Part II: The role of inhalant delivery devices and drug formulations in therapeutic effectiveness in aerosolized medications, 56: 600-612.
- [3] Pilcer, G., Amighi, K. (2010). Formulation strategy and use of excipients in pulmonary drug delivery. International Journal of Pharmaceutics, 392: 1-19.

Submitted by

MEGGLE GmbH & Co. KG

Business Unit Excipients

Megglesstrasse 6-12
83512 Wasserburg
Germany

Tel.: +49 8071 730

info.excipients@meggle.com

www.meggle-excipients.com

Do you need technical support?

inhalation.excipients@meggle.com

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