

Manufacturer/Importer Authorisation ^{1, 2}

1. Authorisation Number DE_BY_05_MIA_2024_0004
2. Name of authorisation holder Labor Ls SE & Co. KG (ORG-100011865 / LOC-100018825)
3. Address(es) of manufacturing site(s) Labor LS SE & Co. KG (ORG-100011865 / LOC-100018825), Mangelsfeld 4-6, Grossenbrach, Bad Bocklet, Bavaria, 97708, Germany
- 3.a Additional details on units inspected of manufacturing site(s) address(es)
4. Legally registered address of authorisation holder Mangelsfeld 4-6, Grossenbrach, Bad Bocklet, Bavaria, 97708, Germany
- 4.a Additional details on units inspected of legally registered address
5. Scope of authorisation and dosage forms² ANNEX 1 and/ or ANNEX 2
6. Legal Basis of authorisation Art. 88 of Regulation (EU) 2019/6
7. Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation confidential
8. Signature
9. Date 2024-01-17
10. Annexes attached Annex 1 and/or Annex 2
Optional Annexes as required:
Annex 3(Addresses of Contract Manufacturing Site(s))
Annex 4(Addresses of Contract laboratories)
Annex 5(Name of Qualified Person)
Annex 6(Name of responsible persons)
Annex 7(Date of inspection on which authorisation granted, scope of last inspection)
Annex 8(Manufactured/ imported products authorised)³

¹The authorisation referred to in paragraph 40(1) of Directive 2001/83/EC as amended and Article 88(1) of Regulation (EU) 2019/6, shall also be required for imports coming from third countries into a Member State.

²Guidance on the interpretation of this template can be found in the Interpretation of the Union format for Manufacturer/Importer Authorisation.

³The Competent Authority is responsible for the appropriate linking of the authorisation with the manufacturer's application (Article 42(3) of Directive 2001/83/EC as amended and Article 90(3) of Regulation (EU) 2019/6).

SCOPE OF AUTHORISATION

ANNEX 1

Name and address of the site: Labor LS SE & Co. KG, Mangelsfeld 4-6, Grossenbrach, Bad Bocklet, Bavaria, 97708, Germany

Additional Details:

Veterinary Medicinal Products

<p>Authorised Operations MANUFACTURING OPERATIONS(according to part 1) IMPORTATION OF MEDICINAL PRODUCTS(according to part 2)</p>
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Part 1 - MANUFACTURING OPERATIONS	
1.1	Sterile products
	<i>1.1.3 Batch certification</i>
1.2	Non-sterile products
	<i>1.2.2 Batch certification</i>
1.6	Quality control testing
	<i>1.6.1 Microbiological: sterility</i> <i>1.6.2 Microbiological: non-sterility</i> <i>1.6.3 Chemical/Physical</i> <i>1.6.4 Biological</i>

Any restrictions or clarifying remarks related to the scope of these Manufacturing operations (for Public users)

The manufacturing authorization is restricted to products that are tested and released by Labor LS SE & Co. KG on a subcontract basis.

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS	
2.1	Quality control testing of imported medicinal products
	<i>2.1.1 Microbiological: sterility</i> <i>2.1.2 Microbiological: non-sterility</i> <i>2.1.3 Chemical/Physical</i> <i>2.1.4 Biological</i>
2.2	Batch certification of imported medicinal products

	<p>2.2.1 <i>Sterile products</i></p> <ul style="list-style-type: none">2.2.1.1 Aseptically prepared2.2.1.2 Terminally sterilised
	<p>2.2.2 <i>Non-sterile products</i></p>

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