Manufacturer/Importer Authorisation

1. Authorisation Number DE_MV_02_MIA_2023_0001

2. Name of authorisation holder Z.A.S. Zentral Archiv Service GmbH (ORG-100033486 /

LOC-100052395)

3. Address(es) of manufacturing site(s) Z.A.S. Zentral Archiv Service GmbH (ORG-100033486 /

LOC-100052395), Justus-Von-Liebig-Strasse 7, Stadtgebiet West, Neubrandenburg, Mecklenburg-Western-Pommerania, 17033,

Germany, GPS: 53.570971, 13.209843

3.a Additional details on units inspected of

manufacturing site(s) address(es)

4. Legally registered address of authorisation

holder

Justus-Von-Liebig-Strasse 7, Stadtgebiet West, Neubrandenburg, Mecklenburg-Western-Pommerania, 17033, Germany, GPS:

53.570971, 13.209843

4.a Additional details on units inspected of

5. Scope of authorisation and dosage forms²

legally registered address

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 2023-08-17

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)

SCOPE OF AUTHORISATION

ANNEX 1

Name and address of the site: Z.A.S. Zentral Archiv Service GmbH,

Justus-Von-Liebig-Strasse 7, Stadtgebiet West,

Neubrandenburg, Mecklenburg-Western-Pommerania, 17033,

Germany, GPS: 53.570971, 13.209843

Additional Details:

Veterinary Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)

Part 1 - MANUFACTURING OPERATIONS	
1.4	Other products or manufacturing activity
	1.4.3 Other: Storage of GMP documents and retained samples of veterinary medicinal products, master and working seeds (viruses, bacteria, fungi, cell lines) on behalf of various manufacturing license holders(en)

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).