

MANUFACTURER'S AUTHORISATION

(This English translation is for reference only. It is not part of the official certificate.)

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| 1. Authorisation number/file number | DE_RP_01_MIA_2024_0016 |
| 2. Name of authorisation holder | HWI pharma services GmbH (LOC-100006965) |
| 3. Address(es) of manufacturing site(s) | HWI pharma services GmbH Rheinzaberner Straße 8 76761 Rülzheim (LOC-100006965) |
| 4. Legally registered address of authorisation holder | Rheinzaberner Strasse 8 76761 Rülzheim |
| 5. Scope of authorisation and dosage forms | ANNEX 1 and ANNEX 2 |
| 6. Legal basis of authorisation | Sect 13 para 1 Arzneimittelgesetz (German Drug Law) Art. 88 of Regulation (EU) 2019/6 and Sect 28 para 1 German Veterinary Medicinal Products Law Art. 61 para 1 to 3 of Regulation (EU) No. 536/2014 in conjunction with Sect 13 para 5 AMG |
| 7. Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation | |
| 8. Signature | On behalf |
| 9. Date | 23/07/2024 |
| 10. Annexes attached | Annex 1 and Annex 2 Annex 4 (Addresses of Contract Laboratories) |

Annex 5 (Name of Qualified Person)
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:

HWI pharma services GmbH, Rheinzaberner Straße 8, 76761 Rülzheim

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| Human Medicinal Products Veterinary Medicinal Products |
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AUTHORISED OPERATIONS

Manufacturing Operations (according to part 1)

| Part 1 - MANUFACTURING OPERATIONS | |
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| 1.1 | Sterile Products |
| | 1.1.3 <i>Batch certification</i> |
| 1.2 | Non-sterile products |
| | 1.2.2 <i>Batch certification</i> |
| 1.3 | Biological medicinal products |
| | 1.3.2 <i>Batch certification</i> |
| | 1.3.2.6 Human or animal extracted products |
| 1.6 | Quality control testing |
| | 1.6.3 <i>Chemical/Physical</i> |

SCOPE OF AUTHORISATION

Annex 2

Name and address of the site:

HWI pharma services GmbH, Rheinzaberner Straße 8, 76761 Rülzheim

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| Investigational Medicinal Products for Human Use |
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| AUTHORISED OPERATIONS Manufacturing Operations (according to part 1) |
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| Part 1 - MANUFACTURING OPERATIONS | |
|--|--------------------------------------|
| 1.1 | Sterile Products |
| | 1.1.3 Batch certification |
| 1.2 | Non-sterile products |
| | 1.2.2 Batch certification |
| 1.3 | Biological medicinal products |
| | 1.3.2 Batch certification |
| | 1.3.2.5 Biotechnology products |
| 1.6 | Quality control testing |
| | 1.6.3 Chemical/Physical |