




MANUFACTURER'S AUTHORISATION

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

- | | |
|--|--|
| 1. Authorisation number/file number | DE_RP_01_MIA_2018_0040 |
| 2. Name of authorisation holder | Popack Logistik GmbH |
| 3. Address(es) of manufacturing site(s) | Popack Logistik GmbH
Achardstraße 1
55127 Mainz |
| 4. Legally registered address of authorisation holder | Achardstraße 1
55127 Mainz |
| 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) |
| 7. Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation | Dr. Ilka Petry |
| 8. Signature | On behalf
 |
| 9. Date | 27/08/2018 |
| 10. Annexes attached | Annex 1 and Annex 2
Annex 7 (Date of inspection on which authorisation granted, scope of last inspection) |





SCOPE OF AUTHORISATION

Name and address of the site:

Popack Logistik GmbH, Achardstraße 1, 55127 Mainz

Human Medicinal Products

AUTHORISED OPERATIONS

Manufacturing Operations (according to part 1)

Part 1 - MANUFACTURING OPERATIONS

1.5 Packaging

1.5.2 Secondary packing





SCOPE OF AUTHORISATION

Name and address of the site:

Popack Logistik GmbH, Achardstraße 1, 55127 Mainz

Investigational Medicinal Products for Human Use

AUTHORISED OPERATIONS

Manufacturing Operations (according to part 1)

Part 1 - MANUFACTURING OPERATIONS

1.5 Packaging

1.5.2 Secondary packing





Date of Inspection on which
authorisation was granted

15/08/2018

Scope of last Inspection

General GMP-Inspection (details see inspection report)

