



## MANUFACTURER'S AUTHORISATION

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

1. Authorisation number/file number	DE_NW_06_MIA_2018_0001/8.87-40.15.07
2. Name of authorisation holder	Stegemann Lohnverpackung & Logistischer Service e.K.
3. Address(es) of manufacturing site(s)	Stegemann Lohnverpackungen & Logistischer Service e. K Up`n Nien Esch 14 48268 Greven
4. Legally registered address of authorisation holder	Up`n Nien Esch 14 48268 Greven
5. Scope of authorisation and dosage forms	ANNEX 1
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	Ricarda Müller
8. Signature	On behalf
9. Date	13/04/2018
10. Annexes attached	Annex 1 Annex 7 (Date of inspection on which authorisation granted, scope of last inspection)

**SCOPE OF AUTHORISATION**

**Annex 1**

Name and address of the site:

Stegemann Lohnverpackungen & Logistischer Service e. K, Up'n Nien Esch 14, 48268 Greven

Veterinary Medicinal Products
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<b>AUTHORISED OPERATIONS</b> Manufacturing Operations (according to part 1)
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<b>Part 1 - MANUFACTURING OPERATIONS</b>
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<b>1.5</b>	<b>Packaging</b>
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	<i>1.5.2 Secondary packing</i>
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Any restrictions or clarifying remarks related to the scope of these Manufacturing operations

to 1.5.2: Includes also labeling
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Date of Inspection on which  
authorisation was granted

25/01/2018

Scope of last Inspection

General GMP Inspection