Regierungspräsidium Darmstadt

CERTIFICATE NUMBER : **DE_HE_01_GMP_2022_0003**

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

Part 1

Issued following an inspection in accordance with:

The competent authority of Germany confirms the following:

The manufacturer : Evonik Operations GmbH

Site address: Kirschenallee 45, Darmstadt, Hessen, 64293, Germany

OMS Location:

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. *DE HE 01 MIA 2020 0062* in accordance with Art. 13 of Directive 2001/20/EC.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on *2021-10-21*, it is considered that it complies with:

- The principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/EC³
- The principles and guidelines of Good Manufacturing Practice laid down in Directive 91/412/EC³

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than three years have elapsed since the date of that inspection. However, this period of validity may be reduced or extended using regulatory risk management principles by an entry in the Restrictions or Clarifying remarks field. This certificate is valid only when presented with all pages and both Parts 1 and 2. The authenticity of this certificate may be verified in EudraGMDP. If it does not appear, please contact the issuing authority.

Online EudraGMDP, Ref key:143664

¹ The certificate referred to in paragraph 111(5) of Directive 2001/83/EC and 80(5) of Directive 2001/82/EC, shall also be required for imports coming from third countries into a Member State.

² Guidance on the interpretation of this template can be found in the Help menu of EudraGMDP database.

³ These requirements fulfil the GMP recommendations of WHO.

Part 2

1 MA	NUFAC	TURING OPERATIONS	
1.2	Non-sterile products		
	1.2.1	on-sterile products (processing operations for the following dosage forms)	
		1.2.1.1 Capsules, hard shell	
		1.2.1.2 Capsules, soft shell	
		1.2.1.8 Other solid dosage forms	
		1.2.1.13 Tablets	
	1.2.2	Batch certification	
1.5	Packaging		
	1.5.1	Primary Packaging	
		1.5.1.1 Capsules, hard shell	
		1.5.1.2 Capsules, soft shell	
		1.5.1.8 Other solid dosage forms	
		1.5.1.13 Tablets	
	1.5.2	Secondary packaging	
1.6	Quality control testing		
	1.6.3	Chemical/Physical	

Clarifying remarks (for public users)

Ref. to 1.2.1.1: Also filling with extracts of animal origin (Antibody from goat's milk extract) Ref. to 1.2.1.2: Coating only Ref. to 1.2.1.8: modified released solid dose forms: - pellets - granules - powder

Name and signature of the authorised person of the Competent Authority of

Confidential
Regierungspräsidium Darmstadt
Tel: Confidential

Fax: Confidential