Ministerium für Soziales, Gesundheit, Frauen und Familie

CERTIFICATE NUMBER : ***DE_SL_01_GMP_2021_0030***

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER [1, 2]

Part 1

Issued following an inspection in accordance with :
Art. 111(5) of Directive 2001/83/EC as amended
Art. 15 of Directive 2001/20/EC

The competent authority of Germany confirms the following:
The manufacturer :***Eurofins PHAST GmbH***
Site address :***Kardinal-Wendel-Strasse 16, Homburg, Saarland, 66424, Germany***

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. ***DE_SL_01_MIA_2021_0025*** in accordance with Art. 40 of Directive 2001/83/EC and Art. 13 of Directive 2001/20/EC .

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on ***2020-10-22*** , it is considered that it complies with :

- The principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/EC [3]

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.

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[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

Human Medicinal Products
Human Investigational Medicinal Products

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 (list of product types)**
	1.3.2 Batch Certification (list of product types) 1.3.2.5 Biotechnology products
1.5	**Packaging**
	1.5.2 Secondary packaging
1.6	**Quality control testing**
	1.6.3 Chemical/Physical *1.6.4 Biological*

2 IMPORTATION OF MEDICINAL PRODUCTS	
2.1	**Quality control testing of imported medicinal products**
	2.1.3 Chemical/Physical *2.1.4 Biological*
2.2	**Batch certification of imported medicinal products**
	2.2.1 Sterile products 2.2.1.1 Aseptically prepared
	2.2.2 Non-sterile products
	2.2.3 Biological medicinal products 2.2.3.5 Biotechnology products

Clarifying remarks (for public users)

Secondary packaging of clinical test samples by contract manufacturers Import of patches with the active pharmaceutical ingredient Rivastigmin Import of lyophilisate with the active ingredient metreleptin (Myalepta) Powder for making an injection, Approval numbers: EU / 1/18/1276/001, 11.3 mg, 1 vial EU / 1/18/1276/002, 11.3 mg, 30 vials EU / 1/18/1276/003, 3 mg, 1 vial EU / 1/18/1276/004, 3 mg, 30 vials EU / 1/18/1276/005, 5.8 mg, 1 vial EU / 1/18/1276/006, 5.8 mg, 30 vials

2021-07-12

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

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Confidential
Ministerium für Gesundheit und Verbraucherschutz
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Fax: ***Confidential***