Competent Regional Authority Direccion De Regulacion Planificacion Y Recursos Sanitarios Departamento De Salud Generalitat De Catalunya

CERTIFICATE NUMBER: ***NCF/2320/001/CAT***

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. 63 of Regulation (EU) 536/2014

The competent authority of Spain confirms the following:
The manufacturer: ***Laboratorios Rubio S.A.***
Site address: ***Calle Industria 29, Poligono Industrial Comte De Sert, Castellbisbal, 08755***
OMS Organisation Id. / OMS Location Id.: ***ORG-100001528*** / ***LOC-100002027***

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. ***0402*** in accordance with Art. 40 of Directive 2001/83/EC and Art. 61 of Regulation (EU) No 536/2014.

Other
artículo 63, Real Decreto Legislativo 1/2015, de 24 de julio, Real Decreto 824/2010, de 25 de junio, Reglamento Delegado (UE) 2017/1569, Directiva (UE) 2017/1572

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on ***2022-09-19***, it is considered that it complies with:

- The principles and guidelines of Good Manufacturing Practice laid down in Directive (EU) 2017/1572 and Commission Delegated Regulation (EU) 2017/1569 [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.

[1] *The certificate referred to in paragraph Art. 111(5) of Directive 2001/83/EC and Art. 15 of Directive 2001/20/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**
	1.2.1 Non-sterile products (processing operations for the following dosage forms) 1.2.1.1 Capsules, hard 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 1.5.1.8 Other solid dosage forms 1.5.1.13 Tablets
	1.5.2 Secondary packaging
1.6	**Quality control testing**
	1.6.2 Microbiological: non-sterility *1.6.3 Chemical/Physical*

Clarifying remarks (for public users)

[F/HV]Manufacture and storage of products with narcotics/psicotropics is certified.; [F/I]Manufacture and storage of products with narcotics/psicotropics is certified.;

2023-04-13

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

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Confidential
Competent Regional Authority. Dirección de Regulación, Planificación y Recursos Sanitarios. Departamento de Salud. Generalitat de Catalunya
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