

Italian Medicines Agency

CERTIFICATE NUMBER: **IT-API/198/H/2024**

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

The competent authority of Italy confirms the following:

The manufacturer: **Advaxia Biologics S.r.l. In Forma Abbreviata Advaxia S.r.l.**

Site address: **Via Pontina Vecchia Km 30600, Pomezia, 00071**

OMS Organisation Id. / OMS Location Id.: **ORG-100022033 / LOC-100030741**

Is an active substance manufacturer that has been inspected in accordance with Art. 111(1) of Directive 2001/83/EC.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **2024-02-05**, it is considered that it complies with::

- The principles of GMP for active substances³ referred to in Article 47 of Directive 2001/83/EC.

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. Updates to restrictions or clarifying remarks can be identified through the EudraGMDP website (<http://eudragmdp.ema.europa.eu/>).

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.

¹The certificate referred to in paragraph Art. 111(5) of Directive 2001/83/EC is also applicable to importers.

²Guidance on the interpretation of this template can be found in the Interpretation of the Union format for GMP certificate.

³These requirements fulfil the GMP recommendations of WHO.

Part 2

Human Medicinal Products

Clarifying remarks (for public users)

Manufactured active substances marked as confidential are for clinical use only. Production and storage of MCB/WCB/EOP and MVB/WVB. According to Italian legislation, all the biological active substances and/or active substances deriving from human and animal tissues, organs, fluids are authorized according to art. 40 of Dir. 2001/83/EC.

2025-01-15

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

Confidential
Italian Medicines Agency
Tel: *Confidential*
Fax: *Confidential*