

Medical Products Agency

CERTIFICATE NUMBER :5.9.1-2021-011687

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. 80(5) of Directive 2001/82/EC as amended

Art. 15 of Directive 2001/20/EC

The competent authority of Sweden confirms the following:

The manufacturer: **Tamro AB**

Site address: **Importgatan 18-20, Hisings Backa, 422 46, Sweden**

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. **5.9.1-2021-011687** in accordance with Art. 40 of Directive 2001/83/EC , Art. 44 of Directive 2001/82/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 **2018-10-04** , 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.

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
Veterinary Medicinal Products
Human Investigational Medicinal Products

1 MANUFACTURING OPERATIONS	
1.5	Packaging
	<i>1.5.2 Secondary packaging</i>

2 IMPORTATION OF MEDICINAL PRODUCTS	
2.3	Other importation activities
	<i>2.3.1 Site of physical importation</i>

Clarifying remarks (for public users)

1.5.2 Re-packaging and re-labelling as well as packaging and labelling of secondary packaging. (Ompackning och ommärkning samt packning och märkning i sekundärförpackning.) Re-labelling and additional labelling for clinical trials. (Ommärkning och tilläggsmärkning för klinisk prövning.) 2.3.1 Quarantine and sampling after importation from 3rd country. (Karantänhållning och provtagning efter import från 3:e land.)

2021-03-04

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



Bengt Berglund

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