MALTA MEDICINES AUTHORITY

IN012/12 Appendix 1 Version 2

Certificate No: MT/030HM/2026

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/
The Malta Medicines Authority confirms the following:
The manufacturer **Pharmadox Healthcare Ltd**
Site address **HHF 303, Hal Far Industrial Estate , Hal Far Birzebbugia BBG3000**

Has been inspected under the national inspection programme in connection with manufacturer's licence no. **ML 013** in accordance with Art. 40 of Directive 2001/83/EC transposed in the following national legislation: Medicines Act 2003 (Ch.458) Part III Title II Art 37, Regulation 3 of Subsidiary Legislation 458.36

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **25th March 2026**, it is considered that it complies with the The principles and guidelines of Good Manufacturing Practice laid down in Directive (EU) 2017/1572[3] and Commission Delegated Regulation (EU) 2017/1569as appropriate.

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.

2nd April 2026

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Dr. Mark Cilia[1]
Director Inspectorate & Enforcement Directorate
Malta Medicines Authority
Tel: 00356 234 39 119

[1] The signature, date and contact details should appear on each page of the certif

[1] The certificate referred to in paragraph 111(5) of Directive 2001/83/EC is also applicable to importers.
[2] Guidance on the interpretation of this template can be found in the Interpretation of the Union format for GMP cert

Part 2

❑ Human Medicinal Products*	
1 MANUFACTURING OPERATIONS – MEDICINAL PRODUCTS *	
1.4	**Other products or processing activity**
	1.4.3 Others (free text) Storage of medicinal products

Any restrictions or clarifying remarks related to the scope of this certificate:
None

2nd April 2026 .....

Dr. Mark Cilia[1]
Director Inspectorate & Enforcement Directorate
Malta Medicines Authority
Tel: 00356 234 39 119

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