

Malta Medicines Authority

Report No: 002/24/MC/kg

STATEMENT OF NON-COMPLIANCE WITH GMP

Exchange of information between National Competent Authorities (NCAs) of the EEA following the discovery of serious GMP non-compliance at a manufacturer¹

Part 1

Issued following an inspection in accordance with
Art. 111(7) of Directive 2001/83/EC as amended

The competent authority of Malta confirms the following:

The manufacturer: **Cubit Lifesciences Limited Liability Partnership**

Site address: **Plot No 39 - 40, Nr Kerala Gidc Tal, Ozone Industrial Park, Bhayla, Gujarat, 382220**

OMS Organisation Id. / OMS Location Id.: **ORG-100047686 / LOC-100078794**

DUNS Number: **86-140-7631**

Other

(Human) Human Medicinal Products

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **2024-02-01**, it is considered that **it does not comply with the Good Manufacturing Practice** requirements referred to in

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Note to receiving authorities: Please contact the issuing authority within 20 working days in case there are critical(2) medicinal products potentially affected by this statement.

Manufacturing Authorisation Holders directly affected by this statement have failed to comply with their obligations under Art. 46 of Directive 2001/83/EC or Art. 93(1)(j) to (l) of Regulation (EU) 2019/6 and as a consequence the Qualified Person referred to in Art. 48 of Directive 2001/83/EC and Art. 97(1) of Regulation (EU) 2019/6 is unable to perform the batch certification referred to in Art. 51 of Directive 2001/83/EC and Art. 97 (6) and (7) of Regulation (EU) 2019/6.

In exceptional circumstances there may be no objection to the Qualified Person certifying affected batches thereby allowing their release provided all of the following conditions are fulfilled:

1. Batch certification is performed in order to maintain supply of critical medicinal products only.
2. A documented risk assessment has been performed by, or on behalf of, the Qualified Person and additional actions have been implemented by the manufacturing and/or batch release site to mitigate the risks posed by the non-compliance. Note: Repeated testing alone is not normally sufficient risk mitigation but, together with other actions, can form part of a strategy commensurate with the nature and the level of risk.
3. A thorough risk-benefit evaluation has been performed for the acceptance of risk and a report prepared that takes full account of the nature of the non-compliance with the involvement of:
 - The Manufacturing Authorisation Holder and the Qualified Person of the site responsible for

batch certification.

- The manufacturing site subject to this Statement of Non-Compliance, if different from the above.
- The relevant Marketing Authorisation Holder(s).

The report has been shared with the National Competent Authorities of the countries in which distribution of the affected batches is anticipated and that any comments from those authorities have been taken into account.

4. Written confirmation has been obtained from the National Competent Authorities in whose territories the affected batches are intended to be distributed that the product is considered critical on its territory, and that there is no objection to distribution.
5. The Supervisory Authority has been informed, if different from the above, and it has not suspended or revoked the relevant Manufacturing Authorisation.
6. The affected Marketing Authorisations have not been revoked or suspended.
7. Any further conditions imposed by the Supervisory Authority and other involved National Competent Authorities are met.

¹The statement of non-compliance referred to in paragraph 111(7) of Directive 2001/83/EC and Art. 94(2) of Regulation (EU) 2019/6, as amended, is also applicable to importers.

²See Appendix 3 of the relevant procedure in the Compilation of Union Procedures.

Part 2

Human Medicinal Products	
1 NON-COMPLIANT MANUFACTURING OPERATIONS	
Include total and partial manufacturing (including various processes of dividing up, packaging or presentation), batch release and certification, storage and distribution of specified dosage forms unless informed to the contrary;	
1.2	Non-sterile products
	<i>1.2.1 Non-sterile products (processing operations for the following dosage forms)</i> 1.2.1.1 Capsules, hard shell 1.2.1.13 Tablets
1.5	Packaging
	<i>1.5.1 Primary Packaging</i> 1.5.1.1 Capsules, hard shell 1.5.1.13 Tablets
	<i>1.5.2 Secondary packaging</i>
1.6	Quality control testing
	<i>1.6.2 Microbiological: non-sterility</i> <i>1.6.3 Chemical/Physical</i>

Clarifying remarks (for public users)

None

Part 3

1.Nature of non-compliance:
A follow-up GMP inspection was conducted at the aforementioned site by the MMA, as an EU Competent Authority, executed during the period of 27th January 2024 – 1st February 2024. The inspection focus, as a follow up inspection, was aligned with the recommendations stemming from a previous GMP inspection conducted between 18th – 23rd February 2023. The previous inspection had resulted in the identification of 2 critical, 3 major and a total of 17 other observed issues. The principal objective of the follow-up inspection was to assess and validate the effective incorporation, spanning the entirety of the facility and the pharmaceutical quality system, of the documented Corrective and Preventive Actions (CAPAs) and commitments presented by Cubit Lifesciences in response to the February 2023 inspection. During the follow-up inspection, the inspectors found two critical, four major and twelve other findings, some of which were already identified in the initial February 2023 inspection and still were found not fully addressed and implemented in the follow-up inspection, indicating an improper implementation of all CAPAs throughout the whole facility. As a result of this outcome, the Inspection Review Group (IRG) at the Malta Medicines Authority has met and decided that a Statement of Non-Compliance with the principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/EC is to be issued for the site.
Action taken/proposed by the NCA
Others

The manufacturer did not previously have EU GMP certification. An EU GMP certificate will not be issued.

Additional comments

Draft supervisory risk assessment has been circulated through the rapid alert network for any comments by NCAs with deadline for responses set to the 27th February 2024 (1 week from date of issue of the draft supervisory risk assessment). A total of 1 reply was received, confirming that there are no medicinal products on the EU market and no critical medicinal products on MRA partner markets. The Malta Medicines Authority issued a statement of GMP non-compliance on 4th March 2024. The 3rd country finished product manufacturer was informed of this decision accordingly.

2024-03-07

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

Confidential
Malta Medicines Authority
Tel: **Confidential**
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