# Medicines and Healthcare products Regulatory Agency

## CERTIFICATE NUMBER : UK MIA 58711 Insp GMP 58711/1966883-0010[H]

## CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER*(1),(2)*

# Part 1

Issued following an inspection in accordance with :
Regulation 331A of The Human Medicines Regulations 2012 (SI 2012/1916)

The competent authority of United Kingdom confirms the following :

The Manufacturer : VANTIVE LIMITED

Site address : VANTIVE LIMITED, 2 WAVERTREE BOULEVARD, WAVERTREE TECHNOLOGY PARK, LIVERPOOL, L7 9PE, UNITED KINGDOM

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. UK MIA 58711 in accordance with Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 10/02/2026 , it is considered that it complies with

- The principles and guidelines of Good Manufacturing Practice laid down in Regulation B17 of the Human Medicines Regulations 2012 (as amended)

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 MHRA-GMDP. If it does not appear, please contact the issuing authority.

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- *(1) Guidance on the interpretation of this template can be found in the Help menu of MHRA-GMDP database.*
- *(2) These requirements fulfil the GMP recommendations of WHO.*

# Part 2

**Human Medicinal Products**

**2. IMPORTATION OF MEDICINAL PRODUCTS**

**[ 2.3 ] Other Importation Activities**

[ 2.3.4 ] Other
Physical importation from an EEA country with financial payment to a Swiss entity

**Restrictions or Remarks**
This certificate is issued based on a desk-based assessment of GMP compliance information provided by the manufacturer. This certificate should be used in combination with the relevant authorisation/registration. A risk-based site inspection programme remains in force.

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| 10/02/2026 | Name and signature of the authorised person of the Competent Authority of United Kingdom<br>Confidential<br>Medicines and Healthcare products Regulatory Agency<br>Tel : Confidential |

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