# Medicines and Healthcare products Regulatory Agency

## CERTIFICATE NUMBER : UK GMP 22481 Insp GMP 22481/366976-0005[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 : MICRO LABS LIMITED

Site address : MICRO LABS LIMITED, 16 VEERASANDRA INDUSTRIAL AREA, ANEKAL TALUK, BANGALORE, IN-560 100, INDIA

Has been inspected in connection with marketing authorisation(s) listing manufacturers located outside of the European Economic Area in accordance with Part 16 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 23/09/2019, 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**

**1. MANUFACTURING OPERATIONS**

**[ 1.2 ] Non-sterile products**

[ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms)

[ 1.2.1.1 ] Capsules, hard shell

[ 1.2.1.13 ] Tablets

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[ 1.2.1.17 ] Other non-sterile medicinal products

Oral powder for reconstruction

**[ 1.5 ] Packaging**

[ 1.5.2 ] Secondary packaging

**[ 1.6 ] Quality control testing**

[ 1.6.2 ] Microbiological: non-sterility

[ 1.6.3 ] Chemical/Physical

**Restrictions or Remarks**

This certificate is valid only for building ML11 on Plot 16 and the Plot 24 warehouse expansion (which included storage areas, sampling and dispensary). This certificate does not include the manufacturing areas in Plot 24 expansion first floor as these were incomplete at the time of inspection. This certificate is restricted to storage and sampling activities in the adjacent ML 25 building. This certificate does not include manufacturing in ML 25.

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| 24/04/2020 | 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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