Medicines and Healthcare products Regulatory Agency

CERTIFICATE NUMBER : UK EXC 204 Insp GMP 204/92237-0010[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 : COLORCON LIMITED

Site address : COLORCON LIMITED, FLAGSHIP HOUSE, VICTORY WAY, CROSSWAYS BUSINESS PARK, DARTFORD, DA2 6QD, UNITED KINGDOM

Has been inspected under the national inspection programme in accordance with Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916)

Is an active substance manufacturer that has been inspected in accordance with Regulation 327 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 03/12/2014, 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)
- The principles of GMP for active substances referred to in Regulation 327 of the Human Medicines Regulations 2012 (SI 2012/1916)

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.

- *(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

Any restrictions related to the scope of this certificate:

Building	Room	Line/equipment	QC Testing	Products
				Excipient Dispersions for Oral Solid Dosage

MHRA-GMDP MHRA

22/12/2014	Name and signature of the authorised person of the Competent Authority of United Kingdom Confidential Medicines and Healthcare products Regulatory Agency Tel : Confidential

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