Medicines and Healthcare products Regulatory Agency

Certificate No: UK API 59275 Insp GDP 59275/36281215-0001

CERTIFICATE OF GDP COMPLIANCE OF A DISTRIBUTOR OF ACTIVE SUBSTANCES FOR USE AS STARTING MATERIALS IN MEDICINAL PRODUCTS FOR HUMAN USE

Issued following an inspection in accordance with Regulation 331 (4) (b) of the Human Medicines Regulations 2012

The wholesale distributor : AIT WORLDWIDE LOGISTICS (UK) LIMITED
Site address : AIT WORLDWIDE LOGISTICS (UK) LIMITED, UNIT 22, ASHFORD INDUSTRIAL ESTATE, SHIELD ROAD, ASHFORD, TW15 1AU, UNITED KINGDOM

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

From the knowledge gained during inspection of this active substance distributor, the latest of which was conducted on 04/03/2025, it is considered that it complies with the principles of good distribution practice for active substances referred to in Regulation B17 and C17 of The Human Medicines Regulations 2012 (SI 2012/1916).

This certificate reflects the status of the premises at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than five years have elapsed since the date of that inspection. However this period of validity may be reduced 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.

The authenticity of this certificate may be verified in MHRA-GMDP. If it does not appear please contact the issuing authority.

Any restrictions or clarifying remarks related to the scope of this certificate(For All Users):

In accordance with the MHRA risk based inspection procedures this site, which was the subject of a new API Registration, will be re-inspected within 24 months of the initial inspection. This certificate should not be relied upon to reflect the compliance status if more than 24 months have elapsed since the date of that initial inspection.

07/05/2025	Name and signature of the authorised person of the Competent Authority of United Kingdom
	Confidential
	Medicines and Healthcare products Regulatory Agency
	Confidential