Medicines and Healthcare products Regulatory Agency

WHOLESALE DISTRIBUTION AUTHORISATION

(MEDICINAL PRODUCTS FOR HUMAN USE)

1. Authorisation Number	UK WDA(H) 56111
2. Name of Authorisation Holder	SUPER B PLUS GROUP LTD
3. Legally registered address of Authorisation Holder	SUPER B PLUS GROUP LTD, RIVERSIDE BUSINESS CENTRE, FORT ROAD, TILBURY, RM18 7ND, UNITED KINGDOM
4. Address(es) of Site(s)	SUPER B PLUS GROUP LTD, OFFICE 2, UNIT C1/B, 30 BRUNEL ROAD, MANOR TRADING ESTATE, BENFLEET, SS7 4PS, UNITED KINGDOM
5. Scope of authorisation (complete for each site under 4)	ANNEX 1
6. Legal basis of authorisation	Regulation 18 of the Human Medicines Regulations 2012
7. Name of responsible officer of the competent authority of the member state granting the wholesaling authorisation	Confidential
8. Date	10/11/2025
9. Annexes attached	Annex 1 Scope of wholesale distribution authorisation Annex 2 (Optional) Address(es) of contract wholesale distribution sites and their authorisation number Annex 3 (Optional) Name(s) of responsible person(s) Annex 4 (Optional) Date of Inspection on which authorisation was granted Annex 5 Additional provisions

ANNEX 1

SCOPE OF WHOLESALE DISTRIBUTION AUTHORISATION

Name and address of the site:

SUPER B PLUS GROUP LTD, OFFICE 2, UNIT C1/B, 30 BRUNEL ROAD, MANOR TRADING ESTATE, BENFLEET, SS7 4PS, UNITED KINGDOM

1. MEDICINAL PRODUCTS

1.1 With “an authorisation” (a UK, Great Britain or Northern Ireland Marketing Authorisation, an Article 126a authorisation, a certificate of registration or traditional herbal registration)

1.2 Without “an authorisation” (a UK, Great Britain or Northern Ireland Marketing Authorisation, an Article 126a authorisation, a certificate of registration or traditional herbal registration) and intended for the UK market

1.3 Without “an authorisation” (a UK, Great Britain or Northern Ireland Marketing Authorisation, an Article 126a authorisation, a certificate of registration or traditional herbal registration) and not intended for the UK market

2. AUTHORISED WHOLESALE DISTRIBUTION OPERATIONS

MHRA-GMDP MHRA-GMDP MHRA

2.1 Procurement

2.3 Supply

2.4 Export

2.6 Products imported from countries on a list

2.6b Products not certified under Article 51 of Directive 2001/83/EC

3. MEDICINAL PRODUCTS WITH ADDITIONAL REQUIREMENTS

3.1.2 Medicinal products derived from blood

3.1.3 Immunological medicinal products

3.3 Cold chain products (requiring low temperature handling)

Any restrictions or clarifying remarks (for all users)

4 Categories of Products Handled at this Site: 4.1 Prescription Only Medicines, 4.2 General Sales List, 4.4 Pharmacy

MHRA-GMDP MHRA MHRA-GMDP MHRA