

Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number UK MIA(IMP) 56952

2: Name of authorisation holder PNR PHARMA CONSULTING LIMITED

3: Address(es) of manufacturing site(s) PNR PHARMA, UPPER GROUND FLOOR ROOM 2, CREUDDYN, PONTFAEN ROAD, LAMPETER, SA48 7BN, UNITED KINGDOM

4: Legally registered address of authorisation holder PNR PHARMA CONSULTING LTD, UNITS 5 AND 6, LIME AVENUE, EBBW VALE, NP23 6GL, UNITED KINGDOM

5: Scope of authorisation and dosage forms ANNEX 1 and/ or ANNEX 2

6: Legal Basis of authorisation Part 6 of The Medicines for Human Use (Clinical Trials) Regulations 2004 [SI 2004/1031]

7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation Confidential

8: Authorisation Date 23/10/2023

9: Annexes attached Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 2

Name and address of the site:

PNR PHARMA, UPPER GROUND FLOOR ROOM 2, CREUDDYN, PONTFAEN ROAD, LAMPETER, SA48 7BN, UNITED KINGDOM

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| Human Investigational Medicinal Products |
| Authorised Operations |
| IMPORTATION OF MEDICINAL PRODUCTS (according to part 2) |
| Part 2 - IMPORTATION OF MEDICINAL PRODUCTS [2.3] Other Importation Activities [2.3.4] Other Importation of QP-certified IMPs from a country on the approved country for import list |

SCOPE OF AUTHORISATION

Annex 2

Name and address of the site:

PNR PHARMA CONSULTING LTD, UNITS 5 AND 6, LIME AVENUE, EBBW VALE, NP23 6GL, UNITED KINGDOM

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| Human Investigational Medicinal Products |
| Authorised Operations |
| MANUFACTURING OPERATIONS (according to part 1) IMPORTATION OF MEDICINAL PRODUCTS (according to part 2) |

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| <p>Part 1 - MANUFACTURING OPERATIONS</p> <p>[1.1] Sterile Investigational Medicinal Products</p> <p>[1.1.3] Batch certification</p> <p>[1.2] Non-sterile investigational medicinal products</p> <p>[1.2.2] Batch certification</p> <p>[1.3] Biological investigational medicinal products</p> <p>[1.3.2] Batch certification</p> <p>[1.3.2.1] Blood products</p> <p>[1.3.2.2] Immunological products</p> <p>[1.3.2.3] Cell therapy products</p> <p>[1.3.2.4] Gene therapy products</p> <p>[1.3.2.5] Biotechnology products</p> <p>[1.3.2.6] Human or animal extracted products</p> <p>[1.5] Packaging</p> <p>[1.5.2] Secondary packaging</p> <p>Part 2 - IMPORTATION OF MEDICINAL PRODUCTS</p> <p>[2.2] Batch certification of imported medicinal products</p> <p>[2.2.1] Sterile Products</p> <p>[2.2.1.1] Aseptically prepared</p> <p>[2.2.1.2] Terminally sterilised</p> <p>[2.2.3] Biological medicinal products</p> <p>[2.2.3.1] Blood products</p> <p>[2.2.3.2] Immunological products</p> <p>[2.2.3.3] Cell therapy products</p> <p>[2.2.3.4] Gene therapy products</p> <p>[2.2.3.5] Biotechnology products</p> <p>[2.2.3.6] Human or animal extracted products</p> <p>[2.3] Other Importation Activities</p> <p>[2.3.1] Site of Physical Importation</p> <p>[2.3.2] Importation of Intermediate which undergoes further processing</p> <p>[2.3.3] Biological Active Substance</p> |
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