Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

- 1: Authorisation Number
- 2: Name of authorisation holder
- 3: Address(es) of manufacturing site(s)
- 4: Legally registered address of authorisation holder
- 5: Scope of authorisation and dosage forms
- 6: Legal Basis of authorisation
- 7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation
- 8: Authorisation Date
- 9: Annexes attached

UK MIA 42760

ARCADIA PHARMA LIMITED

ARCADIA PHARMA LIMITED, UNIT 3, BELL COURT, SWANSEA WEST BUSINESS PARK, FELINFACH, FFORESTFACH, SWANSEA, SA5 4HP, UNITED KINGDOM

ARCADIA PHARMA LIMITED, UNIT 3, BELL COURT, SWANSEA WEST BUSINESS PARK, FELINFACH, FFORESTFACH, SWANSEA, SA5 4HP, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916)

Confidential

04/06/2025 Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1 Name and address of the site:

ARCADIA PHARMA LIMITED, UNIT 3, BELL COURT, SWANSEA WEST BUSINESS PARK, FELINFACH, FFORESTFACH, SWANSEA, SA5 4HP, UNITED KINGDOM

Human Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)

Part 1 - MANUFACTURING OPERATIONS

[1.5] Packaging

[1.5.2] Secondary packaging