# 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 43801** 

SCIECURE PHARMA LIMITED

SCIECURE PHARMA LIMITED, 5 MILLMEAD, GUILDFORD, GU2 4BE, UNITED KINGDOM

SCIECURE PHARMA LIMITED, 5 MILLMEAD, GUILDFORD, GU2 4BE, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations

2012 (SI 2012/1916)

Confidential

26/05/2023

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

### Annex 1

Name and address of the site:

## SCIECURE PHARMA LIMITED, 5 MILLMEAD, GUILDFORD, GU2 4BE, UNITED KINGDOM

**Human Medicinal Products** 

**Authorised Operations** 

IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

### [ 2.2 ] Batch certification of imported medicinal products

[2.2.1] Sterile Products

[2.2.1.2] Terminally sterilised

[2.2.2] Non-sterile products

Issue Date: 26 May 2023

Page 1 of 1