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(IMP) 1811 BARD PHARMACEUTICALS LIMITED

BARD PHARMACEUTICALS LIMITED, UNIT 191, CAMBRIDGE SCIENCE PARK, MILTON ROAD, CAMBRIDGE, CB4 0GW, UNITED KINGDOM

BARD PHARMACEUTICALS LIMITED, UNIT 198, CAMBRIDGE SCIENCE PARK, MILTON ROAD, CAMBRIDGE, CB4 0AB, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

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

Confidential

21/04/2023

Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 2 Name and address of the site:

BARD PHARMACEUTICALS LIMITED, UNIT 191, CAMBRIDGE SCIENCE PARK, MILTON ROAD, CAMBRIDGE, CB4 0GW, UNITED KINGDOM

Human Investigational Medicinal Products	
Authorised Operations	
MANUFACTURING OPERATIONS (according to part 1)	<u> </u>
Part 1 - MANUFACTURING OPERATIONS	
[1.2] Non-sterile investigational medicinal products	
[1.2.1] Non-Sterile Products (processing operations for the following dosage forms)	
[1.2.1.1] Capsul <mark>es</mark> , hard shell	
[1.2.1.13] Tablets	
[1.6] Quality control testing	
[1.6.3] Chemical/Physical	