

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA(IMP) 53174
<b>2: Name of authorisation holder</b>	REACTA BIOTECH LIMITED
<b>3: Address(es) of manufacturing site(s)</b>	REACTA BIOTECH LIMITED T/A REACTA HEALTHCARE, 2 NEWTECH SQUARE, DEESIDE INDUSTRIAL PARK, DEESIDE, CH5 2NT, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	REACTA BIOTECH LIMITED, 2 NEWTECH SQUARE, DEESIDE INDUSTRIAL PARK, DEESIDE, CH5 2NT, UNITED KINGDOM
<b>5: Scope of authorisation and dosage forms</b>	ANNEX 1 and/ or ANNEX 2
<b>6: Legal Basis of authorisation</b>	Part 6 of The Medicines for Human Use (Clinical Trials) Regulations 2004 [SI 2004/1031]
<b>7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation</b>	Confidential
<b>8: Authorisation Date</b>	10/09/2024
<b>9: Annexes attached</b>	Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 2

Name and address of the site:

**REACTA BIOTECH LIMITED T/A REACTA HEALTHCARE, 2 NEWTECH SQUARE, DEESIDE INDUSTRIAL PARK, DEESIDE, CH5  
2NT, UNITED KINGDOM**

Human Investigational Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1)
<b>Part 1 - MANUFACTURING OPERATIONS</b> <b>[ 1.2 ] Non-sterile investigational medicinal products</b> [ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms) [ 1.2.1.11 ] Semi-solids <b>[ 1.5 ] Packaging</b> [ 1.5.1 ] Primary packaging [ 1.5.1.11 ] Semi-solids [ 1.5.2 ] Secondary packaging