# 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 89

CUXSON GERRARD AND COMPANY LIMITED

CUXSON GERRARD & COMPANY LIMITED, 125 BROADWELL ROAD, OLDBURY, B69 4BF, UNITED KINGDOM

CUXSON GERRARD AND COMPANY LIMITED, 125 BROADWELL ROAD, OLDBURY, B69 4BF, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

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

Confidential

01/06/2022

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

Annex 1 Name and address of the site:

### CUXSON GERRARD & COMPANY LIMITED, 125 BROADWELL ROAD, OLDBURY, B69 4BF, UNITED KINGDOM

Human Medicinal Products	
Authorised Operations	
MANUFACTURING OPERATIONS (according to part 1)	
Part 1 - MANUFACTURING OPERATIONS	~
[ 1.2 ] Non-sterile products	
[1.2.1] Non-Sterile Products (processing operations for the following dosage forms)	
[ 1.2.1.5 ] Liquids for external use	
[ 1.2.1.8 ] Other solid dosage forms	
[ 1.2.1.11 ] Semi-solids	
[ 1.5 ] Packaging	
[ 1.5.2 ] Secondary packaging	
[ 1.6 ] Quality control testing	

