# 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

Issue Date: 01 Jun 2022

