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

SCOPE OF AUTHORISATION

Annex 1 Name and address of the site:

SYRI LIMITED, UNIT 4, BRADFIELD ROAD, RUISLIP, HA4 0NU, UNITED KINGDOM

Human Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1) IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

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
 - [1.2.1.6] Liquids for internal use
- [1.2.2] Batch certification

[1.5] Packaging

[1.5.1] Primary packaging

[1.5.1.5] Liquids for external use

[1.5.1.6] Liquids for internal use

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ANNEX 1 and/ or ANNEX 2

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

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Annex 1 and/or Annex 2

- [1.5.2] Secondary packaging
- [1.6] Quality control testing
 - [1.6.2] Microbiological: non-sterility
 - [1.6.3] Chemical/Physical
- Part 2 IMPORTATION OF MEDICINAL PRODUCTS

[2.1] Quality control testing of imported medicinal products

- [2.1.2] Microbiological: non-sterility
- [2.1.3] Chemical/Physical
- [2.2] Batch certification of imported medicinal products
- [2.2.2] Non-sterile products
- [2.3] Other Importation Activities
- [2.3.1] Site of Physical Importation