

Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

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|---|---|
| 1: Authorisation Number | UK MIA 14340 |
| 2: Name of authorisation holder | EUROCAPS LIMITED |
| 3: Address(es) of manufacturing site(s) | EUROCAPS LIMITED, UNITS A, B, C & D, CROWN BUSINESS PARK, DUKESTOWN, TREDEGAR, NP22 4EF, UNITED KINGDOM |
| 4: Legally registered address of authorisation holder | EUROCAPS LIMITED, UNITS A, B, C & D, CROWN BUSINESS PARK, DUKESTOWN, TREDEGAR, NP22 4EF, UNITED KINGDOM |
| 5: Scope of authorisation and dosage forms | ANNEX 1 and/ or ANNEX 2 |
| 6: Legal Basis of authorisation | Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916) |
| 7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation | Confidential |
| 8: Authorisation Date | 09/09/2024 |
| 9: Annexes attached | Annex 1 and/or Annex 2 |

SCOPE OF AUTHORISATION

Annex 1

Name and address of the site:

EUROCAPS LIMITED, UNITS A, B, C & D, CROWN BUSINESS PARK, DUKESTOWN, TREDEGAR, NP22 4EF, UNITED KINGDOM

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| 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) [1.2.1.2] Capsules, soft shell [1.4] Other products or manufacturing activity [1.4.1] Manufacture of: [1.4.1.1] Herbal products [1.6] Quality control testing [1.6.3] Chemical/Physical |