

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

**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** 11/02/2025

**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

Human Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1)
<b>Part 1 - MANUFACTURING OPERATIONS</b> <b>[ 1.2 ] Non-sterile products</b> [ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms) [ 1.2.1.2 ] Capsules, soft shell <b>[ 1.4 ] Other products or manufacturing activity</b> [ 1.4.1 ] Manufacture of: [ 1.4.1.1 ] Herbal products <b>[ 1.6 ] Quality control testing</b> [ 1.6.3 ] Chemical/Physical