

Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number	UK MIA 20093
2: Name of authorisation holder	LIFEPLAN PRODUCTS LIMITED
3: Address(es) of manufacturing site(s)	LIFEPLAN PRODUCTS LIMITED, ELIZABETHAN WAY, LUTTERWORTH, LE17 4ND, UNITED KINGDOM
4: Legally registered address of authorisation holder	LIFEPLAN PRODUCTS LIMITED, ELIZABETHAN WAY, LUTTERWORTH, LE17 4ND, 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	22/08/2024
9: Annexes attached	Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1

Name and address of the site:

LIFEPLAN PRODUCTS LIMITED, ELIZABETHAN WAY, LUTTERWORTH, LE17 4ND, 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) [1.2.1.1] Capsules, hard shell [1.2.1.13] Tablets [1.4] Other products or manufacturing activity [1.4.1] Manufacture of: [1.4.1.1] Herbal products [1.5] Packaging [1.5.1] Primary packaging

[1.5.1.1] Capsules, hard shell

[1.5.1.13] Tablets

[1.5.2] Secondary packaging

[1.6] Quality control testing

[1.6.3] Chemical/Physical

MHRA-GMDP

MHRA

MHRA-GMDP

MHRA