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(IMP) 142 ACCORD-UK LIMITED

ACCORD-UK LIMITED, WHIDDON VALLEY, BARNSTAPLE, EX32 8NS, UNITED KINGDOM

ACCORD-UK LIMITED, WHIDDON VALLEY, BARNSTAPLE, DEVON, EX32 8NS, UNITED KINGDOM ANNEX 1 and/ or ANNEX 2

Part 6 of The Medicines for Human Use (Clinical Trials) Regulations 2004 [SI 2004/1031]

Confidential

09/09/2024

Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 2 Name and address of the site:

ACCORD-UK LIMITED, WHIDDON VALLEY, BARNSTAPLE, EX32 8NS, UNITED KINGDOM

Human Investigational Medicinal Products	
Authorised Operations	
MANUFACTURING OPERATIONS (according to part 1)	
Part 1 - MANUFACTURING OPERATIONS	
[1.2] Non-sterile investigational medicinal products	
[1.2.1] Non-Sterile Products (processing operations for the following dosage for	ms)
[1.2.1.1] Capsules, hard shell	
[1.2.1.12] Suppositories	
[1.2.1.13] Tablets	
Special Requirements	
Hormonal activity Levothyroxine and Liothyronine	
[1.2.1.15] Other non-sterile medicinal products	
Pessaries / hormonal activity - Progesterone	
[1.2.2] Batch certification	
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[1.5] Packaging

[1.5.1] Primary packaging

[1.5.1.1] Capsules, hard shell

[1.5.1.12] Suppositories

[1.5.1.13] Tablets

Special Requirements

- Levothyroxine and Liothyronine
- [1.5.1.15] Other non-sterile medicinal products Pessaries and hormonal activity - Progesterone
- [1.5.2] Secondary packaging
- [1.6] Quality control testing
 - [1.6.3] Chemical/Physical