



ARTWORK

LEO Pharma A/S
Internal Market Access

Scale	Get-up	Material No	Sent by e-mail
100%	HK+MY	064962-XX	▼
Subject	Date		Date
LBL 155 x 125 mm	25/03/20		
Colour	Sign.		Sign.
PMS 485 + 7421 + black	OMA		

Preparation Strength Packsize	Enstilar® Foam 60 g	Place of production	Ireland
Comments: Mock-up for reg. purpose			

LIE025-02 – 155 x 125 mm

One gram contains 50 micrograms of calcipotriol (as monohydrate) and 0.5 mg of betamethasone (as dipropionate).

Excipients:

Liquid paraffin

Polyoxypropylene stearyl ether

All-rac- α -tocopherol

White soft paraffin

Butylhydroxytoluene (E321)

Butane

Dimethyl ether.

See leaflet for further information.

Shake can for a few seconds before use. Hold can (not horizontally) at least 3 cm from skin and spray.

Do not apply directly to the face or eyes.

Wash hands after use.

Read the package leaflet before use.

Keep out of the sight and reach of children.

Do not store above 25°C.

Lot/MAN/EXP - see the bottom of the can.

Caution:

Extremely flammable aerosol. Pressurised container.

May burst if heated. Protect from sunlight. Do not expose

to temperatures exceeding 50°C. Do not pierce or burn,

even after use. Do not spray on an open flame or other

ignition source. Keep away from sparks, open flames and

other ignition sources.

No smoking.

Hong Kong: HK-66143

Malaysia: MAL20026049AZ

LEO Laboratories

Limited

285 Cashel Road

Crumlin, Dublin 12

Ireland

Product registration holder in Malaysia: DKSH Malaysia Sdn Bhd
B-11-01, The Ascent, Paradigm No.1, Jalan SS7/26A, Kelana Jaya,
47301 Petaling Jaya, Selangor, Malaysia.

HAJI UBAT DARI KANAK-KANAK
Controlled medicine/Ubah terkawal

Enstilar®

50 micrograms/g + 0.5 mg/g
cutaneous foam

calcipotriol/betamethasone

Cutaneous use
For external use only
只供外用.

60 g

Net contents: 146.4 g/268 ml



LEO

064962-XX

LIE025-02

5. PROOF OMA FROM	Mock-up Approval Stamp (MAS)		
Date 26/05/20	Graphic Design	Editorial Proof	Second Approver
New proof requested ☐	According to: SOP_000647, SOP_000962, SOP_003993 and SOP_008676 ☐	According to: SOP_000647, SOP_000962 and SOP_008676 ☐	Product name ☐
Sign.:			Dosage form ☐
			Strength/Stripes ☐
			Pack size ☐
	1st Sign.: Date:	Sign.: Date:	Prompts ☐
			Material No./Reg. No. ☐
Date:			Barcode ☐
	2nd Sign.: Date:		Sign.: Date: