Summary of Product Characteristics

1 NAME OF THE MEDICINAL PRODUCT

Monopost Unidose 50 micrograms/ml eye drops, solution in single-dose container

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

1 ml eye drops solution contains 50 micrograms of latanoprost.

One drop contains approximately 1.5 micrograms of latanoprost.

Excipient with known effect: polyethylene glycol glycerol hydroxystearate 40 (castor oil polyoxyl hydrogenated).

For the full list of excipients, see section 6.1.

3 PHARMACEUTICAL FORM

Eye drops, solution in single-dose container.

Product imported from Greece;

The solution is a slightly yellow and opalescent solution.

pH: 6.5 – 7.5

Osmolality: 250-310 mosmol/kg

4 CLINICAL PARTICULARS

As per PA1107/007/001

5 PHARMACOLOGICAL PROPERTIES

As per PA1107/007/001

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

macrogolglycerol hydroxystearate Sorbitol Carbomer 974P Polyethylene glycol 4000 Ethylenediaminetetraacetate disodium salt Sodium hydroxide (for pH adjustment) Water for injections

6.2 Incompatibilities

Not applicable.

6.3 Shelf life

The shelf-life expiry date of this product is the date shown on the container and outer package of the product on the market in the country of origin.

After first opening of the sachet, use the single-dose containers within 10 days.

After first opening of the single-dose container: use immediately and discard the single-dose container after use.

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6.4 Special precautions for storage

Store below 25 °C.

For storage after first opening of the medicinal product, see section 6.3.

6.5 Nature and contents of container

5 or 10 single-dose containers containing 0.2 ml of eye drops solution are packed in sachet. Pack size: 30 single-dose containers.

6.6 Special precautions for disposal

No special requirements.

7 PARALLEL PRODUCT AUTHORISATION HOLDER

IMED Healthcare Ltd. Unit 625 Kilshane Avenue Northwest Business Park Ballycoolin Dublin 15 Ireland

8 PARALLEL PRODUCT AUTHORISATION NUMBER

PPA1463/153/001

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 29th January 2021

10 DATE OF REVISION OF THE TEXT

September 2022

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