Summary of Product Characteristics

1 NAME OF THE MEDICINAL PRODUCT

Zovirax 800 mg Dispersible Tablets

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each dispersible tablet contains 800 mg aciclovir

For the full list of excipients, see section 6.1.

3 PHARMACEUTICAL FORM

Dispersible Tablets

Product imported from Greece

White, oval, film-coated tablets marked with "GXCG1" on one side.

4 CLINICAL PARTICULARS

As per PA1077/084/009

5 PHARMACOLOGICAL PROPERTIES

As per PA1077/084/009

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Microcrystalline cellulose
Aluminium magnesium silicate
Sodium starch glycolate
Povidone K30
Magnesium stearate

Hypromellose
Titanium dioxide (E171)
Polyethylene glycol

6.2 Incompatibilities

Not applicable.

6.3 Shelf life

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

6.4 Special precautions for storage

Do not store above 30°C. Keep the blisters in the outer carton in order to protect from light and moisture.

6.5 Nature and contents of container

Each pack contains 35 dispersible tablets. 7 dispersible tablets per blister.

6.6 Special precautions for disposal and other handling

No special requirements.

7 PARALLEL PRODUCT AUTHORISATION HOLDER

Lexon Pharmaceuticals (Ireland) Limited
Block 3
Harcourt Centre
Harcourt Road
Dublin 2
Ireland

8 PARALLEL PRODUCT AUTHORISATION NUMBER

PPA23176/058/001

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 14th April 2023

10 DATE OF REVISION OF THE TEXT