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 6.1

3 PHARMACEUTICAL FORM

Dispersible Tablets

Product imported from the UK and Greece:
White, oval tablet, 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 glycollate
Povidone K30
Magnesium stearate
hypromellose
polyethylene glycol 400
titanium dioxide
polyethylene glycol 8000

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.

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

Blister packs containing 35 dispersible tablets.

6.6 Special precautions for disposal

No special requirements.

7 PARALLEL PRODUCT AUTHORISATION HOLDER

PCO Manufacturing Ltd.
Unit 10, Ashbourne Business Park
Rath
Ashbourne
Co. Meath
Ireland

8 PARALLEL PRODUCT AUTHORISATION NUMBER

PPA0465/038/007

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 31st March 2017

10 DATE OF REVISION OF THE TEXT

October 2021