Summary of Product Characteristics

1 NAME OF THE MEDICINAL PRODUCT

Zovirax 200 mg dispersible tablets

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each tablet contains 200 mg aciclovir.

For the full list of excipients, see 6.1.

3 PHARMACEUTICAL FORM

Dispersible film-coated tablet (Dispersible tablet)

Product imported from Spain
Zovirax 200 mg tablets are white, round film-coated tablets branded with 'GXCF3' on one side and plain on the other.

4 CLINICAL PARTICULARS

As per PA1077/084/007

5 PHARMACOLOGICAL PROPERTIES

As per PA1077/084/007

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Microcrystalline cellulose (E460)
Aluminium magnesium silicate
Sodium starch glycolate
Povidone K30
Magnesium stearate (E572)
Hypromellose
Macrogol 400
Titanium dioxide (E171)
Macrogol 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. Store in the original package.

6.5 Nature and contents of container

PVC/PVDC/Aluminium/Paper child resistant foil blister packs containing 25 dispersible tablets.

6.6 Special precautions for disposal and other handling

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

PPA0465/038/005

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 16th March 2007

Date of last renewal: 16th March 2012

10 DATE OF REVISION OF THE TEXT

March 2021