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

Excipient(s) with known effect:
This product contains sodium as Sodium starch glycollate

For the full list of excipients, see section 6.1

3 PHARMACEUTICAL FORM

Dispersible Film-coated 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

Core:
Microcrystalline cellulose (E460)
Aluminium magnesium silicate
Sodium starch glycollate (type A)
Povidone K30
Magnesium stearate (E572)

Film Coat:
Hypromellose
Titanium dioxide (E171)
Macrogol 400
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. Pack size: 25 dispersible tablets.

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

PPA1463/199/001

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 2nd December 2022

10 DATE OF REVISION OF THE TEXT