Summary of Product Characteristics

1 NAME OF THE MEDICINAL PRODUCT

Zovirax 800 mg Dispersible Tablets

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each tablet contains 800 mg aciclovir

For a full list of excipients, see section 6.1.

3 PHARMACEUTICAL FORM

Dispersible Tablets

Product imported from Greece.

White, biconvex, elongated, film-coated tablet, impressed with “GX CG1” on one face and plain on the other.

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 for 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

Zovirax Dispersible tablets are stored in blister packs. Each pack contains 35 tablets on 7 blister strips.

6.6 Special precautions for disposal of a used medicinal product or waste materials derived from such medicinal product and other handling of the product

No special requirements

7 PARALLEL PRODUCT AUTHORISATION HOLDER

LTT Pharma Limited
Unit 18
Oxleasow Road
East Moons Moat
Redditch
Worcestershire
B98 0RE
United Kingdom

8 PARALLEL PRODUCT AUTHORISATION NUMBER

PPA1562/183/002

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 11th November 2016