

# Summary of Product Characteristics

## 1 NAME OF THE MEDICINAL PRODUCT

Keflex 250mg Hard capsules

## 2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each capsule contains 250 mg cefalexin anhydrous (as the monohydrate).  
For the full list of excipients, see section 6.1.

## 3 PHARMACEUTICAL FORM

Hard capsule

*Product imported from UK*

Green & white, capsule, printed ‘GP1’

## 4 CLINICAL PARTICULARS

As per PA1226/002/001

## 5 PHARMACOLOGICAL PROPERTIES

As per PA1226/002/001

## 6 PHARMACEUTICAL PARTICULARS

### 6.1 List of excipients

Magnesium stearate  
Titanium dioxide  
Cellulose with sodium carboxymethyl cellulose  
Dimeticone  
Patent blue V  
Quinoline yellow  
Gelatin

### 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.  
Store in the original packaging.

## **6.5 Nature and contents of container**

Packs contain 28 capsules in blister packs.

## **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**

PPA1562/181/001

## **9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION**

Date of first authorisation: 2<sup>nd</sup> September 2016

## **10 DATE OF REVISION OF THE TEXT**