

## Summary of Product Characteristics

### 1 NAME OF THE MEDICINAL PRODUCT

Adalat LA 60mg Prolonged-Release Tablet

### 2 QUALITATIVE AND QUANTITATIVE COMPOSITION

One tablet contains 60 mg nifedipine.

Excipients: sodium

For a full list of excipients, see section 6.1.

### 3 PHARMACEUTICAL FORM

Prolonged-release film-coated tablet.

*Product imported from UK:*

Pink, circular convex tablets with Adalat 60 marked on one side.

### 4 CLINICAL PARTICULARS

As per PA1410/025/007

### 5 PHARMACOLOGICAL PROPERTIES

As per PA1410/025/007

### 6 PHARMACEUTICAL PARTICULARS

#### 6.1 List of excipients

##### Tablet Core

Polyethylene oxide

Hypromellose (5 cp)

Magnesium stearate

Sodium chloride

Ferric oxide, red (E172)

Coating

Cellulose acetate  
Macrogol (3350)  
Hydroxypropylcellulose  
Hypromellose (3 cp)  
Propylene glycol  
Hypromellose (5 cp)  
Titanium dioxide (E171)  
Ferric oxide, red (E172)

Polish and Print

Black ink for printing Opacode S-1-81 06  
(Contains: iron oxide black (E172) and Shellac)

**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 container. The tablets should be protected from strong light. Do not remove the tablet from the blister until immediately before use.

**6.5 Nature and contents of container**

Blisters packed into a cardboard carton of 28 tablets

**6.6 Special precautions for disposal and other handling**

No special requirements.

**7 PARALLEL PRODUCT AUTHORISATION HOLDER**

Clear Pharmacy  
157-173 Roden Street  
Belfast  
BT12 5QA  
United Kingdom

**8 PARALLEL PRODUCT AUTHORISATION NUMBER**

PPA1596/035/003

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

Date of first authorisation: 21st January 2011

**10 DATE OF REVISION OF THE TEXT**

February 2015