

## Summary of Product Characteristics

### 1 NAME OF THE MEDICINAL PRODUCT

Lexapro 15mg film-coated tablets

### 2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each tablet contains 15 mg escitalopram (as oxalate)

For a full list of excipients, see section 6.1.

### 3 PHARMACEUTICAL FORM

Film-coated tablet

*Product imported from France:*

Oval, white, scored, film-coated tablet marked with “E” and “M” on each side of the score on one side of the tablet.

### 4 CLINICAL PARTICULARS

As per PA0805/002/003.

### 5 PHARMACOLOGICAL PROPERTIES

As per PA0805/002/003.

### 6 PHARMACEUTICAL PARTICULARS

#### 6.1 List of excipients

Tablet core:

Microcrystalline cellulose  
Silica, colloidal anhydrous  
Talc  
Croscarmellose sodium  
Magnesium stearate

Coating:

Hypromellose  
Macrogol 400  
Titanium dioxide (E 171)

#### 6.2 Incompatibilities

Not applicable.

#### 6.3 Shelf life

The shelf-life expiry date of this product shall be 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**

The medicinal product does not require any special storage conditions.

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

Over-labelled cardboard carton containing 2 blister strips (14 tablets per strip). Pack size of 28 tablets.

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

No special requirements.

### **7 PARALLEL PRODUCT AUTHORISATION HOLDER**

B & S Healthcare  
Unit 4  
Bradfield Road  
Ruislip  
Middlesex  
HA40NU  
UK

### **8 PARALLEL PRODUCT AUTHORISATION NUMBER**

PPA1328/084/003

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

Date of first authorisation: 14<sup>th</sup> January 2011

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

April 2015