

## Summary of Product Characteristics

### 1 NAME OF THE MEDICINAL PRODUCT

IDEOS 500mg/400 IU Chewable Tablets

### 2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Active ingredients per tablet:

Calcium carbonate ..... 1250 mg  
(equivalent to 500 mg of elemental calcium)

Cholecalciferol (vitamin D3) ..... 400 IU  
(equivalent to 10 µg)

Excipients: Also contains sucrose, sorbitol & hydrogenated soya bean oil.

For a full list of excipients, see section 6.1

### 3 PHARMACEUTICAL FORM

Chewable tablets, for oral administration.

*Product imported from GREECE*

Greyish white, square, chewable tablets.

### 4 CLINICAL PARTICULARS

As per PA1033/001/001

### 5 PHARMACOLOGICAL PROPERTIES

As per PA1033/001/001

### 6 PHARMACEUTICAL PARTICULARS

#### 6.1 List of excipients

Xylitol  
Sorbitol (E420)  
Povidone  
Magnesium stearate  
Lemon flavour\*

\*Composition of the lemon flavouring: Flavouring preparations, natural flavouring substances, maltodextrin, acacia, sodium citrate, citric acid, butylated hydroxyanisole.

Excipients in Vitamin D3 concentrate: alpha-tocopherol, partially hydrogenated soyabean oil, gelatin, sucrose, maize starch

#### 6.2 Incompatibilities

Not applicable.

### **6.3 Shelf life**

The shelf life expiry date for this product shall be the date shown on the tube and outer package of the product on the market in the country of origin.

### **6.4 Special precautions for storage**

Store in the original container.  
Do not store above 25°C.

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

Polypropylene tubes containing 15 tablets. Pack size: 60 tablets.

### **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  
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Redditch  
Worcestershire, B98 0RE  
United Kingdom

## **8 PARALLEL PRODUCT AUTHORISATION NUMBER**

PPA1562/078/001

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

Date of first authorisation: 6<sup>th</sup> July 2012

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

February 2015