

# Summary of Product Characteristics

## 1 NAME OF THE MEDICINAL PRODUCT

Zoton FasTab 30mg oro-dispersible tablets

## 2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each oro-dispersible tablet contains 30mg of lansoprazole.

Excipient(s) with known effect:

Each 30 mg oro-dispersible tablet contains lactose and aspartame (E951).

For a full list of excipients, see section 6.1

## 3 PHARMACEUTICAL FORM

Oro-dispersible tablet

*Product imported from UK:*

White to yellowish white, circular, flat bevelled-edge oro-dispersible tablet with '30' debossed on one side.

Each oro-dispersible tablet contains orange to dark brown microgranules.

## 4 CLINICAL PARTICULARS

As per PA0822/101/003

## 5 PHARMACOLOGICAL PROPERTIES

As per PA0822/101/003

## 6 PHARMACEUTICAL PARTICULARS

### 6.1 List of excipients

Lactose monohydrate

Microcrystalline cellulose

Magnesium carbonate

Low substituted hyprolose

Hyprolose

Hypromellose

Titanium dioxide (E171)

Talc

Mannitol

Methacrylic acid-ethyl acrylate copolymer (1:1) dispersion 30 per cent  
Polyacrylate dispersion 30 per cent  
Macrogol 8000  
Glycerol monostearate  
Polysorbate 80  
Triethyl citrate  
Citric acid anhydrous  
Crospovidone  
Magnesium Stearate  
Aspartame (E951)  
Strawberry Flavour  
Iron oxide red (E172)  
Iron oxide yellow (E172)

## **6.2 Incompatibilities**

Not applicable.

## **6.3 Shelf life**

The shelf-life expiry date of this product is the date shown on the blister strips and outer carton of the product on the market in the country of origin.

## **6.4 Special precautions for storage**

Do not store above 25°C.

Store in the original package in order to protect from moisture

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

Aluminium blisters in an over-labelled cardboard carton containing 28 tablets.

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

No special requirements.

## **7 PARALLEL PRODUCT AUTHORISATION HOLDER**

Primecrown 2010 Limited  
4/5 Northolt Trading Estate  
Belvue Road  
Northolt  
Middlesex  
UB5 5QS

United Kingdom

**8 PARALLEL PRODUCT AUTHORISATION NUMBER**

PPA1633/010/002

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

Date of first authorisation: 4<sup>th</sup> June 2010

**10 DATE OF REVISION OF THE TEXT**

January 2019