

Summary of Product Characteristics

1 NAME OF THE MEDICINAL PRODUCT

Tenoret 50mg/12.5mg Film-coated Tablets

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each film-coated tablet contains atenolol 50 mg and chlortalidone 12.5 mg
For a full list of excipients, see section 6.1.

3 PHARMACEUTICAL FORM

Film-coated tablets.
Product imported from UK:

Brown, biconvex, film-coated tablet imprinted with the name TENORET 50 on one face and AstraZeneca logo on the other face.

4 CLINICAL PARTICULARS

As per PA0970/021/001

5 PHARMACOLOGICAL PROPERTIES

As per PA0970/021/001

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Magnesium carbonate
Maize starch
Sodium laurilsulfate
Gelatin
Magnesium stearate
Methyl hydroxypropyl cellulose
Iron oxide
Macrogol

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

Do not store above 25°C. Store in the original package.
Keep the blister in the outer carton.

6.5 Nature and contents of container

Calendar blister strips (box of 28 tablets) in an over-labelled outer carton.

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
HA4 0NU
United Kingdom

8 PARALLEL PRODUCT AUTHORISATION NUMBER

PPA1328/107/001

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 10th July 2009

10 DATE OF REVISION OF THE TEXT

September 2015