

Summary of Product Characteristics

1 NAME OF THE MEDICINAL PRODUCT

Nexium 20 mg gastro-resistant tablets

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each gastro-resistant tablet contains 20 mg esomeprazole (as magnesium trihydrate)

Excipient(s) with known effect


Each gastro-resistant tablet contains sucrose.

For the full list of excipients, see section 6.1.

3 PHARMACEUTICAL FORM

Gastro-resistant tablet

Product imported from the UK and Croatia

Light pink, oblong, biconvex, film-coated tablet engraved 20 mg on one side and  on the other side

4 CLINICAL PARTICULARS

As per PA2242/013/002

5 PHARMACOLOGICAL PROPERTIES

As per PA2242/013/002

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Glycerol Monostearate 40-55
Hyprolose
Hypromellose
Iron oxide (reddish-brown and yellow) (E172)
Magnesium stearate
Methacrylic acid ethyl acrylate copolymer (1:1) dispersion 30 per cent
Microcrystalline cellulose
Synthetic paraffin
Macrogol
Polysorbate 80
Crospovidone
Sodium stearyl fumarate
Sugar spheres (sucrose and maize starch)
Talc
Titanium dioxide (E171)
Triethyl citrate

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

Product imported from the United Kingdom:

Do not store above 30°C. Store in the original package.

Product imported from Croatia:

Do not store above 30°C. Store in the original package in order to protect from moisture.

6.5 Nature and contents of container

Product imported from the United Kingdom:

Blister packs in wallets of 14 tablets, in an overlabelled outer carton, each containing 2 wallets (28 tablets).

Product imported from Croatia:

Blisters of 28 gastro-resistant tablets

6.6 Special precautions for disposal and other handling

No special requirements for disposal.

Administration through gastric tube

1. Put the tablet into an appropriate syringe and fill the syringe with approximately 25 ml water and approximately 5 ml air. For some tubes, dispersion in 50 ml water is needed to prevent the pellets from clogging the tube.
2. Immediately shake the syringe for approximately 2 minutes to disperse the tablet.
3. Hold the syringe with the tip up and check that the tip has not clogged.
4. Attach the syringe to the tube whilst maintaining the above position.
5. Shake the syringe and position it with the tip pointing down. Immediately inject 5-10 ml into the tube. Invert the syringe after injection and shake (the syringe must be held with the tip pointing up to avoid clogging of the tip).
6. Turn the syringe with the tip down and immediately inject another 5 – 10 ml into the tube. Repeat this procedure until the syringe is empty.
7. Fill the syringe with 25 ml of water and 5 ml of air and repeat step 5 if necessary to wash down any sediment left in the syringe. For some tubes, 50 ml water is needed.

7 PARALLEL PRODUCT AUTHORISATION HOLDER

Merit Pharmaceuticals Limited
Unit C4/C3 Metropoint Point Business Park
Kettles Lane
Swords
Co Dublin
K67 RH92
Ireland

8 PARALLEL PRODUCT AUTHORISATION NUMBER

PPA23080/003/001

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 28th May 2010

10 DATE OF REVISION OF THE TEXT

September 2021