

Summary of Product Characteristics

1 NAME OF THE MEDICINAL PRODUCT

Risedronate Sodium Accord Once a Week 35 mg film-coated tablets

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each film-coated tablet contains 35 mg risedronate sodium, (equivalent to 32.5 mg risedronic acid).

Excipient with known effect:

Lactose monohydrate

For the full list of excipients, see section 6.1

3 PHARMACEUTICAL FORM

Film-coated tablet.

Product imported from Italy, Spain, Greece, the UK, France and Romania:

Oval, light-orange tablet with 'RSN' engraved one side and '35 mg' on the other.

4 CLINICAL PARTICULARS

As per PA2315/140/003

5 PHARMACOLOGICAL PROPERTIES

As per PA2315/140/003

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Product imported from Italy, Spain, Greece, the UK, France:

Tablet core:

Lactose monohydrate

Microcrystalline cellulose

Crospovidone

Magnesium stearate

Film coating:

Dri-Klear (hypromellose, macrogol 400, hypromellose macrogol 400, macrogol 8000 and colloidal hydrated silica)

Chroma-Tone White DDB-7536W (titanium dioxide (E171), hypromellose)

Ferric oxide yellow (E172)

Product imported from Romania:

Tablet core:

Lactose monohydrate

Microcrystalline cellulose

Crospovidone

Magnesium stearate

Film coating:

Macrogol

Hypromellose

Hydroxypropyl cellulose

Polyethylene glycol

Colloidal anhydrous silica
Titanium dioxide (E171)
Ferric oxide yellow (E172)
Ferric oxide red (E172)

6.2 Incompatibilities

Not applicable.

6.3 Shelf life

The shelf life expiry date of this product is the date shown on the container and outer package of the product in the country of origin.

6.4 Special precautions for storage

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

6.5 Nature and contents of container

Blister packs of 4 tablets contained in an outer cardboard carton.

6.6 Special precautions for disposal

No special requirements.

Any urgent medicinal product or waste should be disposed of in accordance with local requirements.

7 PARALLEL PRODUCT AUTHORISATION HOLDER

PCO Manufacturing Ltd.
Unit 10, Ashbourne Business Park
Rath
Ashbourne
Co. Meath
Ireland

8 PARALLEL PRODUCT AUTHORISATION NUMBER

PPA0465/165/001

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 04 November 2005
Date of last renewal: 04 November 2010

10 DATE OF REVISION OF THE TEXT

April 2021