Summary of Product Characteristics

1 NAME OF THE MEDICINAL PRODUCT

Dona 1500 mg powder for oral solution

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each sachet contains:
Glucosamine Sulfate sodium chloride 1884 mg
equivalent to: glucosamine sulphate 1500 mg
sodium chloride 384 mg

Excipients with known effect: Each sachet contains aspartame (E951), sodium and sorbitol (E420).

For the full list of excipients, see section 6.1

3 PHARMACEUTICAL FORM

Powder for oral solution.

Product imported from Spain:
A white, crystalline, odourless powder.

4 CLINICAL PARTICULARS

As per PA2010/022/002

5 PHARMACOLOGICAL PROPERTIES

As per PA2010/022/002

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Aspartame (E951)
Sorbitol (E420)
Citric Acid
Macrogol 4000

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.

6.5 Nature and contents of container

Single-dose sachets in a cardboard box containing 30 sachets.

6.6 Special precautions for disposal and other handling

No special requirements

7 PARALLEL PRODUCT AUTHORISATION HOLDER

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

8 PARALLEL PRODUCT AUTHORISATION NUMBER

PPA0465/401/001

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 12th September 2014

10 DATE OF REVISION OF THE TEXT

May 2021