

# 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 sulfate 1500 mg  
sodium chloride 384 mg

Excipients with known effect:

One sachet contains aspartame, sodium and sorbitol

For the full list of excipients, see section 6.1

## 3 PHARMACEUTICAL FORM

Powder for oral solution.

*Product imported from Portugal*

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 (E330)  
Macrogol 4000

### 6.2 Incompatibilities

Not applicable.

### 6.3 Shelf life

The shelf life expiry date for this product shall be the date shown on the sachets 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**

The single-dose sachet consists of three layered material made of paper, aluminium and polyethylene.  
Cardboard box containing 30 sachets.

### **6.6 Special precautions for disposal**

No special requirements

### **7 PARALLEL PRODUCT AUTHORISATION HOLDER**

Lexon Pharmaceuticals (Ireland) Limited  
Block 3  
Harcourt Centre  
Harcourt Road  
Dublin 2  
Ireland

### **8 PARALLEL PRODUCT AUTHORISATION NUMBER**

PPA23176/035/001

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

Date of first authorisation: 18<sup>th</sup> September 2020

### **10 DATE OF REVISION OF THE TEXT**

February 2022