

# Summary of Product Characteristics

## 1 NAME OF THE MEDICINAL PRODUCT

Metoject 10 mg solution for injection in pre-filled pen

## 2 QUALITATIVE AND QUANTITATIVE COMPOSITION

1 pre-filled pen with 0.20 ml solution contains methotrexate disodium equivalent to 10 mg methotrexate. For the full list of excipients, see section 6.1.

## 3 PHARMACEUTICAL FORM

Solution for injection

Product imported from *Germany*.  
Clear, yellow-brown solution.

## 4 CLINICAL PARTICULARS

As per PA0623/014/003

## 5 PHARMACOLOGICAL PROPERTIES

As per PA0623/014/003

## 6 PHARMACEUTICAL PARTICULARS

### 6.1 List of excipients

Sodium chloride  
Sodium hydroxide  
Water for injections

### 6.2 Incompatibilities

In the absence of compatibility studies, this medicinal product must not be mixed with other medicinal products.

### 6.3 Shelf life

The shelf life expiry date of this product shall be the date shown on the pre-filled pen and outer package of the product on the market in the country of origin.

### 6.4 Special precautions for storage

Store below 25 °C. Keep the pre-filled pens in the outer carton in order to protect from light.

### 6.5 Nature and contents of container

Pre-filled pen containing 0.20 ml (10 mg) solution. Pack size 1 pre-filled pen.

### 6.6 Special precautions for disposal and other handling

The manner of handling and disposal must be in accordance with local requirements. Pregnant health care personnel should not handle and/or administer Metoject.

Methotrexate should not come into contact with the skin or mucosa. In the event of contamination, the affected area must be rinsed immediately with ample amount of water.

For single use only.

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

**7 PARALLEL PRODUCT AUTHORISATION HOLDER**

IMED Healthcare Ltd.  
Unit 625 Kilshane Avenue  
Northwest Business Park  
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Dublin 15  
Ireland

**8 PARALLEL PRODUCT AUTHORISATION NUMBER**

PPA1463/172/002

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

Date of first authorisation: 10<sup>th</sup> September 2021

**10 DATE OF REVISION OF THE TEXT**