

IMPORTANT INFORMATION

ILARIS[®] (canakinumab)

150 mg subcutaneous injection

For the treatment of Gouty Arthritis attacks

EURMP-V11.2 July 2023

You can report any problem or adverse events or request additional copies of the materials through:

Patient Safety Department Novartis Pharma AG - Saudi Arabia -

Toll Free Number: 8001240078

Phone: +966112658100

Fax: +966112658107

Email: adverse.events@novartis.com

Or by online: <http://report.novartis.com/>

Saudi Food and Drug Authority National Pharmacovigilance Center

Unified Contact Center: 19999

Fax: +966112057662

Email: npc.drug@sfd.gov.sa

Or by online: <https://ade.sfda.gov.sa>

This document is approved by SFDA

Before starting canakinumab

- **Infections:** You should not be treated with canakinumab if you have an active infection.
- **Vaccinations:** Talk to your doctor about any vaccinations you may need before starting treatment with canakinumab.

During canakinumab treatment

- **Risk of infections:** Use of canakinumab is associated with an increased risk of infections, including serious infections.
Tell your doctor **immediately** if you have a fever lasting longer than 3 days or other symptoms that might be due to an infection.
- Seek medical attention immediately if you develop symptoms such as:
 - prolonged fever, cough or headache, or
 - localised redness, warmth or swelling of your skin, or
 - persistent cough, weight loss and low-grade fever
- **Pregnancy:** If you received canakinumab while you were pregnant, it is important that you inform the baby's doctor or nurse before any vaccinations are given to your baby. Your baby should not receive live vaccines until at least 16 weeks after you received your last dose of canakinumab before giving birth.

Treatment Indication:

Please make sure to have a list of all medications you are taking when visiting a healthcare professional.

Patient's name:

Date of first dose of canakinumab:.....

Canakinumab dose administered:.....

Doctor's name:.....

Doctor's phone:.....