

## The bearer of this card is treated with Tofacitinib Taro

Patient's name:

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Doctor's name:

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Doctor's phone number:

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Date of starting treatment with Tofacitinib Taro:

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This card contains important safety information about Tofacitinib Taro.

If this information is not clear to you, please ask your doctor or pharmacist to explain it to you. Keep this card and present it to any doctor or pharmacist involved in your care.

If you stop taking Tofacitinib Taro, keep this card for at least 2 months after taking the last dose of Tofacitinib Taro.

Read the patient information leaflet for more information. You should use Tofacitinib Taro according to the patient information leaflet.

**Tell your doctor or your pharmacist about all the medicines you take**, including prescription and non-prescription medicines, vitamins and herbal nutritional supplements. Taking Tofacitinib Taro with certain medicines may increase your risk of side effects, immunosuppression and infections.

The treatment with Tofacitinib Taro may increase the risk of infections and malignancies (including lung cancer, lymphoma, and non-melanoma skin cancer).

Patients aged 65 years and older may be at high risk of infections, heart attack and certain types of cancer. Your doctor may decide that Tofacitinib Taro is not suitable for you.

Contact your doctor **immediately** if you:

- Develop sudden shortness of breath or breathing difficulties, chest or upper back pain, swelling of the leg or arm, leg pain or tenderness, or leg or arm redness or color change while taking Tofacitinib Taro, as these may be signs of a clot in the lungs or veins.
- Develop symptoms of infection, such as fever, persistent cough, weight loss, or excessive tiredness.
- Develop symptoms of herpes zoster, such as blisters or painful rash on the skin.
- Have been in close contact with a person with tuberculosis.
- Develop intense pain or pressure in your chest (which could spread to your arms, jaw, neck and back), shortness of breath, cold sweat, a sensation that you are about to faint or sudden dizziness, as these may be signs of a heart attack.
- Develop swelling of lymph nodes in your neck, armpits or groin, constantly feeling tired, develop fever, night sweats, experience persistent or worsening cough, develop breathing difficulties, hoarseness or wheezing, or unexplained weight loss.
- Notice any new lesion on the skin or any changes in existing moles or spots.

- Develop symptoms of interstitial lung disease, such as shortness of breath.
- Develop abdominal signs and symptoms such as stomach pain, abdominal pain, blood in your stool, or any change in your bowel habits accompanied by fever.
- Develop yellow skin, nausea or vomiting.
- Are planning to receive any vaccine. There are certain types of vaccines that you must not receive during treatment with Tofacitinib Taro.
- Become pregnant or plan to become pregnant.

Tofacitinib Taro must not be used during pregnancy. Women of childbearing age should use effective contraception methods during treatment with Tofacitinib Taro and for at least 4 weeks after the last dose.

Do not breastfeed during treatment with Tofacitinib Taro.

### **Reporting side effects**

Side effects can be reported to the Ministry of Health by the portal for reporting side effects found on the homepage of the Ministry of Health: [www.health.gov.il](http://www.health.gov.il), or by entering the link: <https://sideeffects.health.gov.il>

Side effects can also be reported to Taro by email: [drug.safety@sunpharma.com](mailto:drug.safety@sunpharma.com)

This card was approved according to the guidelines of the Ministry of Health in April 2025.

**Safety Information Card  
for a patient treated with Tofacitinib Taro  
(Tofacitinib 5 mg, 10 mg)**