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TREATMENT WITH KINERET®

Patient's name:

Caregiver's name (for children):

Patient's dose:

Physician's name and phone number:

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Call for Safety Reporting

Suspected Adverse Drug Reactions (side effects) or medication errors may be reported using the Medicines Authority ADR reporting form, which is available online at: http://www.medicinesauthority.gov.mt/adrportal and sent by post or email to:
P: Pharmacovigilance Section at Post-Licensing Directorate, Medicines Authority, Sir Temi Zammit Buildings, Malta Life Sciences Park, San Gwann SGN 3000
E: postlicensing.medicinesauthority@gov.mt

Alternatively, the contact details of the company may be used for reporting:
Sobi IKE; E-mail: safety@pharmassist.gr; Tel: 00302106561435

www.sobi.com
HCC-2183 - September 2018

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Important safety information regarding Still's disease

Macrophage activation syndrome (MAS) is a serious complication of Still's disease. Left untreated MAS can be life threatening. The risk for developing MAS is increased if you have an infection or if your Still's disease symptoms are poorly controlled. Symptoms of MAS can be e.g. persistent high fever, swelling of lymph nodes, and persistent rash.

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Kineret® (anakinra) can increase the risk for a serious infection. Symptoms might be persistent high fever, shivers, cough, headache, and redness and tenderness of the skin. Also, persistent low-grade fever, weight loss, and persistent cough can be signs of an infection.

If you develop signs of an infection or worsening of your Still's disease symptoms you should contact your health care provider as soon as possible.