



ENBREL PATIENT CARD

This card contains important safety information that you need to be aware of before you are given Enbrel and during treatment with Enbrel. If you do not understand this information, please ask your doctor to explain it to you.

- Show this card to any doctor involved in your treatment.
- See the Enbrel package leaflet for more information.
- Keep this card with you for 2 months after the last Enbrel dose, since side effects may occur after your last dose of Enbrel.

It is important that you and your doctor record the brand name and batch number of your medication

Enbrel batch number:

Infections			
Enbrel may increase you	ar risk of getting infections, w	which could be serious.	
If you develop listlessness, seeYou should be	suse Enbrel if you have an in symptoms suggestive of infects k medical attention immedia evaluated for tuberculosis (T eening for TB below:	ctions, such as fever, persist tely.	ent cough, weight loss, or
Test:_		Test:	
Date:		Date:	
Result	s:	Results:	
Please ask your	physician to list your other i	nedications that may increa	se your risk of infection.

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Other Information (please complete)

Patient's Name:		
Doctor's Name:		
Doctor's Phone:		

Call for 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 https://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, you may also report such events promptly to Pfizer Hellas Pharmacovigilance Department, contact details: +30 210 67 85 908 and +30 210 67 85 808 (24-hour line), or via the webportal <u>Pfizer Adverse Event Safety Reporting</u> (Pfizersafetyreporting.com), or contact the local representative Vivian Corporation Ltd. Tel. + 356 21344610/ +356 2258 8600.)