

ConSERF

CONSUMER SIDE EFFECT REPORTING FORM NATIONAL CENTRE FOR ADVERSE DRUG REACTIONS MONITORING Help us make medicines safer



Please fill in all sections marked with * and give as much other information as you can.

All personal data will remain **confidential**.

Report No. (for official use):

All personal data will remain confi	dential.			
Information about the person	1 who had the side effect		Rep	orter details
*Age :	Nationality:	se	Repor	of report: rter's name: Number : address:
*Any health problems / allergies / pregnancy? (please specify): E.g.: Diabetes, high blood pressure, asthma, allergy to painkiller, or 16 weeks pregnant				
e.g Diabetes, ingit blook pressore, astrima, altergy to parmitter, or 10 weeks pregnant				
Information about the medication(s) suspected to cause the side effect, and other medications				
*Suspected Medicine(s): (plea			(please a	attach additional sheets if necessary)
Suspected medicine name	Dosage	Dates:		Reason for use
(include MAL number if known)	(e.g. 250mg three times daily)	Started DD/MM/YY	Stopped DD/MM/YY	
		DD/IVIIVI/ I I	DD/WWW/TT	
*Were any other medicines tak	an at the same time? \ \\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\	logso givo th	no dotails hold	ow) □No
*Were any other medicines taken at the same time?: Other medicine(s) name Dosage Dates:				Reason for use
(include MAL number if known)	(e.g. 250mg three times daily)	Started	Stopped	Reason for ose
-		DD/MM/YY	DD/MM/YY	-
Information on the side effe	rt(s)			
		v v h	\ Donation su	sheidad an D. D. W. W. W. W.
1. * Date of side effect(s): a) Reaction started on D D M M Y Y b) Reaction subsided on D D M M Y Y				
2. * Please describe the side effect(s) experienced:				
3. * How long was the medication(s) taken before the side effect appeared? minutes/hours/days/months/years (choose)				
4. * Did the side effect subside when the medication(s) was stopped ?				
5. * Did the side effect reappear when the medication(s) was <u>taken again</u> ?□ Yes □ No □ Did not take again				
6. * How serious was the side effect? (select all that apply below)				
☐ Mild or slightly uncomfortable ☐ Had to seek medical advice ☐ Admitted to the hospital				
☐ Uncomfortable but could carry out daily activities ☐ Bad, interferes with daily activities ☐ Other:				
7. * Was any treatment given / medication taken to overcome the side effect? — Yes (please specify) — No				
8. * What is the current outcome of	of the side effect?			
□ Fully recovered		de effects co	ontinuing	□ Caused death
Thank you for reporting				

ConSERF

CONSUMER SIDE EFFECT REPORTING FORM

Help us make medicines safer

If you think you have a side effect to your medicine, please seek advice from your pharmacist or doctor.

What is ConSERF?

- This form is used to report a suspected side effect to any medicine or vaccine (including prescription, over-the-counter, or traditional products, health supplements, cosmetic products, etc.).
- A side effect (or adverse drug reaction ADR) is defined as any unintended effect of a medicine which occurs at the normal dose used.
- Please report any side effect you find troubling, even if you are not certain it is due to the medicine or vaccine.
- Your identity and the information provided will be kept confidential.

Why report a side effect?

- This will help improve the safe use of medicines
- This may identify new side effects of a medicine

Every report will be analysed and entered into the Malaysian and World Health Organisation (WHO) databases of medication side effects.

How to report?

- Obtain this form from your local pharmacist or from our website (http://npra.moh.gov.my --> Orang Awam). Please complete as many sections as possible to ensure your report is useful. Consult your pharmacist for assistance.
- Please return the form to your pharmacist to be sent to us, submit online, or post/email directly to us.
- Please provide your contact details to allow us to obtain further information about your report if necessary.

Questions or comments?

Contact us::

National Pharmaceutical Regulatory Agency

Ministry of Health Malaysia

| http://npra.moh.gov.my | fv@npra.gov.my | Tel: 03-7801 8464/ 8470 | Fax: 03-7956 7151 |

ConSERF

Consumer Side Effect Reporting Form

Bahagian Regulatori Farmasi Negara (NPRA) Kementerian Kesihatan Malaysia

PUSAT PEMONITORAN KESAN ADVERS UBAT KEBANGSAAN
BAHAGIAN REGULATORI FARMASI NEGARA (NPRA)

PETI SURAT 319, JALAN SULTAN

46730 PETALING JAYA

SELANGOR