

Adverse Drug Reaction (ADR) Reporting Form

A. Patient Details

Patient name (Optional):	Age:	Gender: <input type="checkbox"/> Male <input type="checkbox"/> Female	Weight:	Height:
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B. Reporter Details

Reporter name:	Relationship to the patient:
Address:	E-mail:
Phone / Mobile:	

C. Suspected Product Information

Type of product:	<input type="checkbox"/> Medicinal product	<input type="checkbox"/> Herbal product	<input type="checkbox"/> Vaccine	<input type="checkbox"/> Food Supplement	<input type="checkbox"/> Other	
Trade Name	Active ingredient name:					
Batch Number:	Manufactured Name:					
How did you obtain the product:	<input type="checkbox"/> From the pharmacy with prescription		<input type="checkbox"/> From the pharmacy without prescription			
	<input type="checkbox"/> From other places (non-pharmacy), specify:					
Name and address of the establishment from which the product was dispensed:						
Dosage form of the product:	<input type="checkbox"/> Tablets	<input type="checkbox"/> Capsules	<input type="checkbox"/> Syrup	<input type="checkbox"/> Suppositories	<input type="checkbox"/> Eye/Ear drops	<input type="checkbox"/> Injection
	<input type="checkbox"/> Topical ointment/cream	<input type="checkbox"/> Herbs/Herbal mixture	<input type="checkbox"/> Eye/Ear ointment	<input type="checkbox"/> Spray	<input type="checkbox"/> Patches	
	<input type="checkbox"/> Other, specify:					
Date of starting use of the product:	Route of administration:					
Purpose of use:	Dosage:					
Did you stop using the product:	<input type="checkbox"/> Yes	<input type="checkbox"/> No				

D. Adverse Drug Reaction (ADR) Description

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Date of event started:	Date of event disappeared, if applicable:

E. Seriousness of ADR Serious (If the case is serious, specify one of the conditions listed below) Non-serious Un-known

<input type="checkbox"/> Patient died, date:	<input type="checkbox"/> Life threatening	<input type="checkbox"/> Permanent disability
<input type="checkbox"/> Hospitalization	<input type="checkbox"/> Prolonged hospitalization more than 24 hr	<input type="checkbox"/> Congenital anomaly
<input type="checkbox"/> Required intervention to prevent permanent impairment/ damage	<input type="checkbox"/> Required Emergency Room (ER) visit	
<input type="checkbox"/> Other.....		

Date:
Case Serial No:



F. Action Taken

Drug withdrawal Dose reduced Dose increased Dose not changed Unknown Not applicable

G. Outcome of ADR (Tick all applicable)

The patient Recovered, date: Recovering No improvement Unknown

G. Additional Information

Was the doctor or pharmacist informed about these symptoms? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Un-known		
If yes, did they complete the adverse reaction reporting form? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Un-know		
May we obtain additional information from your treating physician? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Un-know		
Doctor's Name:	Hospital:	Phone Number:

- **Privacy:**

The information provided in the report is treated with strict confidentiality and is fully protected, including the identity of the patient and the person submitting the report. This information cannot be used against the reporter under any circumstances.