

PHILLIPS PHARMA GROUP – REPORT AN ADVERSE REACTION*

* any untoward medical occurrence that may present during treatment with a medicine, but which does not necessarily have a causal relationship with this treatment – coincidence in time without any suspicion of causal relationship, or a response to a medicine which is noxious (harmful) and unintended, and which occurs at doses normally used in humans, including:

- Abnormal tests or laboratory findings
- Signs and symptoms of clinical significance
- Changes in physical examination findings
- Hypersensitivity
- Progression of underlying disease
- Drug interaction
- Drug dependency

Please fill out the form below and send to the email address indicated in the footer:

Patient Initials		First Initial:	Last Initial:
Country			
Date of Birth		DD-MMM-YYYY	
Gender	<input type="checkbox"/> Male <input type="checkbox"/> Female		
Date of reaction onset		DD-MMM-YYYY	
Describe Reaction			
Please check all results applicable to adverse reaction		<input type="checkbox"/> Patient died <input type="checkbox"/> Involved or prolonged inpatient hospitalization <input type="checkbox"/> Involved persistence or significant disability or incapability <input type="checkbox"/> Life threatening	
Suspect drug(s) including generic name			
Batch number of suspect drug			
Source of medicine			
Daily dose(s)			
Route(s) of administration		<input type="checkbox"/> Oral <input type="checkbox"/> Intravenous (IV) <input type="checkbox"/> Intramuscular (IM) <input type="checkbox"/> Intrathecal <input type="checkbox"/> Subcutaneous <input type="checkbox"/> Buccal <input type="checkbox"/> Rectal <input type="checkbox"/> Vaginal <input type="checkbox"/> Ocular	
		<input type="checkbox"/> Otic <input type="checkbox"/> Transnasal <input type="checkbox"/> Inhalation <input type="checkbox"/> Nebulization <input type="checkbox"/> Topical <input type="checkbox"/> Transdermal <input type="checkbox"/> Intraosseous <input type="checkbox"/> Other:	
Indication(s) for use			
Therapy dates		FROM:	TO:
Therapy duration			
Did reaction abate after stopping drug?		<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Not Applicable	

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Did reaction reappear after reintroduction?	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Not Applicable	
Concomitant drug(s) and dates of administration (exclude those used to treat reaction)		
Other relevant history (eg: diagnostics, allergies, pregnancy with last month of period, etc.)		
Name and address of manufacturer		
Report source	<input type="checkbox"/> Patient <input type="checkbox"/> Study <input type="checkbox"/> Literature <input type="checkbox"/> Health Professional <input type="checkbox"/> Other	
Date of report		
Report type	<input type="checkbox"/> Initial <input type="checkbox"/> Follow Up	
Your email		We may need to get in touch with you for follow up or details
Your phone number		
Upload any relevant attachments		Prescription, results, images, etc.
Consent	<input type="checkbox"/> We may share your data with third parties but keep your information confidential.	

Please note that any data provided herein, in any form, is collected and will be processed by us in accordance with the prevailing laws and regulations in the land. By submitting your information, you expressly consent to the collection and processing of your data.

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