



LADY READING HOSPITAL (MTI) PESHAWAR
DEPARTMENT OF PHARMACY SERVICES

Form No. _____

Date: _____

ADVERSE DRUG REACTION REPORTING FORM
For Voluntary reporting of Adverse Drug Reactions by healthcare professionals

IMPORTANT: DO NOT PLACE IT IN PATIENT'S RECORD

Advice about Reporting:

- Report adverse experiences with medications
- Report serious adverse reactions. A reaction is serious when the patient outcome is:
 - death
 - life-threatening (real risk of dying)
 - hospitalization (initial or prolonged)
 - required intervention to prevent permanent impairment or damage
 - disability (significant, persistent or permanent)
 - congenital anomaly
- Report even if you're not certain that the product caused adverse drug reaction

Who can report?

- Any healthcare professional (Doctors including Dentists, Nurses and Pharmacists)

Where to report?

- Please return the completed form to Pharmacovigilance Centre (PVC), Department of Pharmacy, Lady Reading Hospital (MTI) Peshawar

What happens to the submitted information?

- Information provided in this form is handled in strict confidence. The causality assessment is carried out at the Pharmacovigilance centre (PVC) LRH-MTI Peshawar by using specific scale (recommended by WHO). The analysed forms are forwarded to Drug regulatory authority of Pakistan (DRAP) and further reported to WHO ADR monitoring centre.
- The reports are periodically reviewed by the Pharmacovigilance centre (PVC) LRH-MTI Peshawar. The information generated on the basis of these reports helps in continuous assessment of the benefit-risk ratio of medicines.
- The information is submitted to the Pharmacy & Therapeutic Committee (P&TC) LRH-MTI Peshawar. The Committee is entrusted with the responsibility to review the data and suggest any interventions that may be required.

Date:	Reporting Person & Designation:		
Patient Name:	MR No.	Ward:	
Gender:	Age (Years):	Weight (Kg):	
Date of ADR:	Suspected Drug(s):		
Date of Recovery:			

DRUG TYPE	<input type="checkbox"/> Analgesic	<input type="checkbox"/> Bronchodilator	<input type="checkbox"/> IV Solution	<input type="checkbox"/> Other (specify)
	<input type="checkbox"/> Antimicrobial	<input type="checkbox"/> Cardiovascular	<input type="checkbox"/> Laxative	
	<input type="checkbox"/> Anticoagulant	<input type="checkbox"/> Contrast Media	<input type="checkbox"/> Narcotic	
	<input type="checkbox"/> Anticonvulsant	<input type="checkbox"/> Corticosteroid	<input type="checkbox"/> Oxytocics	
	<input type="checkbox"/> Antidepressant	<input type="checkbox"/> Diuretic	<input type="checkbox"/> Radionuclide	
	<input type="checkbox"/> Antiemetic	<input type="checkbox"/> Immunization	<input type="checkbox"/> Sedative	
	<input type="checkbox"/> Antihistamine	<input type="checkbox"/> Insulin	<input type="checkbox"/> Vasodilator	
	<input type="checkbox"/> Antineoplastic	<input type="checkbox"/> Investigational	<input type="checkbox"/> Vasopressor	

Description of Reaction:	Seriousness of the Reaction:
Reaction Management:	
Reaction severity: <input type="checkbox"/> Fatal <input type="checkbox"/> Recovering <input type="checkbox"/> Unknown <input type="checkbox"/> Continuing <input type="checkbox"/> Recovered <input type="checkbox"/> Other (specify)	
	<input type="checkbox"/> Death <input type="checkbox"/> Life Threatening <input type="checkbox"/> Prolonged Hospitalization <input type="checkbox"/> Disability <input type="checkbox"/> Congenital abnormality <input type="checkbox"/> Required intervention to prevent permanent impairment/damage <input type="checkbox"/> Other (specify)
	Type of ADR: <input type="checkbox"/> Augmented (A) <input type="checkbox"/> Bizarre (B) <input type="checkbox"/> False Alarm (F) <input type="checkbox"/> Hypersensitivity

Confidentiality: The patient's identity is held in strict confidence and protected to the fullest extent. Programme staff is not expected to and will not disclose the reporter's identity in response to a request from the public. Submission of a report does not constitute an admission that medical personnel or manufacturer or the product caused or contributed to the reaction.

Reverse side to be completed by PVC-Pharmacist and to be returned to DPIC