ADR Reporting Form

health
Department:
Health
REPUBLIC OF SOUTH AFRICA

NDoH Pharmacovigilance Centre for Public Health Programmes (NPC)
Adverse Drug Reaction (ADR) / Product Quality Problem Report Form
This report will be shared with the
South African Health Products Authority (SAHPRA)
adr@sahpra.org.za or call 012501031

Reporting Health Care Facility/Practice				
Tel: 012 395 9506 (NPC) **Fax:** 086 241 2473 **Email:** npc@health.gov.za	Facility/Practice			
	District		Tel	
	Province		Fax	

Patient Details									
Patient Initials		File/Reference Number			Date of Birth/Age				
Sex	□M □F □Unk	Race		Weight (kg)		Height (cm)		Pregnant?	□N □Y
Allergies				Estimated Gestational Age at time of reaction					

Suspect Medicine(s) **[Medicines suspected to have caused the ADR]**

Trade Name [Generic Name if Trade Name is unknown]	Name of Manufacturer	Route	Dose (mg) and Interval	Date Started	Date Stopped	Reason for use	Batch Number / Expiry Date

All other Medicines Patient was taking at time of reaction **[Including over-the-counter and herbal products]**

Trade Name [Generic Name if Trade Name is unknown]	Name of Manufacturer	Route	Dose (mg) and Interval	Date Started	Date Stopped	Reason for use	Batch Number / Expiry Date

Adverse Drug Reaction/Product Quality Problem			
Date and time of onset of reaction		Date reaction resolved/duration	

Please describe Adverse Reaction/Product Quality Problem: (kindly add as much clinical information as possible)

Intervention **[tick all that apply]**

□ No intervention
□ Intervention unknown
□ Patient counselled/non-medical treatment
□ Discontinued Suspect Drug; **Replaced with**:_____
□ Decreased Suspect Drug Dosage; **New Dose**:_____
□ Treated ADR with:_____
□ Referred to hospital; **Hospital Name**_____
□ Other Intervention (e.g. dialysis): _____

Patient Outcomes **[tick all that apply]**

□ Patient recovered □ Patient recovering
□ Patient not recovering □ Outcome unknown
□ Patient died; Date of death:_____
□ Impairment/Disability □ Congenital Anomaly
□ Patient hospitalised or hospitalisation prolonged
□ Life threatening □ Other:_____
□ ADR reappeared after restarting suspect drug/similar drug (rechallenge)?: □ N □ Y □ Not done □ Unknown

Laboratory Results

Lab Test	**Test Result**	**Test Date**	**Lab Test**	**Test Result**	**Test Date**

Co-morbidities/Other Medical Condition(s) **[tick all that apply]**

□ Hypertension □ Diabetes □ Asthma □ Tuberculosis □ HIV/AIDS □ Other:

Reported by

Name			E-mail		
Designation	□ Nurse □ Pharmacist □ Doctor □ Other:			Date Reported	
Telephone		Signature		**VERSION 35.0 May 2021**	

THIS ADR REPORT IS NOT A CONFIRMATION THAT THE REPORTER OR THE SUSPECT MEDICINE(S) CAUSED THE ADR