



sa military health service

Department:
Defence
REPUBLIC OF SOUTH AFRICA

SOUTH AFRICAN MILITARY HEALTH SERVICE (SAMHS): PATIENT CLINICAL MOTIVATION FORM FOR MEDICINE NOT INCLUDED IN THE SAMHS MEDICINE FORMULARY

- After completion of this clinical motivation form, please submit it to the SAMHS Medicine Formulary (SAMHS MF) Committee. Reference to supporting evidence, e.g. trials, studies, research papers and/or professional body guidelines must be attached. It is important that the motivation indicates why pharmaceuticals available are not acceptable.
- Relevant test results, a **DD3638 or private prescription** and the SAHPRA Adverse Drug Reaction/Product Quality Problem Report Form, if applicable, must be attached. For non SAMHS prescribers, the referring SAMHS HCP must be indicated. As per the National Treasury Regulations, pharmaceutical trade names cannot be requested.

1

PATIENT'S DETAILS (Please complete in full or place patient's sticker)

Force Number									
Initials & Surname									
Gender	M	F	Date of Birth						
Contact Details: Telephone/email									

2

PRESCRIBER'S DETAILS (Please complete in full)

Prescriber's name									
Specialist Prescriber?	Yes	No	Specialist Discipline						
Signature	C	C	Y	Y	M	M	D	D	
	Tel No								

3

MEDICAL DETAILS (Prescribing Specialist/Consultant to complete in full)

Medicine (generic name) to be authorised									
Duration of treatment									
Patient co-morbidities									



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Previous medicine used

	PRODUCT NAME	QUANTITY USED PER MONTH	START DATE	DURATION OF TREATMENT	TREATMENT OUTCOME
1					
2					
3					

Motivation from the prescribing doctor (Please indicate if the medicine motivated for is aligned with accepted and published professional guidelines, eg Hypertension Society of South Africa, National Osteoporosis Foundation.) Add pages and list relevant tests if necessary.

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4 FOR SAMHS USE ONLY (To be completed by SAMHS MF Committee)

Comments and recommendations

Is the motivated for medicine on proprietary contract?	<input type="checkbox"/> Y	<input type="checkbox"/> N	or quotation?	<input type="checkbox"/> Y	<input type="checkbox"/> N
Is the motivation aligned with the SAMHS Medicine Formulary (SAMHS MF)?	<input type="checkbox"/> Y	<input type="checkbox"/> N			
Is the motivation aligned with best practise (Professional body guidelines)?	<input type="checkbox"/> Y	<input type="checkbox"/> N			
Cost per month=	R				
Was the Adverse Drug Reaction/Product Quality Problem Report Form included?	<input type="checkbox"/> Y	<input type="checkbox"/> N			
Refer back to prescriber?	<input type="checkbox"/> Y	<input type="checkbox"/> N			
to Director Medicine/HOD Clinical Speciality/Pharmacologist?			<input type="checkbox"/> Y	<input type="checkbox"/> N	
Approved?	<input type="checkbox"/> Y	<input type="checkbox"/> N			
Did you communicate the approximate supply time to: The prescriber?	<input type="checkbox"/> Y	<input type="checkbox"/> N	Patient?	<input type="checkbox"/> Y	<input type="checkbox"/> N

Force Number	Rank	Name	Tel No					
Signature	C	C	Y	Y	M	M	D	D



ADVERSE DRUG REACTION (ADR)/PRODUCT QUALITY PROBLEM REPORT FORM
(PUBLIC AND PRIVATE SECTOR) (Including Herbal Products)

Reporting Health Care Facility/Practice							
Tel: 012 842 7609/10 (SAHPRA) 021 447 1618 (NADEMC) Fax: 021 448 6181 E-mail: adr@sahpra.org.za	Facility/Practice						
	District				Tel		
	Province				Fax		
Patient Details							
Force/ID Number	Initials & Surname			Date of Birth/Age			
Sex	<input type="checkbox"/> M <input type="checkbox"/> F <input type="checkbox"/> Unk	Race	Weight (kg)	Height (cm)	Pregnant?	<input type="checkbox"/> N <input type="checkbox"/> 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]	Route	Dose (mg) and Interval	Date Started/Given	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]	Route	Dose (mg) and Interval	Date Started/Given	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)				Patient Outcomes (tick all that apply)			
<input type="checkbox"/> No intervention <input type="checkbox"/> Intervention unknown <input type="checkbox"/> Patient counselled/non-medical treatment <input type="checkbox"/> Discontinued Suspect Drug. Replaced with: _____ <input type="checkbox"/> Decreased Suspect Drug Dosage. New Dose: _____ <input type="checkbox"/> Treated ADR - with: _____ <input type="checkbox"/> Referred to hospital. Hospital Name: _____ <input type="checkbox"/> Other intervention (e.g. dialysis): _____				<input type="checkbox"/> ADR recovered/resolved <input type="checkbox"/> Recovering/resolving <input type="checkbox"/> Not recovered/not resolved <input type="checkbox"/> Life threatening <input type="checkbox"/> Impairment/disability <input type="checkbox"/> Congenital anomaly <input type="checkbox"/> Patient hospitalised or hospitalisation prolonged <input type="checkbox"/> Patient died: Date of death: _____ <input type="checkbox"/> Other: _____ <input type="checkbox"/> ADR reappeared after restarting suspect drug/similar drug (rechallenge)?: <input type="checkbox"/> N <input type="checkbox"/> Y <input type="checkbox"/> Not done <input type="checkbox"/> Unknown			
Laboratory Results				Additional Laboratory Results			
Lab Test	Test Result	Test Date		Lab Test	Test Result	Test Date	
Co-morbidities/Other Medical Condition(s)							
Reported by							
Name				E-mail			
Designation	<input type="checkbox"/> Nurse <input type="checkbox"/> Pharmacist <input type="checkbox"/> Doctor <input type="checkbox"/> Other:			Telephone			
Date reported				Signature			
THIS ADR REPORT IS NOT A CONFIRMATION THAT THE REPORTER OR THE SUSPECT MEDICINE(S) CAUSED THE ADR							V6.0 05/19

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CONSULTANT'S RESTRICTED MEDICINE FOLLOW UP AUTHORITY ¹		SG0006
A	PATIENT INFORMATION	Place of service
	No.....: _____ Rank/Title _____	Institute.....: _____
	Inits & surname: _____	Location.....: _____
	Gender.....: _____ (F/M) DOB.: _____	Service point: _____
	Admin unit.....: _____	Discipline.....: _____
	Contact tel.....: _____ Cons no.....: _____	
B	CONSULTATION PARTICULARS	
	Consultation date	Consultation time
	Laymans diagnosis (with consent of patient)	
	Allergies	
C	CONSULTANT'S FOLLOW-UP RECOMMENDATIONS	
	Medicine	Dose
	Date of next visit to consultant	
	Special Instructions	
D	Notes to Patient	HCP Information
	<p>This authority form is not a prescription, but an authority for a Medical Officer to prescribe restricted medication started by a consultant. Hand this form to the Medical Officer you are seeing at your Home Unit.</p> <p>This follow-up form is valid for <u>1 year only</u>.</p> <p>Ensure that you keep a copy of this authority form. Ask the pharmacist to return the form to you after your medication has been dispensed.</p>	HCP no.....: _____ Inits & surname : _____ HCP discipline..: _____ Duration of cons: _____ min Tel.: _____ Signature.....: _____ Qualification.....: _____

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¹ Military Medical Code List (MMCL)