Pharmacovigilance Department Form: Collection of ADR information.

ADVERSE EVENT REPORT

PATIENT INFORMATION:

Patient initials:	
COUNTRY:	

REPORT TYP	<u>e:</u>	Initial [up								
DATE OF BIR	TH AGE	RACE	SEX			HEIGH	T WEIG	GHT ONSET		ECOVERY ATE		
DD/MM/YYYY				Male				DD/MN	1/YYYY D	D/MM/YYYY		
				emale								
ADVERSE EV		IATION :										
ADVERSE EVENT(S) IN MEDICAL TERMS (diagnosis, if possible)						Seriousness criteria Check all appropriate to event						
Description of event:						Patient died						
						Involved or prolonged inpatient hospitalization						
								Involved persistent or significant disability or incapacity				
								Life-threatening				
								Congenital anomaly/birth defect				
								Other significant medical events				
HISTORY :					I				r only those f	indings		
						TEST / LABORATORY FINDINGS (enter only those findings necessary for AE diagnosis or course description)						
	RELEVANT M conditions su			-	5							
experien			,									
SUSPECT DRUG INFORMATION :												
Produc		Manufact urer	Batch/lo t	Expiry date	Dose	Route of use	Frequency	Therapy dates		Indication		
Brand	Generic		number					(from) DD/MM/YYYY	(To) DD/MM/YYYY	-		
name	Name											



Date:

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MACLEODS

L I								
CONCOMITAN	T DRUGS							
DRUG NAME(S)		DOSE		THERAPY DATES		REASON FOR USE		
			(from)	(То)				
Brand name	Generic Name		DD/MM/YYYY		DD/MM/YYYY			
ACTION TAKEN	I WITH SUSPE	CT DRUG (mark	all as appropria	ate)				
No Action Taken Withdrawn				Treatment taken				
			Did Rea	action Reappeared After Restarting of Drug?				
OUTCOME OF		ΔF						
		Date of recovery:			Condition still	present and unchanged		
Recovered	with sequela	е			Condition deteriorated			
Condition	improving				Death Autopsy: No Yes			
ASSESSMENT (OF CAUSALITY							
🗌 Probab		Possible		🗌 No	ot Related	Unknown		
REPORTER'S	INFORMATIC	DN :						
NAME, ADDRESS, TELEPHONE NUMBER AND EMAIL OF REPORTER					DATE OF THIS REPORT DD/MM/YYYY			
					∐ нср ∐ со	DNSUMER 🗌 OTHER		
					Signature:			
				Senders Contact details:				
				Email ID: eusafety@macleodspharma.com				

ADVERSE EVENT REPORT

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