Lenalidomide

Adverse Event Report Form

Reporter's details							
Title: (Mr, Mrs, Miss, Dr, etc)	First Name(s):	Surname:					
Job Title:							
Address:							
City, Town:	Country:						
Post code:	Country:						
Phone Number:	Fax Number:						
Email address:							

Patient information		
Patient ID (initials):	Age:	Date of birth: DD MM YYYY
Weight (Kg):		Height (cm):

Adverse event	
Overall diagnosis of the event	Event onset date: DD MM YYYY
	Event stop date: DD MM YYYY
	Or ongoing at time of reporting HR MIN
	(if less than 24 hours)

Description of adverse event	Outcome of adverse ever	nt
Symptoms and treatment	Recovered	TICK
	Recovered with sequele	TICK
	Not recovered	TICK
	Unknown	TICK
	Death	TICK
	Date of death	DD MM YYYY
	Possible cause of death	

If autopsy is performed please forward report. Please attach relevant clinical laboratory assessments to confirm the event.

Seriousness of adverse event (tick all that apply)				
Death	TICK			
Life-threatening	TICK			
Hospitalization or prolonged hospitalization	TICK			

Company

Address City, Town Country

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Adverse Event Report Form related to EU	Lenalidomide			
Persistent or significant disability or				

Persistent or significant disability or	HCK
incapacity	
Congenital anomaly/birth defect	TICK
Other medically important condition or	TICK
event	
Non-serious	TICK

Tel: Fax: Email:

Medical history (May be supplied as a copy of Medical file if up to date)						
Current or past relevant medical history (including concurrent illness, allergy, smoking, alcohol abuse)	YES	ΝΟ				
If YES please specify						

Suspect drug						
Drug, Dosage- form, Strength, Route (e.g. Tab 5mg, oral)	Dose & frequency	Batch no.	Therapy Start date	Therapy Stop date	Causal relationship 1= Not related 2 = Related	Indication for use of drug
			DD MM YY	DD MM YY		
			DD MM YY	DD MM YY		
			DD MM YY	DD MM YY		

Other medication (Medication taken during the past 3 months prior to the event - May be supplied as a copy of Medical file if up to date)						
Drug, Dosage- form, Strength, Route (e.g. Tab 5mg, oral)	Dose & frequency	Batch no.	Therapy Start date	Therapy Stop date	Causal relationship 1= Not related 2 = Related	Indication for use of drug
			DD MM YY	DD MM YY		
			DD MM YY	DD MM YY		
			DD MM YY	DD MM YY		

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Adverse Event Report Form related to EU RMP version 2.2 Lenalid						alidomide	
Action taken, suspect drug							
Continued unchanged TICK Continued, dose or dose TICK Withdrawn TICK N/A TICK regimen changed							TICK
Please specify if dose or dose regimen changed:				·			

Notification			
Initial report	TICK Final report	TICK Follow-up report	TICK
Name:			
Title:			

Signature:

Data Privacy statement:

All personal information will be strictly confidential and not used for any other purposes than preparing a report form.

Ver 1