Lenalidomide Grindeks (lenalidomide)

Patient card

Lenalidomide Grindeks (lenalidomide) - key elements of the patient card

Patient date of birth:	Patient name, surname:
Physician name:	
Institution:	;
Telephone number during office hours:	······
Telephone number after office hours:	
Information for Healthcare Prof	fessionals
1. This patient is receiving lenalidomid	le for the treatment of:
2. Type of patient (tick one):	
☐ Woman of non-childbearing po	otential
Male	
☐ Woman of childbearing potent * fill out a pregnancy test schedule	ial*
3. Before the first prescription, the lenalidomide's expected teratogenici	patient has been given instructions about ty and the need to avoid pregnancy:
Physician name	
Signature	
Date	

4. Pregnancy test schedule

	Patient is					
Date of Visit	using one effective method of	Date of pregnancy test	Pregnancy test result***	Next appointment date	Date of pomalidomide prescription	Physician signature
	contraception		□n :::			
			Positive			
☐ Yes ☐No	☐ Yes ☐No		☐Negative			
			Unclear			
			□Not done			
☐ Yes ☐No		Positive				
	☐ Yes ☐No	\exists_{No}	Negative			
			Unclear			
			Not done			
			Positive			
	☐ Yes ☐No		Negative			
			Unclear			
			Not done			
			Positive			
	☐ Yes ☐No		Negative			
			Unclear			
			Not done			
☐ Yes ☐No		Positive				
	□ Ves □No		Negative			
			Unclear			
			Not done			
			Positive			
☐ Yes ☐No		■ Negative				
			Unclear			
			☐Not done			
☐ Yes ☐No		Positive				
		■Negative				
		1110	Unclear			
			☐Not done			
☐ Yes ☐No		Positive				
	□ Ves □No		Negative			
			Unclear			
			☐Not done			
☐ Yes ☐			Positive			
	☐ Yes ☐No		Negative			
			Unclear			
			☐Not done			
			Positive			
	☐ Yes ☐No		Negative			
			Unclear			
			Not done			

^{***}Women of childbearing potential must have a medically supervised pregnancy tests with a minimum sensitivity of 25 mIU/mL prior to starting the treatment once she has been established on effective contraception for at least 4 weeks, every 4 weeks and 4 weeks after the end of therapy except in the case of confirmed tubal sterilisation. This requirement includes women of childbearing potential who practice absolute and continuous abstinence. For further information see Summary of Product Characteristics.

Suspected adverse reactions that may occur during treatment with Lenalidomide Grindeks should be reported via adverse drug reactions (ADRs) to Malta Medicines Authority via the ADR Reporting Website: www.medicinesauthority.gov.mt/adrportal.

For further information regarding this medication please contact local representative: EJ Busuttil Ltd.

Phone: 2147184 (service is available 24/7)

Email: safety@ejbusuttil.com

Office Adress:

Busuttil Buildings, Triq I-Ghadam, Central Business District Zone 1, Birkirkara CBD1060 MALTA

Adverse events should also be reported to AS GRINDEKS. Krustpils iela 53, Rīga, LV-1057, Latvia (Marketing Authorisation Holder) at: +371 67083644, E-mail: vigilance@grindeks