LENALIDOMIDE PATIENT CARD

Australia

Patient initials:	Date of birth:
Doctor's name:	
Doctor's phone number:	

Information for Patients and Healthcare Professionals

Lenalidomide is structurally related to thalidomide and is expected to cause severe birth defects or death to an unborn baby, therefore:

- Female patients of childbearing potential must always use effective contraception
- Female patients of childbearing potential must have a negative pregnancy test every four weeks before each prescription refill to ensure the patient is not pregnant.
- Male patients with pregnant partners or with partners of childbearing potential not using effective contraception must always use condoms (even if the man has had a vasectomy)
- ➤ If a female patient or female partner of a male patient suspects they are pregnant, they must contact their prescriber immediately
- > You MUST immediately contact your prescriber if you experience any adverse effects.

For complete information on the side effects of Lenalidomide, patients should read the Consumer Medicine Information (CMI), while HCPs should read the Product Information (PI).

Approved by TGA: November 2022

LENALIDOMIDE PATIENT CARD

Australia

Prescriber to complete the following:				
1. INDICATIONS				
Multiple myeloma:	☐ Patient with Newly Diagnosed Multiple Myeloma (NDMM) who has undergone autologous stem cell transplantation. ☐ Patient with previously untreated multiple myeloma who is not eligible for transplant ☐ Patient who has received at least one prior therapy			
Myelodysplastic Syndromes (MDS):	☐ Transfusion-dependent anaemia due to low- or intermediate risk myelodysplastic syndromes associated with a deletion 5q cytogenetic abnormality with or without additional cytogenetic abnormalities			
Other:	If others, Please Specify			
2. STATUS OF PATIENT (tic	k one)			
Male				
		Using condoms:		
		Yes/No/Not Applicable		
Women of Non-Childbearing Potential				
Women of childbearing potential				
(Complete Section 4)		Using effective		
		contraception: Yes/No		
3. Before the first administration, counselling was provided on the expected teratogenicity of Lenalidomide in humans and the need to avoid pregnancy.				
Patient's Signature: Doctor's Signature:				

LENALIDOMIDE PATIENT CARD

	-	
Δ	ustra	lıa

Date:	Date:

4. FOR WOMEN OF CHILDBEARING POTENTIAL							
Date of visit	An effective method of contraception is being used (Yes/No)	Date of pregnancy test	Pregnancy Test Result IF POSITIVE, DO NOT DISPENSE	Prescription Date	Doctor's signature	Date of Dispensing	Pharmacist's Name and Signature

A completed electronic Prescription Authorisation Form (ePAF) must accompany each prescription to confirm that the patient continues to use effective contraception (if required) and, in the case of a female patient of childbearing potential, has a negative pregnancy test performed every four weeks prior each prescription.

Emergency contact information:

Emergency Prescriber Contact:

Further information is available in the patient brochure.

Cipla

For information and questions on the Risk Management of Cipla products, the Pregnancy Prevention Programme, pharmacy registrations, the pregnancy reporting form and for Adverse events, please contact Cipla:

Email: Lenalidomide.cipla@cipla.com

Approved by TGA: November 2022