

# Pomalidomide Rowex Pregnancy Prevention Programme

## Risk Awareness From-

### Women with Non-Childbearing Potential

Pomalidomide

Please complete this form to report a pregnancy in a patient (or in a female partner of a male patient) treated with pomalidomide. Please send them immediately. Contact details are given below.

#### **REPORTING OF ADVERSE REACTIONS**

Suspected adverse reactions and medication errors should be reported either to:

**ADR Reporting, The Medicines Authority, Post-Licensing Directorate,**  
Sir Temi Zammit Buildings, Malta Life Sciences Park, San Gwann SGN 3000, Malta  
Website:  
[www.medicinesauthority.gov.mt](http://www.medicinesauthority.gov.mt)  
e-mail: [postlicensing.medicinesauthority@gov.mt](mailto:postlicensing.medicinesauthority@gov.mt)

OR

**Marketing Authorisation Holder:**  
Rowex Limited Medical Information  
Tel: 0877941968  
Email: [adverse.event.ireland@sandoz.net](mailto:adverse.event.ireland@sandoz.net)

**Approved by the Malta Medicines Authority on the 13<sup>th</sup> January 2026.**



5. Of the thromboembolic risk and possible requirement to take thromboprophylaxis during treatment with pomalidomide.	Tick
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**Patient: please read thoroughly and initial the adjacent box if you agree with the statement**

I understand that severe birth defects may occur with the use of pomalidomide. I have been warned by my prescriber that any unborn baby has a high risk of birth defects and could even die if a woman is pregnant or becomes pregnant while taking pomalidomide.	Patient Initials
I understand that pomalidomide will be prescribed ONLY for me. I must not share it with ANYONE.	Patient Initials
I have read the Pomalidomide Patient Guide and understand the contents, including the information about other possible important health problems (side effects) associated with the use of pomalidomide.	Patient Initials
I know that I cannot donate blood while taking pomalidomide (including dose interruptions) and for at least 7 days after stopping treatment.	Patient Initials
I understand that I must return any unused pomalidomide capsules to my pharmacy at the end of my treatment	Patient Initials
I have been informed about the thromboembolic risk and possible requirement to take thromboprophylaxis during treatment with pomalidomide	Patient Initials
I understand that my prescriber will send or may provide me with a completed 'Prescription Authorisation Form' with each pomalidomide prescription for the pharmacy	Patient Initials
I understand that the 'Prescription Authorisation Form' contains non-identifiable information about me, which will ensure pomalidomide is dispensed safely. This information may be used by the Marketing Authorisation Holder and the distributor for the product I receive and the Malta Medicines Authority (MMA) evaluate the safe use of pomalidomide.	Patient Initials

**Patient Confirmation**

I confirm that I understand and will comply with the requirements of the POMALIDOMIDE Pregnancy Prevention Programme, and I agree that my prescriber can initiate my treatment with pomalidomide.

Your personal data is used solely for the purpose of entering you into the Pomalidomide Pregnancy Prevention Programme and is processed by the marketing authorisation holder (MAH) of the pomalidomide product you receive, its third party service providers and any worldwide Affiliates the MAH may have, for the purposes of compliance with the Risk Management Plan legal obligations and for storage purposes. Third party service providers include, for example, the distributor of the pomalidomide product you receive.

Your pharmacist can confirm the details of the MAH for the pomalidomide product you are given and this will also be mentioned on the packaging and package leaflet. Queries on how your personal data will be processed can be directed to the MAH in question, by consulting their publicly available information (e.g. on their website) which details how they process your personal data and provides a contact point for any queries in relation to their use of your personal data.

Patient Signature		Date	DD	MM	YYYY
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**Statement of the interpreter (where appropriate)**

I have interpreted the information above to the patient/parent/carer to the best of my ability and in a way in which I believe she/he/they can understand. She/he/they agree to follow the necessary precautions to prevent an unborn child being exposed to pomalidomide.

Signed		Name: (Print)		Date	DD	MM	YYYY
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**Important Safety Information:**

This risk minimisation material is being provided by Rowex Ltd. For Further Information, please refer to the Summary of Product Characteristics (SmPC) for the respective medicinal product from the relevant Marketing Authorisation holder.