



This medicinal product is subject to additional monitoring. This will allow quick identification of new safety information.

Pomalidomide Pregnancy Report Form

Pregnancy reports must be sent to the relevant Medical information team IMMEDIATELY

This form must be returned to the MAH who provided the product. Please see contact details below:

NOTE: Please use the first three letters of the month (e.g.: JAN)

Date of awareness: DDMMYYYY

Patient Data

Sex of Patient: ☐ Female ☐ Male

☐ Pregnancy of Patient

☐ Pregnancy of Patient's Partner **OR**

☐ Exposure of a Pregnant Female (complete information below)

Pregnant Woman's Initials (F, M, L):

Date of Birth: DDMMYYYY

Age:

Patient Initials (F, M, L): (Who received drug)

Date of Birth: DDMMYYYY

Age:

Drug Name:

Date of First Dose: DDMMYYYY

Date of Last Dose: DDMMYYYY

Pregnancy Initially Diagnosed By:

☐ Home Urine Test

☐ Office Urine Test

☐ Serum Test

Date of Pregnancy Test: DDMMYYYY

Last Menstrual Period: DDMMYYYY

Female is Currently: weeks pregnant **OR** ☐ No longer Pregnant ☐ Unknown

Female has Elected to: ☐ Carry Pregnancy to Term

Expected Date of Delivery: DDMMYYYY

☐ Terminate Pregnancy

Date Performed or Pending: DDMMYYYY

Reporter's Information:

Reporter's Name:	Date: DDMMYYYY
Reporter's Contact Information/ Address:	Reporter's Signature:
Reporter's E-mail Address:	Reporter's Phone Number:

Patient's Prescribing Physician's Information:

Physician's Name:	Date: DDMMYYYY
Physician's Contact Information/ Address:	Physician's Signature:
Physician's E-mail Address:	Physician's Phone Number:
	Physician's Fax Number:

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Background Information on Reason for Pregnancy

Was patient erroneously considered not to be of childbearing potential? ☐ Yes ☐ No

If yes, state reason for considering not to be of childbearing potential

- | | | |
|---|---------------------------|--------------------------|
| <input type="radio"/> Age ≥ 50 years and naturally amenorrhoeic* for ≥ 1 year
*amenorrhoea following cancer therapy or during breastfeeding does not rule out childbearing potential | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Premature ovarian failure confirmed by a specialist gynaecologist | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Previous bilateral salpingo-oophorectomy, or hysterectomy | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> XY genotype, Turner syndrome, uterine agenesis. | <input type="radio"/> Yes | <input type="radio"/> No |

Indicate from the list below what contraception was used

- | | | |
|--|---------------------------|--------------------------|
| <input type="radio"/> Implant | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Levonorgestrel-releasing intrauterine system (IUS) | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Medroxyprogesterone acetate depot | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Tubal sterilization (specify below) | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Tubal ligation | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Tubal diathermy | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Tubal chips | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Sexual intercourse with a vasectomized male partner only; vasectomy must be confirmed by two negative semen analyses | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Ovulation inhibitory progesterone-only pills (i.e. desogestrel) | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Other progesterone-only pills | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Combined oral contraceptive pill | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Other intra-uterine devices | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Condoms | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Cervical cap | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Sponge | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Withdrawal | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Other | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> None | <input type="radio"/> Yes | <input type="radio"/> No |

Indicate from the list below the reason for contraceptive failure

- | | | |
|---|---------------------------|--------------------------|
| <input type="radio"/> Missed oral contraception | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Other medication or intercurrent illness interacting with oral contraception | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Identified mishap with barrier method | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Unknown | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Had the patient committed to complete and continuous abstinence | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Was the drug started despite patient already being pregnant | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Did patient receive educational materials on the potential risk of teratogenicity | <input type="radio"/> Yes | <input type="radio"/> No |
| <input type="radio"/> Did patient receive instructions on need to avoid pregnancy | <input type="radio"/> Yes | <input type="radio"/> No |

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Background Information on Reason for Pregnancy

Prenatal information

Date of Last Menstrual Period:

D	D	M	O	N	Y	Y	Y	Y
---	---	---	---	---	---	---	---	---

 Expected Delivery Date:

D	D	M	O	N	Y	Y	Y	Y
---	---	---	---	---	---	---	---	---

Pregnancy test

Urine Qualitative ☐ Reference Range:

--	--	--	--	--	--	--	--	--

 Date:

D	D	M	O	N	Y	Y	Y	Y
---	---	---	---	---	---	---	---	---

Serum Quantitative ☐ Reference Range:

--	--	--	--	--	--	--	--	--

 Date:

D	D	M	O	N	Y	Y	Y	Y
---	---	---	---	---	---	---	---	---

Past Obstetric History

Year of Pregnancy	Outcome	Gestational Age	Type of Delivery																						
<table border="1" style="display: inline-table; vertical-align: middle;"><tr><td>Y</td><td>Y</td><td>Y</td><td>Y</td></tr></table>	Y	Y	Y	Y	<input type="radio"/> Spontaneous abortion <input type="radio"/> Therapeutic abortion <input type="radio"/> Live birth <input type="radio"/> Still birth	<table border="1" style="display: inline-table; vertical-align: middle;"><tr><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td></tr></table>										<table border="1" style="display: inline-table; vertical-align: middle;"><tr><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td></tr></table>									
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Y	Y	Y	Y																						

Birth defects

Was there any birth defect from any pregnancy? ☐ Yes ☐ No ☐ Unknown

Is there any family history of any congenital abnormality abstinence? ☐ Yes ☐ No ☐ Unknown

If yes to either of these questions, please provide details below:

--

Maternal Past Medical History

Condition	Dates										Treatment	Outcome
	From:	D	D	M	O	N	Y	Y	Y	Y		
	To:	D	D	M	O	N	Y	Y	Y	Y		
	From:	D	D	M	O	N	Y	Y	Y	Y		
	To:	D	D	M	O	N	Y	Y	Y	Y		
	From:	D	D	M	O	N	Y	Y	Y	Y		
	To:	D	D	M	O	N	Y	Y	Y	Y		
	From:	D	D	M	O	N	Y	Y	Y	Y		
	To:	D	D	M	O	N	Y	Y	Y	Y		
	From:	D	D	M	O	N	Y	Y	Y	Y		
	To:	D	D	M	O	N	Y	Y	Y	Y		
	From:	D	D	M	O	N	Y	Y	Y	Y		
	To:	D	D	M	O	N	Y	Y	Y	Y		

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Maternal Current Medical Conditions

Condition	From	Treatment
	D D M O N Y Y Y Y	
	D D M O N Y Y Y Y	
	D D M O N Y Y Y Y	
	D D M O N Y Y Y Y	
	D D M O N Y Y Y Y	
	D D M O N Y Y Y Y	
	D D M O N Y Y Y Y	

Maternal Social History

Alcohol ☐ Yes ☐ No Tobacco ☐ Yes ☐ No IV or recreational drug use ☐ Yes ☐ No

If yes, amount/units per day: If yes, amount per day: If yes, provide details:

Maternal medication during pregnancy and in 4 weeks before pregnancy

(including herbal, alternative and over the counter medicines and dietary supplements)

Medication/treatment	Dates	Indication
	Start Date: D D M O N Y Y Y Y Stop Date/Continuing: D D M O N Y Y Y Y	
	Start Date: D D M O N Y Y Y Y Stop Date/Continuing: D D M O N Y Y Y Y	
	Start Date: D D M O N Y Y Y Y Stop Date/Continuing: D D M O N Y Y Y Y	
	Start Date: D D M O N Y Y Y Y Stop Date/Continuing: D D M O N Y Y Y Y	
	Start Date: D D M O N Y Y Y Y Stop Date/Continuing: D D M O N Y Y Y Y	
	Start Date: D D M O N Y Y Y Y Stop Date/Continuing: D D M O N Y Y Y Y	
	Start Date: D D M O N Y Y Y Y Stop Date/Continuing: D D M O N Y Y Y Y	

Name of person completing this form

Name: Signature:

Date: D D M O N Y Y Y Y

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Data Privacy Notice

Your personal data will be processed by the relevant marketing authorisation holder, and its worldwide affiliates, to the extent and for as long as necessary, for the purposes of the compliance with drug safety legal obligations and for storage purposes. Should you have any queries in relation to the use of your personal data please contact the relevant marketing authorisation holder.

Reporter's Signature (required):

Signature:

Date signed:

D	D	M	O	N	Y	Y	Y	Y
---	---	---	---	---	---	---	---	---

Thank you for providing information that will assist us in our commitment to patient safety.