

**Lenalidomide**  
**Event-Specific Questionnaire for HCP - Pregnancy Outcome Form**  
(Patient or Partner of Patient)

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)

**Reporter information**

Reporter Name:	
Address:	
City, County, Country:	
Phone No.:	
Fax No.:	

**Patient information**

Patient ID:		Date of Birth:	<table><tr><td>D</td><td>D</td><td>M</td><td>O</td><td>N</td><td>Y</td><td>Y</td><td>Y</td><td>Y</td></tr></table>	D	D	M	O	N	Y	Y	Y	Y	Ethnicity:	
D	D	M	O	N	Y	Y	Y	Y						

**Partner of patient information**

<input type="radio"/> Not applicable	Ethnicity:	
--------------------------------------	------------	--

**Pregnancy outcome**

Date of delivery:	<table><tr><td>D</td><td>D</td><td>M</td><td>O</td><td>N</td><td>Y</td><td>Y</td><td>Y</td><td>Y</td></tr></table>	D	D	M	O	N	Y	Y	Y	Y	Gestation age at delivery:	
D	D	M	O	N	Y	Y	Y	Y				

- Normal ☐ No ☐ Yes
- C-section ☐ No ☐ Yes
- Induced ☐ No ☐ Yes
- Ectopic pregnancy ☐ No ☐ Yes
- Elective termination ☐ No ☐ Yes
- Spontaneous abortion ( $\leq 20$  weeks) ☐ No ☐ Yes
- Foetal death/stillbirth ( $>20$  weeks) ☐ No ☐ Yes
- Were the products of conception examined? ☐ No ☐ Yes

Date:	<table><tr><td>D</td><td>D</td><td>M</td><td>O</td><td>N</td><td>Y</td><td>Y</td><td>Y</td><td>Y</td></tr></table>	D	D	M	O	N	Y	Y	Y	Y
D	D	M	O	N	Y	Y	Y	Y		
Weeks from LMP:										

If yes, was the foetus normal? ☐ No ☐ Yes ☐ Unknown If no, describe below:

--

**Obstetrics information**

Complications during pregnancy	<input type="radio"/> No <input type="radio"/> Yes	If yes, please specify	
Complications during labour/delivery	<input type="radio"/> No <input type="radio"/> Yes	If yes, please specify	
Post-partum maternal complications	<input type="radio"/> No <input type="radio"/> Yes	If yes, please specify	

**Foetal outcome**

Live normal infant	<input type="radio"/> No <input type="radio"/> Yes		
Foetal distress	<input type="radio"/> No <input type="radio"/> Yes		
Intra-uterine growth retardation	<input type="radio"/> No <input type="radio"/> Yes		
Neonatal complication	<input type="radio"/> No <input type="radio"/> Yes	If yes, please specify	
Birth defect noted?	<input type="radio"/> No <input type="radio"/> Yes	If yes, please specify	

Sex: ☐ Male ☐ Female Birth weight: \_\_\_\_ lbs \_\_\_\_ oz. or \_\_\_\_ kg Length: \_\_\_\_ inches or \_\_\_\_ cm.

Apgar score: 1 min: \_\_\_\_ 5 min: \_\_\_\_ 10 min: \_\_\_\_ ☐ Unknown

**Signature of person completing this form**

Signature:		Date:	<table><tr><td>D</td><td>D</td><td>M</td><td>O</td><td>N</td><td>Y</td><td>Y</td><td>Y</td><td>Y</td></tr></table>	D	D	M	O	N	Y	Y	Y	Y
D	D	M	O	N	Y	Y	Y	Y				

**Lenalidomide**  
**Event-Specific Questionnaire for HCP - Pregnancy Outcome Form**  
(Patient or Partner of Patient)

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)

**Drug Safety 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.