

Ireland

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

The form must be returned to the Marketing Authorisation Holder (MAH) who provided the product. Please see the relevant MAH details below:

Accord Healthcare Ireland Ltd.	Email: medinfo@accord-healthcare.com	Tel: 0044 1271 385257
AS Grindeks.	Email: adrian.curley@grindeks.ie	Tel: +353 (0)87 298 8226
Teva Pharmaceuticals Ireland.	Email: medinfo@tevauk.com	Tel: 0044 207 540 7117
Clonmel Healthcare	Email: medicalinformation@clonmel-health.ie	Tel: (052) 6177777
Viartis Limited	Email: pv.ireland@viartis.com	Tel: 0044 1707 853000 (select option 5)
Rowex Ltd	Email: mi.ireland@sandoz.net	Tel: 087 794 1968

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

Date of awareness:	D	D	M	O	N	Y	Y	Y	Y
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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: Age:

Patient Initials (F, M, L): (Who received drug) Date of Birth: Age:

Drug Name:

Date of First Dose: Date of Last Dose:

Pregnancy Initially Diagnosed By:

- Home Urine Test
 Office Urine Test
 Serum Test

Date of Pregnancy Test: Last Menstrual Period:

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

Female has Elected to: Carry Pregnancy to Term Expected Date of Delivery:

Terminate Pregnancy Date Performed or Pending:

Reporter's Information:

Reporter's Name:	<input type="text"/>	Date:	<input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>
Reporter's Contact Information/ Address:	<input type="text"/>	Reporter's Signature:	<input type="text"/>
Reporter's E-mail Address:	<input type="text"/>	Reporter's Phone Number:	<input type="text"/>

Prescriber's Information:

Prescriber's Name:	<input type="text"/>	Date:	<input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>
Prescriber's Contact Information/ Address:	<input type="text"/>	Prescriber's Signature:	<input type="text"/>
Prescriber's E-mail Address:	<input type="text"/>	Prescriber's Phone Number:	<input type="text"/>
		Prescriber's Fax Number:	<input type="text"/>

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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

Age ≥ 50 years and naturally amenorrhoeic* for ≥ 1 year
*amenorrhoea following cancer therapy or during breastfeeding does not rule out childbearing potential Yes No

Premature ovarian failure confirmed by a specialist gynaecologist Yes No

Previous bilateral salpingo-oophorectomy, or hysterectomy Yes No

XY genotype, Turner syndrome, uterine agenesis. Yes No

Indicate from the list below what contraception was used

Implant Yes No

Levonorgestrel-releasing intrauterine system Yes No

Medroxyprogesterone acetate depot Yes No

Tubal sterilization (specify below) Yes No

Tubal ligation Yes No

Tubal diathermy Yes No

Tubal chips Yes No

Sexual intercourse with a vasectomized male partner only; vasectomy must be confirmed by two negative semen analyses Yes No

Ovulation inhibitory progesterone-only pills (i.e. desogestrel) Yes No

Other progesterone-only pills Yes No

Combined oral contraceptive pill Yes No

Other intra-uterine devices Yes No

Condoms Yes No

Cervical cap Yes No

Sponge Yes No

Withdrawal Yes No

Other Yes No

None Yes No

Indicate from the list below the reason for contraceptive failure

Missed oral contraception Yes No

Other medication or intercurrent illness interacting with oral contraception Yes No

Identified mishap with barrier method Yes No

Unknown Yes No

Had the patient committed to complete and continuous abstinence Yes No

Was the drug started despite patient already being pregnant Yes No

Did patient receive educational materials on the potential risk of teratogenicity Yes No

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Did patient receive instructions on need to avoid pregnancy Yes No

Background Information on Reason for Pregnancy

Prenatal information

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Pregnancy test

<input type="radio"/>		
<input type="radio"/>		D M O N Y Y Y Y

Past Obstetric History

Year of Pregnancy	Outcome	Gestational Age	Type of Delivery
Y Y Y Y	<input type="radio"/> Spontaneous abortion <input type="radio"/> Therapeutic abortion <input type="radio"/> Live birth <input type="radio"/> Still birth		
Y Y Y Y	<input type="radio"/> Spontaneous abortion <input type="radio"/> Therapeutic abortion <input type="radio"/> Live birth <input type="radio"/> Still birth		
Y Y Y Y	<input type="radio"/> Spontaneous abortion <input type="radio"/> Therapeutic abortion <input type="radio"/> Live birth <input type="radio"/> Still birth		
	<input type="radio"/> Spontaneous abortion <input type="radio"/> Therapeutic abortion <input type="radio"/> Live birth <input type="radio"/> Still birth		
	<input type="radio"/> Spontaneous abortion <input type="radio"/> Therapeutic abortion <input type="radio"/> Live birth <input type="radio"/> Still birth		

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 O N Y Y Y Y		
	To: D D O N Y Y Y Y		

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	To:	D	D	O	N	Y	Y	Y	Y	Y	Y	
	To:	D	D	O	N	Y	Y	Y	Y	Y	Y	

Maternal Current Medical Conditions

Condition	From	Treatment
	D D O N Y Y Y Y Y	
	D D O N Y Y Y Y Y	
	D D O N Y Y Y Y Y	
	D D O N Y Y Y Y Y	
	D D O N Y Y Y Y Y	
	D D O N Y Y Y Y Y	
	D D O N Y 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:

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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	

Name of person completing this form

Name:	Signature:
Date: D D M O N Y Y Y Y Y	

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:	
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On behalf of Accord Healthcare Ireland Ltd., Clonmel Healthcare Ltd., Viatriis Limited, AS Grindeks, Teva Pharmaceuticals Ireland, and Rowex Ltd thank you for providing information that will assist us in our commitment to patient safety.

