

Revlimid®

Information for Healthcare
Professionals

Checklist

**Checklist for commencing Revlimid treatment in
Women of Childbearing Potential**

Patient Name

Date of Birth

Counselling

| | Insert <input type="checkbox"/> or N/A |
|---|--|
| Inform of expected teratogenic risk to the unborn child | |
| Inform of the need for effective contraception 4 weeks before starting treatment, during treatment interruption, throughout the entire duration of treatment and for 4 weeks after the end of treatment <u>or</u> absolute and continued abstinence | |
| Inform that that even if she has amenorrhoea she must comply with advice on contraception | |
| Confirm patient is capable of complying with contraceptive measures | |
| Inform of the expected consequences of pregnancy and the need to stop treatment and consult rapidly if there is a risk of pregnancy | |
| Confirm patient agrees to undergo pregnancy testing at 4 weekly intervals unless confirmed tubal sterilisation | |
| Inform of hazards and necessary precautions associated with use of Revlimid | |
| Inform patient not to share medication | |
| Inform to return unused capsules to pharmacist | |
| Inform not to donate blood whilst taking Revlimid or for one week after stopping | |
| Inform about the thromboembolic risk and possible requirement to take thromboprophylaxis during treatment with Revlimid | |

Contraceptive referral

| | |
|--------------------------------------|--|
| Contraceptive referral required | |
| Contraceptive referral made | |
| Contraceptive consultation completed | |

Contraception

Patient is currently established on one of the following for at least 4 weeks

| | |
|--|--|
| Implant | |
| Levonorgestrel-releasing intrauterine system (IUS) | |
| Medroxyprogesterone acetate depot | |
| Tubal Sterilisation | |
| Sexual intercourse with a vasectomised male partner only: vasectomy must be confirmed by two negative semen analyses | |
| Ovulation inhibitory progesterone only pill (i.e. desogestrel) | |

Abstinence

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|---|--|
| Patient commits to complete and absolute abstinence | |
|---|--|

Pregnancy test

| | |
|---|--|
| Negative pregnancy test before starting treatment even if absolute and continued abstinence | |
|---|--|

TREATMENT CANNOT START UNTIL PATIENT IS ESTABLISHED ON EFFECTIVE METHOD OF CONTRACEPTION FOR 4 WEEKS OR COMMITS TO COMPLETE AND CONTINUED ABSTINENCE AND PREGNANCY TEST IS NEGATIVE

**Checklist for commencing Revlimid treatment in
Women NOT of Childbearing Potential**

Patient Name

Date of Birth

Counselling

| | Insert \checkmark or N/A |
|---|--|
| Inform of expected teratogenic risk to the unborn child | |
| Inform patient not to share medication | |
| Inform to return unused capsules to pharmacist | |
| Inform not to donate blood whilst taking Revlimid or for one week after stopping | |
| Inform of hazards and necessary precautions associated with use of Revlimid | |
| Inform about the thromboembolic risk and possible requirement to take thromboprophylaxis during treatment with Revlimid | |

**The following criteria have been met to determine patient is woman NOT of
childbearing potential**

| | |
|---|--|
| Age \geq 50 years and naturally amenorrhoeic* for \geq one year not induced by chemotherapy | |
| Premature ovarian failure confirmed by specialist gynaecologist | |
| Bilateral salpingo-oophorectomy | |
| XY genotype, Turner's syndrome, uterine agenesis | |

*Amenorrhoea following cancer therapy or during lactation does not rule out childbearing potential.

Checklist for commencing Revlimid treatment in Men

Patient Name

Date of Birth

Counselling

| | Insert √ or N/A |
|--|----------------------------|
| Inform of expected teratogenic risk to the unborn child | |
| Inform of need to use condoms (even if he has had vasectomy) throughout treatment duration, during dose interruption, and for one week after cessation of treatment if partner is pregnant or of childbearing potential who is not using effective contraception. | |
| Inform patient not to share medication | |
| Inform to return unused capsules to pharmacist | |
| Inform not to donate blood whilst taking Revlimid or for one week after stopping | |
| Inform of hazards and necessary precautions associated with use of Revlimid | |
| Inform about the thromboembolic risk and possible requirement to take thromboprophylaxis during treatment with Revlimid | |
| Inform about which are effective contraceptive methods that the female partner of a male patient can use | |
| Inform that if his female partner becomes pregnant whilst he is taking Revlimid or shortly after he has stopped taking Revlimid, he should inform his treating physician immediately and that it is recommended to refer the female partner to a physician specialised or experienced in teratology for evaluation and advice. | |