Treatment Initiation Form Revlimid
Men 28 April 2010

Treatment Initiation Form

Men

This Treatment Initiation Form must be completed for each male patient prior to the initiation of their Revlimid treatment. The form should be retained with their medical records, and a copy provided to the patient.

The aim of the Treatment Initiation Form is to assist both prescribers and patients to ensure all necessary steps are taken to prevent foetal exposure to lenalidomide and to assist in ensuring that patients are fully informed of and understand the risk of teratogenicity and other adverse effects associated with the use of lenalidomide. It is not a contract and does not absolve anybody from his/her responsibilities with regard to the safe use of the product and prevention of foetal exposure.

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Patient Name Date of Birth

I have fully explained to the patient named above the nature, purpose and risks of the treatment associated with Revlimid, especially the risks to women of childbearing potential. I will comply with all my obligations and responsibilities as the prescribing physician of Revlimid

	Insert √
Inform of need to use condoms throughout treatment duration, during	
dose interruption, and for one week after cessation of treatment if partner	
is pregnant or of childbearing potential.	
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	

Physician Name

Physician Signature

Date

Patient Name Date of Birth Patient: please read thoroughly and initial the adjacent box if you agree with the statement

My doctor has explained to me and I have understood the possible risks and the possible benefits associated with Revlimid® (lenalidomide). I have had the opportunity to ask questions and I have understood the answers provided to those questions.	Patient initials
I have received, read and understood the Patient Information Brochure	Patient initials
I understand that Revlimid® (lenalidomide) has been prescribed for me personally and that I should not share it with any other person even if they have the same condition as me. I should store Revlimid® (lenalidomide) out of the reach of children.	Patient initials
I will return any unused capsules for my pharmacist.	Patient initials
I will not donate blood during treatment or for one week after stopping treatment	Patient initials
I understand that Revlimid® (lenalidomide) is expected to be harmful to the unborn child	Patient initials
I agree to use condoms throughout treatment duration, during dose interruption, and for one week after cessation of treatment if my partner is pregnant or of childbearing potential.	Patient initials
If my partner were to become pregnant during my treatment with Revlimid® (lenalidomide), I will advise her to seek medical advice immediately.	Patient initials

Patient Confirmation

I confirm that I understand and will comply with the requirements of the Revlimid Pregnancy Prevention Programme, and I agree that my doctor can initiate my treatment with Revlimid

Patient Signature Date