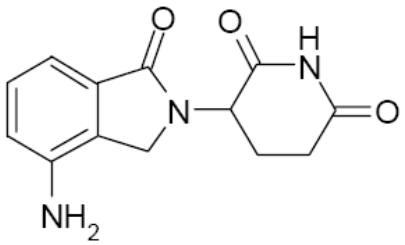
Attachment 1: Product information for AusPAR Lenalidomide (Revlimid) Celgene Pty Ltd PM-2014-02792-1-4 Final 5 February 2016. This Product Information was approved at the time this AusPAR was published.

Revlimid® (lenalidomide) capsules Product Information

Teratogenic Effects:
Revlimid (lenalidomide) is structurally related to thalidomide. Thalidomide is a known human teratogen that causes severe life-threatening human birth defects. If lenalidomide is taken during pregnancy, it may cause birth defects or death to an unborn baby. Women should be advised to avoid pregnancy whilst taking Revlimid (lenalidomide), during dose interruptions, and for 4 weeks after stopping the medication.

1. Name of the Medicine

Australian approved name:	Lenalidomide
Molecular formula:	$C_{13}H_{13}N_3O_3$
Molecular weight:	259.25 g/mol
ATC code:	L04 AX04
Chemical name:	3-(4'-amino-1,3-dihydro-1-oxo-2H-isoindol-2-yl)-2,6-piperidinedione
Chemical Abstract Service (CAS) registry number:	191732-72-6
Chemical structure:	

2. Description

Lenalidomide is an off-white to pale-yellow solid, with a melting point between 265ºC and 270ºC. Lenalidomide is generally more soluble in organic solvents but exhibits the greatest solubility in 0.1N HCl buffer. The solubility of lenalidomide in water and at pH 1.21 is < 1.5 mg/mL and 18 mg/mL, respectively.

Lenalidomide has an asymmetric carbon atom and can therefore exist as the optically active forms S(-) and R(+). Lenalidomide is produced as a racemic mixture with a net optical rotation of zero.

List of Excipients

Revlimid capsules contain the following excipients: lactose, anhydrous; cellulose, microcrystalline; croscarmellose sodium; and magnesium stearate.

The capsule shells contain gelatin, titanium dioxide (E171), black ink, and the following colourants: 2.5 mg (FD&C Blue#2 [indigo carmine; E132], and yellow iron oxide [E172]); 7.5 mg (yellow iron oxide [E172]); 10 mg (FD&C Blue#2 [E132], and yellow iron oxide [E172]); 15 mg (FD&C Blue#2 [E132]); and 20 mg (FD&C Blue#2 [E132], and yellow iron oxide [E172]).

The black printing ink used on the capsules contains shellac, ethanol, isopropyl alcohol, butan-1-ol, propylene glycol, water-purified, ammonium hydroxide, potassium hydroxide, and black iron oxide [E172].

3. Pharmacology

3.1. Pharmacodynamic Properties

Pharmacotherapeutic group: Immunomodulating agent.

3.2. Mechanism of Action

Lenalidomide has a pleiotropic mechanism of action including immunomodulatory, anti-neoplastic, anti-angiogenic and pro-erythropoietic properties. Specifically, lenalidomide inhibits proliferation of certain haematopoietic tumour cells (including multiple myeloma [MM] plasma tumour cells and those with deletions of chromosome 5), enhances T cell- and Natural Killer (NK) cell-mediated immunity and increases the number of NK T cells, inhibits production of pro-inflammatory cytokines (e.g. TNF-α and IL-6) by monocytes, inhibits angiogenesis by blocking the migration and adhesion of endothelial cells and the formation of microvessels, and augments foetal haemoglobin production by CD34+ haematopoietic stem cells.

3.2.1. Cardiac Electrophysiology

A QTc study was conducted to evaluate the effects of lenalidomide on QT interval at single doses of 10 mg and 50 mg. A single dose of lenalidomide up to 50 mg is not associated with prolongation of the QT interval in healthy male subjects. This indicates that lenalidomide is not expected to result in clinically significant prolongation of the QT interval in patients at the approved therapeutic doses.

3.3. Pharmacokinetic Properties

3.3.1. Absorption

In healthy volunteers, lenalidomide is rapidly absorbed following oral administration with maximum plasma concentrations occurring between 0.6 and 1.5 hours post-dose. The maximum concentration (C_{max}) and area-under-the-concentration versus time curve (AUC) increase proportionately with increases in dose. Multiple dosing does not cause marked drug accumulation. In plasma, the relative exposures of the S- and R- enantiomers of lenalidomide are approximately 56% and 44%, respectively. The absolute bioavailability of lenalidomide has not been determined.

Co-administration with a high-fat and high-calorie meal in healthy volunteers reduces the extent of absorption, resulting in an approximately 20% decrease in AUC and 50% decrease in the C_{max} in plasma.

The pharmacokinetics of lenalidomide were very similar in subjects with myelodysplastic syndrome (MDS) compared to subjects with MM. In patients with low- or intermediate-1-risk MDS, a single 10 mg oral dose of lenalidomide was rapidly absorbed with a median time to maximum concentration (t_{max}) of around 1 hour post-dose. The mean terminal half-life was approximately 4 hours. Following multiple dosing of 10 mg per day for 14 days there was no accumulation of lenalidomide in plasma, with the mean plasma exposure (C_{max} and AUC) and renal clearance at the steady-state comparable to those observed with a single dose. The plasma concentrations 1 hour after dosing were relatively stable for 280 days.

3.3.2. Distribution

In vitro (^{14}C)-lenalidomide binding to plasma proteins was low with mean plasma protein binding at 22.7% and 29.2% in MM patients and healthy volunteers, respectively.

Lenalidomide is present in semen (< 0.01% of the dose) after administration of 25 mg/day and the drug is undetectable in semen 3 days after stopping the drug.

3.3.3. Metabolism and Excretion

In vitro studies indicate that lenalidomide has no inhibitory effect on CYP1A2, CYP2C9, CYP2C19, CYP2D6, CYP2E1 and CYP3A.

A majority of lenalidomide is eliminated unchanged through urinary excretion. The contribution of renal excretion to total clearance in subjects with normal renal function was 65 - 85%. The half-life ($t_{½}$) of elimination has been observed to increase with dose, from approximately 3 hours at 5 mg up to approximately 9 hours at doses of 400 mg (the higher dose is believed to provide a better estimate of $t_{½}$). Steady-state levels are achieved by Day 4.

Following a single oral administration of [^{14}C]-lenalidomide (25 mg) to healthy volunteers, approximately 90% and 4% of the radioactive dose is eliminated in urine and faeces, respectively. Approximately 82% of the radioactive dose is excreted as lenalidomide, almost exclusively via the urinary route. Hydroxy-lenalidomide and N-acetyl-lenalidomide represent 4.59% and 1.83% of the excreted dose, respectively. The renal clearance of lenalidomide exceeds the glomerular filtration rate and therefore is at least actively secreted to some extent.

Pharmacokinetic analyses in patients with impaired renal function indicate that as renal function decreases (< 50 mL/min), the total drug clearance decreases proportionally resulting in an increase in AUC. The half-life of lenalidomide increased from approximately 3.5 hours in subjects with creatinine clearance > 50 mL/min to more than 9 hours in subjects with reduced renal function (< 50 mL/min). However, renal impairment did not alter the oral absorption of lenalidomide. The C_{max} was similar between healthy subjects and patients with renal impairment. Recommended dose adjustments in patients with impaired renal function are described in Section 10 [Dosage and administration].

Pharmacokinetic analyses based on MM studies indicate that lenalidomide is rapidly absorbed at all dose levels, with maximum plasma concentrations occurring between 0.5 and 4.0 hours post-dose both on Days 1 and 28. The C_{max} and AUC values increase proportionally with dose following single and multiple doses in MM patients. Exposure in MM patients is slightly higher based on C_{max} and AUC values as compared to healthy male volunteers since the clearance/bioavailable fraction of a drug (CL/F) in MM patients is lower (approximately 200 mL/min compared to 300 mL/min) than it is in healthy volunteers. This is consistent with the compromised renal function in the MM patients, possibly as a consequence of their age (average patient age of 58 vs. 29 for healthy volunteers) and their disease.

4. Clinical Trials

4.1. Newly Diagnosed Multiple Myeloma (NDMM) in Patients Not Eligible for Stem-cell Transplantation

Study MM-020 was a Phase III, multicenter, randomised, open-label, 3-arm study to compare the efficacy and safety of lenalidomide and low-dose dexamethasone (Rd) given for 2 different durations of time (i.e., [Arm A: Continuous Rd, until progressive disease] or [Arm B: Rd18, for up to eighteen 28-day cycles {72 weeks}]), to Arm C (melphalan, prednisone and thalidomide [MPT] for a maximum of twelve 42-day cycles [72 weeks]). A total of 1623 subjects with newly diagnosed multiple myeloma (NDMM) (non-eligible for autologous stem cell transplant [ASCT]) were enrolled and randomised in a 1:1:1 ratio to Arm A (n = 535), Arm B (n = 541), or Arm C (n = 547).

Patients in the Continuous Rd arm and the Rd18 arm received lenalidomide 25 mg once daily on Days 1 to 21 of 28-day cycles. Dexamethasone 40 mg was dosed once daily on Days 1, 8, 15, and 22 of each 28-day cycle. Initial dose and regimen for the Continuous Rd and Rd18 arms were adjusted according to age and renal function. Patients > 75 years received a dexamethasone dose of 20 mg once daily on Days 1, 8, 15, and 22 of each 28-day cycle. All patients received prophylactic anticoagulation (low molecular weight heparin, warfarin, heparin, or low-dose aspirin) during the study.

The primary efficacy endpoint in the study was progression free survival (PFS). The demographics and disease-related baseline characteristics of the patients were well balanced in all 3 arms. In general, study subjects had advanced-stage disease: of the total study population, 41% had ISS stage III, and 9% had severe renal insufficiency (creatinine clearance [CLcr] < 30 mL/min). The median age was 73 in the 3 arms, with 35% of total patients > 75 years of age.

The study showed a statistically significant prolongation of PFS benefit in patients receiving Continuous Rd (Arm A) compared to MPT (Arm C). The Hazard Ratio was 0.72 ([95% CI: 0.61, 0.85]; p = 0.00006), indicating a 28% decrease in the risk of disease progression for patients treated with Continuous Rd compared with those treated with MPT. The median follow-up time for all surviving subjects at the interim analysis was 37.0 months. The overall response rate (≥ partial response [PR]) was higher in Continuous Rd (75.1%) than in MPT (62.3%) (p < 0.00001). A greater percentage of patients achieved at least a complete response (CR) in Continuous Rd than in MPT (15.1% versus 9.3 %, respectively).

The preliminary analysis of the primary comparison of overall survival (OS) shows a reduction of risk of death of 22% in the Continuous Rd Arm compared with the MPT arm. In an updated analysis of OS where the median follow-up time for all surviving subjects was 45.5 months, a further improvement in the reduction of risk of death was noted in the Continuous Rd arm compared with the MPT arm (HR 0.75; p = 0.002). The efficacy results are summarised in Table 1 below.

PFS2 (an exploratory endpoint) was defined for all patients as the time from randomisation to second objective progressive disease (PD), or death from any cause, whichever occurred first. PFS2 was significantly longer in the Continuous Rd arm compared to arm MPT (HR 0.77 [95% CI: 0.65, 0.92]; p = 0.003). The results show a difference in patients who have started 2nd line treatment and type of therapy received: for Continuous Rd, of the 43% who started 2nd line treatment, 62% received bortezomib-containing therapy vs. 12% lenalidomide therapy; for MPT, of the 57% who started 2nd line therapy, 49% received bortezomib-containing therapy vs. 34% lenalidomide therapy.

Table 1. Summary of Efficacy Data from Study MM-020

Endpoint	Continuous Rd (N = 535)	Rd18 (N = 541)	MPT (N = 547)
PFS (months)			
Median [95% CI]	25.5 [20.7, 29.4]	20.7 [19.4, 22.0]	21.2 [19.3, 23.2]
HR [95% CI]; p-value			
Rd vs. MPT	0.72 [0.61, 0.85]; p = 0.00006		
Rd vs. Rd18	0.70 [0.60, 0.82]; p = 0.00001		
Rd18 vs. MPT	1.03 [0.89, 1.20]; ns		
Overall Survival (months)*			
Median [95% CI]	58.9 [56.0, NE]	56.7 [50.1, NE]	48.5 [44.2, 52.0]
HR [95% CI]; p-value			
Rd vs. MPT	0.75 [0.62, 0.90]; p = 0.002		
Rd vs. Rd18	0.91 [0.75, 1.09]; ns		
Rd18 vs. MPT	0.83 [0.69, 0.99]; p = 0.034		
Myeloma Response, n (%)			
Complete Response	81 (15.1)	77 (14.2)	51 (9.3)
Very Good Partial Response	152 (28.4)	154 (28.5)	103 (18.8)
Partial Response	169 (31.6)	166 (30.7)	187 (34.2)
Overall response (CP, VGPR or PR)	402 (75.1)	397 (73.4)	341 (62.3)
Duration of response (months)			
Median [95% CI]	35.0 [27.9, 43.4]	22.1 [20.3, 24.0]	22.3 [20.2, 24.9]

* OS data is based on an updated analysis (03 March 2014); NE = not estimable; ns = not significant

4.2. Previously Treated Multiple Myeloma

The efficacy and safety of lenalidomide were evaluated in two Phase III, multi-centre, randomised, double-blind, placebo-controlled, parallel-group controlled studies (MM-009 and MM-010) of lenalidomide plus high dose dexamethasone therapy versus high dose dexamethasone alone in patients with MM who have received at least one prior treatment. Out of 353 patients in the MM-009 and MM-010 studies who received lenalidomide/dexamethasone, 45.6% were aged 65 or over. Of the 704 patients evaluated in the MM-009 and MM-010 studies, 44.6% were aged 65 or over.

In both studies, patients in the lenalidomide/dexamethasone (len/dex) group took 25 mg of lenalidomide orally once daily on Days 1 to 21 and a matching placebo capsule once daily on Days 22 to 28 of each 28-day cycle. Patients in the placebo/dexamethasone (placebo/dex) group took 1 placebo capsule on Days 1 to 28 of each 28-day cycle. Patients in both treatment groups took 40 mg of dexamethasone orally once daily on Days 1 to 4, 9 to 12, and 17 to 20 of each 28-day cycle for the first 4 cycles of therapy. The dose of dexamethasone was reduced to 40 mg orally once daily on Days 1 to 4 of each 28-day cycle after the first 4 cycles of therapy. In both studies, treatment was to continue until disease progression. In both studies, dose adjustments were allowed based on clinical and laboratory finding.

The primary efficacy endpoint in both studies was time-to-progression (TTP). In total, 353 patients were evaluated in the MM-009 study; 177 in the len/dex group and 176 in the placebo/dex group and, in total, 351 patients were evaluated in the MM-010 study; 176 in the len/dex group and 175 in the placebo/dex group.

In both studies, the baseline demographic and disease-related characteristics were comparable between the len/dex and placebo/dex groups. Both patient populations presented a median age of 63 years, with a comparable male to female ratio. The ECOG performance status was comparable between both groups, as was the number and type of prior therapies.

Pre-planned interim analyses of both studies showed that len/dex was statistically significantly superior ($p < 0.00001$) to dexamethasone alone for the primary efficacy endpoint, TTP. CR and overall response (OR) rates in the len/dex arm were also significantly higher than the placebo/dex arm in both studies. An extended follow-up efficacy analysis was conducted with a median follow-up of 30.2 months. Table 2 summarises the results of the follow-up efficacy analyses.

Table 2: Summary of Efficacy Analysis - Studies MM-009 and MM-010

Endpoint	MM-009		MM-010	
	Len/Dex N = 177	Placebo/Dex N = 176	Len/Dex N = 176	Placebo/Dex N = 175
TTP (months)				
Median [95% CI]	13.9 [9.5, 17.1]	4.6 [3.7, 5.1]	12.1 [9.4, 19.8]	4.6 [3.8, 4.8]
Hazard ratio [95% CI]	0.33 [0.25, 0.44]		0.36 [0.27, 0.48]	
p-value	< 0.001		< 0.001	
Response Rate				
CR n (%)	28 (16)	4 (2)	30 (17)	7 (4)
PR n (%)	79 (45)	30 (17)	75 (43)	34 (19)
p-value	< 0.001		< 0.001	
PFS (months)				
Median [95% CI]	12.3 [8.4, 16.7]	4.6 [3.7, 4.7]	10.2 [7.4, 15.2]	4.6 [3.7, 4.7]
Hazard ratio [95% CI]	0.36 [0.27, 0.47]		0.42 [0.32, 0.55]	
p-value	< 0.001		<0.001	

4.3. Myelodysplastic Syndromes (MDS)

The efficacy and safety of Revlimid were evaluated in low- or intermediate-1-risk MDS patients with a deletion-5q (q31-33) cytogenetic abnormality, with or without additional cytogenetic abnormalities.

Study MDS-004 was a Phase III, multi-centre, randomised, double-blind, placebo-controlled study in red blood cell (RBC) transfusion-dependent subjects. The 52-week double-blind treatment phase included 205 subjects who were randomised to receive 10 mg lenalidomide for 21 days of a 28-day cycle, 5 mg lenalidomide continuously, or placebo. The primary efficacy endpoint was transfusion independence at 182 days. The median age of patients was 68.0 years (range 36 to 86), the median duration of MDS was 2.6 years (range 0.2 to 29.2) and 76.1% of patients were females. The study enrolled patients with absolute neutrophil counts (ANC) $\geq 0.5 \times 10^9$/L, platelet counts $\geq 25 \times 10^9$/L, serum creatinine ≤ 2.0 mg/dL, serum SGOT/AST or SGPT/ALT ≤ 3.0 x upper limit of normal (ULN), and serum total bilirubin ≤ 1.5 mg/dL. An overview of the efficacy results for the Intent-to-Treat (ITT) populations from MDS-004 receiving either cyclic lenalidomide dosing at 10 mg, or placebo, is presented in Table 3.

Study MDS-003 was a Phase II, multi-centre, open-label, single-arm study of 148 patients who were RBC transfusion-dependent. Dosing was primarily at a continuous dose of 10 mg once daily for 28 days, with some experience at a dose of 10 mg daily for 21 of 28 days. The primary efficacy endpoint was RBC transfusion independence of at least 2 months duration, as defined by the MDS International Working Group (IWG) criteria. The median age of patients was 71.0 years (range 37 to 95), the median duration of MDS was 2.5 years (range 0.1 to 20.7) and 65.5% of patients were females. The study enrolled patients with absolute neutrophil counts (ANC) $\geq 0.5 \times 10^9$/L, platelet counts $\geq 50 \times 10^9$/L, serum creatinine ≤ 2.5 mg/dL, serum SGOT/AST or SGPT/ALT ≤ 3.0 x upper limit of normal (ULN), and serum direct bilirubin ≤ 2.0 mg/dL. Table 3 summarises the efficacy results for the ITT population from MDS-003.

In both MDS-003 and MDS-004, granulocyte colony-stimulating factor was permitted for patients who developed neutropenia or fever in association with neutropenia.

Table 3. Summary of Results of Efficacy Analyses for Studies MDS-003 and MDS-004

Endpoint	MDS-003	MDS-004*	
	10 mg Cont	10 mg Cyclic	Placebo
Number RBC-Transfusion Independent at 56 days [a]	97 (65.5%) N=148	42 (60.9%) N=69	5 (7.5%) N=67
Number RBC-Transfusion Independent at 182 days [b]	86 (58.1%) N=148	38 (55.1%) N=69	4 (6.0%) N=67
Median time (range) to transfusion independence (weeks) [c]	4.1 (0.3, 49.0) N=97	4.6 (0.3, 14.7) N=42	0.3 (0.3, 24.1) N=5
Median [95% CI] duration of RBC-transfusion independence (weeks)	114.4 [78.4 – 153.7] N=97	NE [98.3 – NE] N=42	NE [9.1 – NE] N=5
Durability of response – subjects who maintained transfusion independence [d]	40 (41.2%) N=97	30 (71.4%) N=42	4 (80.0%) N=5
Median rise in haemoglobin (g/dL) (range)	5.6 (2.2, 40.7) N=97	6.4 (1.8, 10.0) N=42	2.6 (1.5, 4.4) N=5

Cont = continuous (28 days of a 28-day cycle); Cyclic = (21 days of a 28-day cycle)

*: Based on RBC-transfusion independence response for subjects who became RBC-transfusion independent for at least 56 days.

a: transfusion independence was defined as the absence of any RBC transfusion during any consecutive 56 days during the treatment period accompanied by at least a 1 g/dL increase in Hb from screening/baseline.

b: RBC-transfusion independence response for subjects who became RBC-transfusion independent for at least 182 days.
c: Measured from the day of the first dose of study drug to the first day of the first 56-day RBC transfusion-free period.
d: Measured from the first of the consecutive 56 days during which the subject was free of RBC transfusions to the date of the first RBC transfusion after this period.

5. Indications

5.1. Multiple Myeloma (MM)

Revlimid is indicated for the treatment of patients with newly diagnosed multiple myeloma who are ineligible for autologous stem cell transplantation.

Revlimid in combination with dexamethasone is indicated for the treatment of multiple myeloma patients whose disease has progressed after one therapy.

5.2. Myelodysplastic Syndromes (MDS)

Revlimid is indicated for treatment of patients with transfusion-dependent anaemia due to low- or intermediate-1 risk myelodysplastic syndromes associated with a deletion 5q cytogenetic abnormality with or without additional cytogenetic abnormalities.

6. Contraindications

- Women who are pregnant.
- Women of childbearing potential unless all of the conditions of the *i-access*® Program are met (see Section 7 [Precautions]).
- Hypersensitivity to the active substance or to any of the excipients.

7. Precautions

7.1. Effects on Fertility

A fertility and early embryonic development study in male and female rats, with administration of lenalidomide up to 500 mg/kg/day, produced no parental toxicity and no adverse effects on fertility or early embryonic development. The systemic exposure in rats at 500 mg/kg was > 70-fold higher than the human exposure at 25 mg/day, based on AUC.

7.2. Use in Pregnancy (Pregnancy Category X)

For lenalidomide, no clinical data on exposed pregnancies are available. Because lenalidomide is a structural analogue of thalidomide (a known human teratogen that causes severe, life-threatening birth defects), and has shown teratogenic effects in animal studies, lenalidomide must not be used in pregnant women. Women of childbearing potential must use effective means of contraception.

Embryofoetal development studies were conducted in monkeys and rabbits. In monkeys, lenalidomide was teratogenic at systemic exposures (based on plasma AUC) well below that anticipated clinically, and a NOEL for the teratogenic effects could not be established in the study.

In rabbits treated with 3, 10 and 20 mg/kg/day orally, maternal and developmental toxicity was noted at ≥ 10 mg/kg/day. The toxicity was characterised by slightly reduced foetal body weights, increased incidences of post-implantation loss, and gross external findings in the foetuses associated with maternal toxicity of lenalidomide. Increased incidences of soft tissue and skeletal variations in the foetuses were also observed at 10 and 20 mg/kg/day. The NOEL for developmental toxicity of lenalidomide in rabbits was identified as 3 mg/kg/day, which is associated with a plasma AUC value equivalent to that anticipated clinically at the 25 mg/day dose in humans.

If lenalidomide is taken during pregnancy, a teratogenic effect of lenalidomide in humans cannot be ruled out. Therefore, the conditions of the *i-access*® Program must be fulfilled for all patients.

7.2.1. The *i-access*® Program Conditions for Pregnancy Prevention

Revlimid is available under a restricted distribution program (*i-access*®). Only physicians and pharmacists registered with this Program can prescribe and dispense the product. In addition, Revlimid must only be dispensed to patients who are registered and meet all the conditions of the Program.

7.2.1.1. Females of Non-Childbearing Potential

A female patient or a female partner of a male patient is considered to have childbearing potential unless she meets at least one of the following criteria:

- Age ³ 50 years and naturally amenorrhoeic for ³ 1 year*
- Premature ovarian failure confirmed by a specialist gynaecologist
- Previous bilateral salpingo-oophorectomy, or hysterectomy
- XY genotype, Turner syndrome, uterine agenesis.

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

Female patients of non-childbearing potential are only required to comply with the General Conditions listed within the *i-access*® Program (see Section 7.2.1.5).

7.2.1.2. Females of Childbearing Potential

Female patients of childbearing potential must comply with the following requirements on counselling, contraception and pregnancy testing.

If pregnancy occurs in a female patient treated with lenalidomide, treatment must be stopped and the patient should be referred to a physician specialised or experienced in teratology for evaluation and advice. Similarly, if pregnancy occurs in a partner of a male patient taking lenalidomide, the female partner should be referred to a physician specialised or experienced in teratology for evaluation and advice.

Counselling

For female patients of childbearing potential, lenalidomide is contraindicated unless all of the following are met:

- She understands the potential teratogenic risk to the unborn child.
- She understands and agrees to comply with the need for effective contraception, without interruption, 4 weeks before starting treatment, throughout the entire duration of treatment, and 4 weeks after the end of treatment.
- Even if a woman of childbearing potential has amenorrhea she must follow all the advice on effective contraception.
- She should be capable of complying with effective contraceptive measures.
- She is informed and understands the potential consequences of pregnancy and the need to rapidly consult her physician if there is a risk of pregnancy.
- She understands the need to commence the treatment as soon as lenalidomide is dispensed following a negative pregnancy test.
- She understands the need and accepts to undergo medically supervised pregnancy testing every 4 weeks.
- She acknowledges that she understands the hazards and necessary precautions associated with the use of lenalidomide.

Contraception

Female patients of childbearing potential must use one effective method of contraception for 4 weeks before therapy, during therapy, and until 4 weeks after lenalidomide therapy, even in case of dose interruption, unless the patient commits to absolute and continuous abstinence confirmed on a monthly basis. If not established on effective contraception, the patient must be referred to an appropriately trained health care professional for contraceptive advice in order that contraception can be initiated.

Table 4. Recommended Methods of Contraception

Contraceptive method	Comments
Contraceptive implant	Contraceptive implants and levonorgestrel-releasing intrauterine systems are associated with an increased risk of infection at the time of insertion and irregular vaginal bleeding. Prophylactic antibiotics should be considered particularly in patients with neutropenia. Copper-releasing intrauterine devices are generally not recommended due to the potential risks of infection at the time of insertion and menstrual blood loss which may compromise patients with neutropenia or thrombocytopenia.
Levonorgestrel-releasing intrauterine system (IUS)	
Medroxyprogesterone acetate depot	
Tubal ligation	
Sexual intercourse with a vasectomised male partner only	Vasectomy must be confirmed by two negative semen analyses.
Ovulation inhibitory progesterone-only pills (i.e. desogestrel).	Because of the increased risk of venous thromboembolism in patients with MM taking lenalidomide and dexamethasone, combined oral contraceptive pills are not recommended. If a patient is currently using combined oral contraception, the patient should switch to one of the effective methods listed above. The risk of venous thromboembolism continues for 4-6 weeks after discontinuing combined oral contraception. The efficacy of contraceptive steroids may be reduced during co-treatment with dexamethasone.

Pregnancy Testing

Medically supervised pregnancy tests with a minimum sensitivity of 25 mIU/mL must be performed for females of childbearing potential as outlined below.

This requirement includes females of childbearing potential who practice absolute and continuous abstinence. Ideally, pregnancy testing, issuing a prescription and dispensing should occur on the same day. For females of childbearing potential, dispensing of lenalidomide must occur within a maximum of 7 days of the negative pregnancy test.

· *Prior to Starting Treatment*

A medically supervised pregnancy test should be performed when lenalidomide is prescribed. The test should occur either at the time of consultation, or in the 3 days prior to the visit to the prescriber and at a point where the patient has been using effective contraception for at least 4 weeks. The test should ensure the patient is not pregnant when she starts treatment with lenalidomide. This requirement includes women of childbearing potential who practice absolute and continuous abstinence.

· *Follow-up and End of Treatment*

A medically supervised pregnancy test should be repeated every 4 weeks, including 4 weeks after the end of treatment. These pregnancy tests should be performed on the day of the prescribing visit or in the 3 days prior to the visit to the prescriber. This requirement includes women of childbearing potential who practice absolute and continuous abstinence.

7.2.1.3. Male Patients

Male patients must comply with the following requirements on counselling and contraception as clinical data has demonstrated the presence of lenalidomide in human semen.

Counselling and contraception

- He understands the potential teratogenic risk if engaged in sexual activity with a female of childbearing potential.
- He understands and complies with the need for the use of a condom (if engaged in sexual activity with a female of childbearing potential) throughout treatment duration, during dose interruption, and for 1 week after cessation of treatment if their partner is of childbearing potential and has no contraception.
- He understands that if his partner becomes pregnant whilst he is taking lenalidomide or during the 1st week after he discontinues taking lenalidomide, he should inform his treating physician immediately.
- He understands that he must not donate sperm during therapy (including during dose interruptions) or for 1 week following discontinuation of lenalidomide.

7.2.1.4. Prescribers

- Ensure that females patients of childbearing potential comply with the conditions of the *i-access*® Program, including confirmation that they have an adequate level of understanding of the Program requirements
- Provide full patient information about the potential teratogenic risk and the strict pregnancy prevention measures as specified in the *i-access*® Program to female patients of childbearing potential and, as appropriate, to male patients
- Ensure that all patients acknowledge and agreed to comply with the conditions of the *i-access*® Program.

7.2.1.5. General Conditions

All patients should be instructed never to give this medicinal product to another person and to return any unused capsules to their pharmacist at the end of treatment.

All patients should not donate blood during therapy (including during dose interruptions), or for 1 week following discontinuation of lenalidomide. In Australia, patients with myeloma are permanently excluded from donating blood.

7.3. Use during Lactation

It is not known whether lenalidomide is excreted in human milk. Because of the potential for adverse reactions in nursing infants from lenalidomide, a decision should be made whether to discontinue nursing or to discontinue the medicine, taking into account the importance of the drug to the mother.

7.4. Paediatric Use

There is no experience in treating children and adolescents with Revlimid. Therefore, lenalidomide should not be used in the paediatric age group (0-18 years).

7.5. Use in the Elderly

Population pharmacokinetic analyses included patients with ages ranging from 39 to 85 years old and show that age does not influence the disposition of lenalidomide. No dose adjustments are needed for lenalidomide.

For NDMM patients > 75 years of age, a reduced starting dose of dexamethasone is recommended (see Section 10 [Dosage and administration]).

Lenalidomide has been used in clinical trials in previously treated MM patients up to 86 years of age (see Section 3 [Pharmacology]). The percentage of patients aged 65 or over was not significantly different between the len/dex and placebo/dex groups. No overall difference in effectiveness was observed between these patients and younger patients. However, overall serious adverse events, in particular the serious vascular events (including DVT and pulmonary embolism) and serious cardiovascular events (including atrial fibrillation), were all more frequent in lenalidomide-treated patients 65 years and over.

Lenalidomide has also been used in MDS clinical trials in patients up to 95 years of age. Of the 395 patients in the MDS clinical trials who received 10 mg lenalidomide, 72.2% were aged 65 and over. No overall difference in safety was observed between these patients and younger patients, but greater pre-disposition of older individuals to drug-related toxicities cannot be ruled out.

Lenalidomide is known to be substantially excreted by the kidney. The risk of adverse reactions to this drug may be greater in patients with impaired renal function. Elderly patients are more likely to have decreased renal function, so care should be taken in dose selection for such patients. Renal function should therefore be monitored (see Section 10 [Dosage and administration]).

7.6. Genotoxicity/Carcinogenicity

In vitro (mutation in bacteria, chromosomal aberration in human lymphocytes, mutation in mouse lymphoma cells, Syrian Hamster Embryo cell transformation) and *in vivo* (rat micronucleus) genotoxicity studies revealed no drug related-effects at either the gene or chromosomal level. Carcinogenicity studies with lenalidomide have not been conducted.

7.7. Second Primary Malignancies

In clinical trials in NDMM patients not eligible for ASCT, a 4.9-fold increase in incidence rate of haematologic second primary malignancies (SPM) (cases of AML and MDS) has been observed in patients receiving lenalidomide in combination with melphalan and prednisone until progression (1.75 per 100 person-years) compared with melphalan in combination with prednisone (0.36 per 100 person-years). A 2.12-fold increase in incidence rate of solid tumour SPM has been observed in patients receiving lenalidomide (9 cycles) in combination with melphalan and prednisone (1.57 per 100 person-years) compared with melphalan in combination with prednisone (0.74 per 100 person-years).

In NDMM patients receiving lenalidomide in combination with dexamethasone until progression or for 18 months, the haematologic SPM incidence rate (0.16 per 100 person-years) was not increased as compared to thalidomide in combination with melphalan and prednisone (0.79 per 100 person-years). A 1.3-fold increase in incidence rate of solid tumour SPM has been observed in patients receiving lenalidomide in combination with dexamethasone until progression or for 18 months (1.58 per 100 person-years) compared to thalidomide in combination with melphalan and prednisone (1.19 per 100 person-years).

In clinical trials of NDMM patients eligible for transplant, an increased incidence rate of haematologic SPM has been observed in patients receiving lenalidomide immediately following high-dose melphalan and autologous stem cell transplant (ASCT) (1.27 to 1.56 per 100 patient-years) compared with patients who received placebo (0.46 to 0.53 per 100 person-years, respectively). Cases of B-cell malignancies (including Hodgkin's lymphoma) observed in the clinical trials were in patients who received

lenalidomide in the post-ASCT setting. Please note that lenalidomide is not indicated for the treatment of NDMM patients eligible for ASCT.

Based on a low number of cases, a numerical imbalance in SPM (comprising mainly of basal cell and squamous cell skin cancers) has been observed in clinical trials in previously treated MM patients with len/dex (3.98 per 100 patient-years) compared with placebo/dex (1.38 per 100 patient-years).

Subjects who received lenalidomide-containing therapy until disease progression did not show a higher incidence of invasive SPM than subjects treated in the fixed duration lenalidomide-containing arms. These results suggest that duration of lenalidomide treatment is not associated with an increased risk for the occurrence of invasive SPM.

Both the benefit achieved with Revlimid and the risk of SPM should be considered and discussed with patients, before initiating treatment with the product. Physicians should also carefully evaluate patients before and during treatment using standard cancer screening for occurrence of SPM and institute treatment as appropriate.

7.8. Myocardial Infarction

Myocardial infarction has been reported in patients receiving lenalidomide, particularly in those with known risk factors. Patients with known risk factors (including prior thrombosis) should be closely monitored, and action should be taken to try to minimise all modifiable risk factors (e.g. smoking, hypertension, and hyperlipidaemia).

7.9. Venous and Arterial Thromboembolism

The combination of lenalidomide with dexamethasone is associated with an increased risk of venous thromboembolism (deep vein thrombosis [DVT] and pulmonary embolism [PE]) and arterial thromboembolism (predominantly myocardial infarction and cerebrovascular event), in patients with MM. In clinical studies of patients with del 5q MDS, lenalidomide as monotherapy was also associated with an increased risk of DVT. Concomitant administration of erythropoietic agents or previous history of DVT may also increase thrombotic risk in these patients. Therefore, erythropoietic agents, or other agents that may increase the risk of thrombosis, such as hormone replacement therapy, should be used with caution in MM patients receiving lenalidomide with dexamethasone. A haemoglobin concentration above 120 g/L should lead to discontinuation of erythropoietic agents. Patients and physicians are advised to be observant for the signs and symptoms of thromboembolism. Patients should be instructed to seek medical care if they develop symptoms such as shortness of breath, chest pain, arm or leg swelling. Prophylactic antithrombotic medicines, such as low molecular weight heparins or warfarin, should be recommended, especially in patients with additional thrombotic risk factors. The decision to take antithrombotic prophylactic measures should be made after careful assessment of an individual patient's underlying risk factors.

If a patient experiences any thromboembolic events, treatment must be discontinued and standard anticoagulation therapy started. Once the patient has been stabilised on the anticoagulation treatment and any complications of the thromboembolic event have been managed, the lenalidomide treatment may be restarted at the original dose dependent upon a benefit-risk assessment. The patient should continue anticoagulation therapy during the course of lenalidomide treatment.

7.10. Neutropenia and Thrombocytopenia

The major dose-limiting toxicities of lenalidomide include neutropenia and thrombocytopenia. Patients with neutropenia should be monitored for signs of infection. Patients and physicians are advised to be observant for signs and symptoms of bleeding, including petechiae and epistaxis, especially with use of concomitant medication that may increase risk of bleeding. Therefore, co-administration of lenalidomide with other myelosuppressive agents should be undertaken with caution. Appropriate management should be instituted if such toxicity is observed. Patients taking Revlimid should have their

complete blood counts (CBC) assessed periodically as described below. A dose interruption and/or dose reductions may be required.

7.10.1. Newly Diagnosed Multiple Myeloma (NDMM) in Patients Not Eligible for ASCT

CBC should be assessed every 7 days (weekly) for the first 2 cycles, day 1 and day15 of cycle 3, and every 28 days (4 weeks) thereafter.

7.10.2. Previously Treated Multiple Myeloma

The combination of lenalidomide with dexamethasone in previously treated MM patients is associated with a higher incidence of Grade 4 neutropenia (4.8% in len/dex-treated patients compared with 0.6% in placebo/dex-treated patients; see Section 9 [Adverse Effects]). Grade 4 febrile neutropenia episodes were observed infrequently (0.6% in len/dex-treated patients compared to 0.0% in placebo/dex treated patients; see Section 9 [Adverse Effects]). Patients should be advised to promptly report febrile episodes. In case of neutropenia, the physician should consider the use of growth factors in patient management.

The combination of lenalidomide with dexamethasone in previously treated MM patients is associated with a higher incidence of Grade 3 and Grade 4 thrombocytopenia (10.8% and 1.4%, respectively, in len/dex-treated patients compared to 5.4% and 0.9% in placebo/dexamethasone-treated patients; see Section 4 [Clinical trials]).

CBC, including white blood cell count with differential count, platelet count, haemoglobin, and haematocrit should be performed at baseline, every 2 weeks for the first 12 weeks of lenalidomide treatment, and monthly thereafter to monitor for cytopenias.

7.10.3. Myelodysplastic Syndrome (MDS)

In clinical studies of patients with del 5q MDS, lenalidomide as monotherapy was associated with significant neutropenia and thrombocytopenia. Grade 3 or 4 haematologic toxicity was seen in 80% of patients. In the 48% of patients who developed Grade 3 or 4 neutropenia, the median time to onset was 42 days (range, 14-411 days), and the median time to documented recovery was 17 days (range, 2-170 days). In the 54% of patients who developed Grade 3 or 4 thrombocytopenia, the median time to onset was 28 days (range, 8-290 days), and the median time to documented recovery was 22 days (range, 5-224 days).

Patients on therapy for del 5q MDS should have their CBCs monitored weekly for the first 8 weeks of therapy and at least monthly thereafter. Patients may require the use of blood product support and/or growth factors (see Section 10 [Dosage and administration]).

7.11. Peripheral Neuropathy

Lenalidomide is structurally related to thalidomide, which is known to induce severe peripheral neuropathy. At this time, the neurotoxic potential of lenalidomide associated with long-term use cannot be ruled out.

7.12. Tumour Lysis Syndrome and Tumour Flare Reaction

Tumour lysis syndrome (TLS) and tumour flare reaction (TFR) have commonly been observed in patients with chronic lymphocytic leukemia (CLL), and uncommonly in patients with other lymphomas, who were treated with lenalidomide. Fatal instances of TLS have been reported during treatment with lenalidomide. Patients at risk of TLS and TFR are those with high tumour burden prior to treatment. Caution should be practiced when introducing these patients to lenalidomide. These patients should be monitored closely, especially during the first cycle or dose-escalation, and appropriate precautions taken. There have been rare reports of TLS in patients with MM treated with lenalidomide, and no reports in patients with MDS treated with lenalidomide.

7.13. Angioedema and Other Dermatological Reactions

Rare cases of angioedema and serious dermatological reactions including Stevens-Johnson syndrome and toxic epidermal necrolysis have been reported from post-marketing experience. These events have the potential to be fatal. Patients with a prior history of Grade 4 rash associated with thalidomide treatment should not receive lenalidomide. Revlimid interruption or discontinuation should be considered for Grade 2-3 skin rash. Revlimid must be discontinued for angioedema, Grade 4 rash, exfoliative or bullous rash, or if Stevens-Johnson syndrome or toxic epidermal necrolysis is suspected. Revlimid should not be resumed following the discontinuation for these reactions.

7.14. Atrial Fibrillation

In the two pivotal randomised controlled trials in previously treated (relapsed/refractory) MM patients, atrial fibrillation occurred in 14 (4.0%) subjects treated with len/dex compared to 4 (1.1%) subjects treated with placebo/dex (unadjusted for the longer on-study observation time for patients receiving lenalidomide). Careful review of these cases revealed the presence of multiple risk factors for atrial fibrillation (e.g. infections, hypertension, congestive heart failure, electrolyte imbalance), and a causal relationship to lenalidomide treatment has not yet been determined.

7.15. Use in Patients with Impaired Thyroid Function

Cases of hypothyroidism have been reported and monitoring of thyroid function should be considered.

7.16. Use in Patients with Lactose Intolerance

Revlimid capsules contain lactose. Patients with rare hereditary problems of galactose intolerance, the Lapp lactase deficiency or glucose-galactose malabsorption should not take Revlimid.

7.17. Hepatic Disorders

Hepatic failure, including fatal cases, has been reported in patients treated with lenalidomide in combination with dexamethasone: acute hepatic failure, toxic hepatitis, cytolytic hepatitis, cholestatic hepatitis, and mixed cytolytic/cholestatic hepatitis have been reported. The mechanisms of severe drug-induced hepatotoxicity remain unknown although, in some cases, pre-existing viral liver disease, elevated baseline liver enzymes, and possibly treatment with antibiotics might be risk factors.

Abnormal liver function tests were commonly reported and were generally asymptomatic and reversible upon dosing interruption. Once parameters have returned to baseline, treatment at a lower dose may be considered.

Lenalidomide is excreted by the kidneys. It is important to dose adjust patients with renal impairment in order to avoid plasma levels which may increase the risk for higher haematological side effects or hepatotoxicity. Monitoring of liver function is recommended, particularly when there is a history of or concurrent viral liver infection, or when lenalidomide is combined with medications known to be associated with liver dysfunction.

7.18. Increased Mortality in Chronic Lymphocytic Leukemia

In a prospective, randomized (1:1) clinical trial in the first line treatment of patients with chronic lymphocytic leukemia (CLL), single agent Revlimid therapy was associated with an increased risk of death as compared to single agent chlorambucil. Lenalidomide is not recommended for use in CLL outside of controlled clinical trials.

7.19. Effects on Ability to Drive and Use Machines

No studies on the effects on the ability to drive and use machines have been performed. Lenalidomide may have minor or moderate influence on the ability to drive and use machines. Fatigue, dizziness, somnolence and blurred vision have been reported with the use of lenalidomide. Therefore, caution is recommended when driving or operating machines.

8. Interaction with Other Medicines

Erythropoietic agents, or other agents that may increase the risk of thrombosis, such as hormone replacement therapy, should be used with caution in MM patients receiving lenalidomide with dexamethasone (see Section 7 [Precautions], and Section 9 [Adverse Effects]).

In vitro, lenalidomide does not inhibit UGT1A1-mediated bilirubin glucuronidation in human liver microsomes derived from donors representing genotypes UGT1A1*1/*1, UGT1A1*1/*28, and UGT1A1*28/*28.

The major dose-limiting toxicities of lenalidomide include neutropenia and thrombocytopenia. Therefore, co-administration of lenalidomide with other myelosuppressive agents should be undertaken with caution.

8.1. Oral Contraceptives

No interaction study has been performed with oral contraceptives. Lenalidomide is not an inducer of cytochrome P450 enzymes (see below). Dexamethasone is known to be a weak to moderate inducer of CYP3A4 and is likely to also affect other enzymes as well as transporters. It may not be excluded that the efficacy of oral contraceptives may be reduced during treatment. Effective measures to avoid pregnancy must be taken.

Lenalidomide is not a substrate, inhibitor or inducer of cytochrome P450 enzymes *in vitro*. Hence, co-administration of cytochrome P450 substrates (including hormonal contraceptives), inhibitors or inducers with lenalidomide is not likely to result in clinically relevant drug-drug interactions.

8.2. Dexamethasone

In patients with MM, co-administration of single or multiple doses of dexamethasone (40 mg/day) had no significant effect on the multiple dose pharmacokinetics of lenalidomide (25 mg/day).

8.3. Warfarin

Co-administration of multiple doses of 10 mg of lenalidomide had no effect on the single dose pharmacokinetics of R- and S- warfarin. Co-administration of a single 25 mg dose of warfarin had no effect on the pharmacokinetics of lenalidomide. However, it is not known whether there is an interaction during clinical use (concomitant treatment with dexamethasone). Dexamethasone is a weak to moderate enzyme inducer and its effect on warfarin is unknown. Close monitoring of warfarin concentration is advised during the treatment.

8.4. Digoxin

Concomitant administration with lenalidomide 10 mg/day increased the plasma exposure of digoxin (0.5 mg, single dose) by 14%. Therefore, periodic monitoring of the digoxin concentration is advised during lenalidomide treatment. In the same study, the co-administration of digoxin (a P-glycoprotein substrate) did not significantly affect the pharmacokinetics of lenalidomide.

8.5. Human Efflux Transporters

Lenalidomide is a weak substrate but not an inhibitor of P-glycoprotein (P-gp). Co-administration of multiple doses of P-gp inhibitor, quinidine (600 mg, twice daily) had no effect on the single dose pharmacokinetics of lenalidomide (25 mg). Single dose co-administration of lenalidomide (25 mg) and P-gp inhibitor/substrate, temsirolimus (25 mg), does not affect the pharmacokinetics of either drug.

Lenalidomide is not an inhibitor of bile salt export pump (BSEP), MRP2, OAT1, OAT3, OATP1B1, OATP1B3, or OCT2.

8.6. Renal Drug Interactions

Renal drug-drug interaction studies have not been performed. The renal clearance of lenalidomide is slightly greater than the glomerular filtration rate, suggesting that active secretion contributes to a minor extent (≤ 25%) of renal clearance. Hence, the inhibition of the active secretion of lenalidomide will most likely not result in a clinically relevant drug-drug interaction.

9. Adverse Effects

9.1. Multiple Myeloma (MM)

9.1.1. Newly Diagnosed Multiple Myeloma (NDMM) in Patients Not Eligible for ASCT

In the large phase III, controlled study (MM-020), data were evaluated from 1072 patients who received at least one dose of Revlimid with low dose dexamethasone, either given Continuously (Rd) or for 18 cycles (Rd18) (see Section 4 [Clinical trials]).Median treatment duration was notably longer in the Continuous Rd arm (80.1 weeks) when compared to the Rd18 (72 weeks) and the MPT arms (67.1 weeks), as the Continuous Rd arm sustained treatment until disease progression, while treatments in arms Rd18 and MPT were both capped at 72 weeks. The median average daily dose of lenalidomide was 21.8 mg in the Continuous Rd arm and 24.4 mg in the Rd18 arm.

In general, the most frequently reported adverse events (AEs) were comparable in Arm Rd until progression and Arm Rd18, and included diarrhoea, anaemia, constipation, peripheral oedema, neutropenia, fatigue, back pain, nausea, asthenia, insomnia, decreased appetite and muscle spasms. The most frequently reported Grade 3 or 4 events included neutropenia, anaemia, thrombocytopenia, pneumonia, asthenia, fatigue, back pain, hypokalaemia, rash, cataract, lymphopenia, dyspnoea, DVT, hyperglycaemia, and leukopenia. No particular AE led to discontinuation of any study drug in more than 2% of subjects in either arm. Over time, the Rd regimen was generally better tolerated than MPT. Subjects in Arm MPT discontinued treatment sooner and more frequently prior to disease progression than subjects receiving Rd. Subjects in Arm MPT also more frequently experienced AEs leading to study drug discontinuation. Extended treatment in the Rd arm beyond 18 months generally resulted in a limited increase in most AEs compared with Rd18 or MPT.

For the Rd regimen, 66.4% of patients experienced at least one AE leading to Revlimid interruption, 60.0% experienced at least one AE leading to dexamethasone interruption, and 69.2% experienced at least one AE leading to Revlimid or dexamethasone interruption, compared to 77.4% in the MPT arm for thalidomide or melphalan or prednisone interruption.

9.1.2. Previously Treated Multiple Myeloma

In two Phase III placebo-controlled studies (MM-009 and MM-010), 353 patients with previously treated MM were exposed to the len/dex combination and 352 to the placebo/dex combination. The median duration of exposure to study treatment was significantly longer (44.0 weeks) in the len/dex group as compared to placebo/dex (23.1 weeks). The difference was accounted for by a lower rate of discontinuation from study treatment due to lower progression of disease in patients exposed to len/dex (39.7%) than in placebo/dex patients (70.4%).

The most serious adverse events were:

- Venous thromboembolism (deep vein thrombosis, pulmonary embolism) (see Section 7 [Precautions])
- Grade 4 neutropenia (see Section 7 [Precautions]).

Table 5 below collectively shows the treatment-emergent adverse events that occurred at a frequency of greater than or equal to 10% in any of the study arms for the MM-009, MM-010, and MM-020 studies in MM patients:

Table 5.Treatment-emergent Adverse Events Reported for at least 10% of Subjects in Any Arm – Studies MM-009/010 and MM-020

System Organ Class Preferred term	% frequency				
	MM-009/010		MM-020		
	RD	Pbo	Continuous[a] Rd	Rd18[b]	MPT
Number of patients (N)	**353**	**350**	**532**	**540**	**541**
Blood and Lymphatic System Disorders					
Anaemia	36.0	25.1	43.8	35.7	42.3
Neutropenia	44.8	8.3	35.0	33.0	60.6*
Thrombocytopenia	22.9	12.0	19.5	18.5	25.0*
Leukopenia	9.9	4.0	11.8	11.1	17.4*
Lymphopenia	5.9	2.6	11.1	8.0	13.1*
Ear and Labyrinth Disorders					
Vertigo	4.2	2.3	5.1	3.7	6.5*
Eye disorders					
Cataract	8.2	2.9	13.7	5.7	0.9
Vision blurred	18.4	12.3	5.5	3.7	4.4
Gastrointestinal Disorders					
Diarrhoea	45.3	29.7	45.5	38.5	16.5
Constipation	41.9	21.7	43.0	39.3	52.7*
Nausea	27.8	21.7	28.6	23.7	30.5*
Vomiting	13.3	9.7	17.5	12.6	20.1*
Abdominal pain	11.0	6.6	13.0	7.6	5.5
Abdominal pain upper	9.3	6.0	8.5	6.9	5.4
Dyspepsia	17.0	14.6	10.7	5.2	6.7
Dry mouth	7.6	3.7	7.0	7.0	11.5*
General Disorders and Administration Site Conditions					
Oedema peripheral	32.3	24.9	39.7	31.3	39.7*
Fatigue	47.0	42.0	32.5	32.8	28.5
Asthenia	30.6	26.9	28.2	22.8	22.9
Pyrexia	29.5	24.0	21.4	18.9	14.0
Oedema	10.8	9.1	7.1	5.2	5.9
Infections and Infestations					
Bronchitis	16.1	9.1	16.9	10.9	7.9
Nasopharyngitis	22.1	9.7	15.0	10.0	6.1
Urinary tract infection	9.9	5.7	14.3	11.7	7.6
Upper respiratory tract infection	26.1	15.7	13.0	9.8	5.7
Pneumonia	16.7	8.6	12.4	12.6	7.4
Pharyngitis	16.7	9.7	1.3	1.5	1.5
Investigations					
Weight decreased	19.8	15.4	13.5	14.4	8.9
Blood creatinine increased	4.2	3.4	4.6	6.6	4.4
Metabolism and Nutrition Disorders					
Decreased appetite	22.7	14.0	23.1	21.3	13.3
Hypokalaemia	15.9	6.6	17.1	11.5	7.0
Hyperglycaemia	16.7	14.3	11.7	9.6	3.5
Hypocalcaemia	9.9	3.1	10.7	10.4	5.7
Musculoskeletal and Connective Tissue Disorders					
Back pain	28.6	19.4	32.0	26.9	21.4
Muscle spasms	38.2	23.1	20.5	18.9	11.3

System Organ Class Preferred term	% frequency				
	MM-009/010		MM-020		
	RD	Pbo	Continuous[a] Rd	Rd18[b]	MPT
Number of patients (N)	**353**	**350**	**532**	**540**	**541**
Arthralgia	20.7	18.3	19.0	13.1	12.2
Bone pain	17.0	11.7	16.4	14.3	11.5
Pain in extremity	15.9	11.1	14.8	12.2	11.3
Musculoskeletal pain	3.4	2.9	12.6	10.9	6.7
Musculoskeletal chest pain	8.8	6.9	11.3	9.4	7.2
Muscular weakness	16.7	16.3	8.1	6.5	5.4
Myalgia	11.3	11.1	5.1	3.5	3.1
Nervous System Disorders					
Peripheral sensory neuropathy	4.2	3.1	20.5	17.0	35.3*
Paraesthesia	15.0	13.4	16.0	13.7	19.0*
Dizziness	24.9	16.9	15.8	13.0	21.1*
Headache	27.2	25.4	14.1	9.6	10.4
Tremor	21.2	7.7	14.1	13.5	18.5*
Hypoaesthesia	11.6	7.7	8.3	4.4	7.6
Dysgeusia	15.3	10.0	7.3	8.3	4.1
Neuropathy peripheral	15.3	10.9	6.4	4.1	11.5*
Psychiatric Disorders					
Insomnia	38.0	37.7	27.6	23.5	9.8
Depression	13.6	10.9	10.9	8.5	5.5
Anxiety	12.5	9.7	7.7	6.7	7.6
Confusional state	10.8	6.9	7.1	5.4	4.6
Respiratory, Thoracic and Mediastinal Disorders					
Cough	26.9	25.4	22.7	17.4	12.6
Dyspnoea	25.8	18.0	22.0	16.5	20.9
Skin and Subcutaneous Tissue Disorders					
Rash	23.8	11.1	21.4	24.3	17.2
Pruritus	8.8	5.4	8.8	9.1	4.4
Hyperhidrosis	10.2	7.4	4.7	3.5	2.4
Vascular Disorders					
Deep vein thrombosis	9.3	4.6	10.2	6.7	3.7

a = Continuous Rd arm where patients were dosed with Revlimid + low-dose dexamethasone until progressive disease
b = Rd18 arm where patients were dosed with Revlimid + low-dose dexamethasone for up to eighteen 28-day cycles (72 weeks)

* Events where frequency in the comparator arm was the same or higher than the treatment arm(s) in the same study. If more than one treatment arm was present, the highest frequency in any of the treatment arms was used for the comparison.

9.2. Myelodysplastic Syndromes (MDS)

Data from the placebo-controlled MDS-004 study demonstrate that lenalidomide is also well tolerated in subjects with low- or intermediate-1-risk MDS with a deletion 5q cytogenetic abnormality with or without other cytogenetic abnormalities. The most frequently reported adverse events were related to blood and lymphatic system disorders, skin and subcutaneous tissue disorders, gastrointestinal disorders, and general disorders and administrative site conditions.

In study MDS-004, neutropenia in 76.8% (106/138) of subjects and thrombocytopenia in 46.4% (64/138) of subjects were the most frequently reported adverse events. The next most common adverse events observed were diarrhoea (34.8%), constipation (19.6%) and nausea (19.6%); pruritus (25.4%) and rash (18.1%); fatigue (18.1%) and oedema peripheral (15.2%); and muscle spasms (16.7%). Table 6

summarises the adverse events that were reported in ≥ 10% of the Revlimid-treated patients and more frequent than in the placebo patients, in the MDS-004 clinical study.

Table 6. Most Frequently Reported (≥ 10% in lenalidomide arm) Adverse Events in MDS-004

System Organ Class Preferred term	% with lenalidomide[a] (N=138)	% with placebo (N=67)
General Disorders & Administration Site Conditions		
Fatigue	18.1	7.5
Oedema Peripheral	15.2	7.5
Pyrexia	13.8	6.0
Gastrointestinal Disorders		
Diarrhoea	34.8	17.9
Nausea	19.6	9.0
Constipation	19.6	7.5
Abdominal pain	10.9	6.0
Musculoskeletal and Connective Tissue Disorders		
Muscle spasms	16.7	9.0
Nervous System disorders		
Headache	14.5	9.0
Respiratory, Thoracic and Mediastinal Disorders		
Cough	11.6	6.0
Infections and Infestations		
Nasopharyngitis	11.6	7.5
Bronchitis	11.6	4.5
Upper respiratory tract infection	10.9	6.0
Skin and Subcutaneous Tissue Disorders		
Pruritus	25.4	4.5
Rash	18.1	1.5
Dry Skin	10.1	1.5
Blood and Lymphatic System Disorders		
Neutropenia	76.8	17.9
Thrombocytopenia	46.4	3.0
Leukopenia	12.3	4.5

a. Combined 5 mg and 10 mg lenalidomide treatment arms from MDS-004.

The safety results (N=148) from the Phase 2 open-label study MDS-003 are consistent with the findings from MDS-004. Neutropenia (66.2%) and thrombocytopenia (64.9%) were the most frequently reported AEs, followed by diarrhoea (60.1%), pruritus (44.6%), fatigue (42.6%), rash (37.8%) and arthralgia (31.8%).

The most serious Grade 3 and Grade 4 adverse events from the MDS-004 study (N=138, 5 mg and 10 mg doses combined) were neutropenia (5.8%), thrombocytopenia (5.8%), venous thromboembolism (deep vein thrombosis [3.6%] and pulmonary embolism [2.9%]), and altered mood (0.7%). The frequency of these events in the open-label MDS-003 study (N=148) were neutropenia (64.9%), thrombocytopenia (54.7%), and venous thromboembolism (deep vein thrombosis [4.7%] and pulmonary embolism [3.4%]).

In the 10 mg group from study MDS-004, the dose of Revlimid was reduced or interrupted at least once due to an AE in 43 (62.3%) patients, which occurred a mean of 50.1 days into the study and lasted a mean of 26.8 days. Twenty-four (34.8%) subjects had a second dose reduction or interruption. In study MDS-003, the dose of Revlimid was reduced or interrupted at least once due to an AE in 127 (85.8%)

patients, which occurred a mean of 75.2 days into the study and lasted a mean of 30.4 days. Eighty-two (55.4%) subjects had a second dose reduction or interruption. The mean interval between the first and second dose reduction/interruption was 198.2 days. The second dose reduction/interruption due to an AE lasted a mean of 44.5 days.

9.3. Adverse Drug Reactions – Multiple Myeloma (MM) and Myelodysplastic Syndromes (MDS)

The adverse drug reactions observed in MM patients, and in MDS patients treated with at least one dose of 10 mg lenalidomide, are tabulated below by system organ class and frequency (Table 7). Frequencies are defined as: very common (≥ 1/10); common (≥ 1/100, < 1/10); uncommon (≥ 1/1,000, < 1/100).

The following table is derived from data gathered during the main clinical trials in newly diagnosed/previously treated MM and MDS. The data were not adjusted according to differences in duration of treatment across the MM studies.

Adverse drug reactions have been included under the appropriate category in the table below according to the highest frequency observed in the lenalidomide arm of any of the main clinical trials.

Table 7: Adverse Drug Reactions (including Grade 3 and 4) observed in Patients in Main MM and MDS Studies*

Frequency	All ADRs	Grade 3 and 4 ADRs
Infections and Infestations[#]		
Very common	Pneumonia; Bronchitis; Bacterial, viral and fungal infections (including opportunistic infections); Upper respiratory tract infection	Pneumonia
Common	Sepsis; Sinusitis	Bacterial, viral and fungal infections (including opportunistic infections); Sepsis; Bronchitis
Neoplasms benign, malignant and unspecified (including cysts and polyps)		
Common		Squamous cell carcinoma of skin
Blood and Lymphatic System Disorders		
Very common	Neutropenias; Thrombocytopenia; Anaemia, Leukopenias	Neutropenias; Thrombocytopenia; Anaemia; Leukopenias
Common	Febrile neutropenia; Pancytopenia	Febrile neutropenia; Pancytopenia
Endocrine Disorders		
Common	Hypothyroidism	
Metabolism and Nutrition Disorders		
Very common	Decreased appetite; Hypokalaemia; Hyperglycaemia; Hypocalcaemia	
Common	Dehydration; Hypomagnesaemia; Iron overload	Hypokalaemia; Hypocalcaemia; Hypophosphataemia; Diabetes mellitus; Hyperglycaemia; Hyponatraemia; Gout; Decreased appetite
Psychiatric Disorders		
Very common	Insomnia; Depression	
Common		Depression; Insomnia
Nervous System Disorders		
Very common	Peripheral neuropathies (excluding motor neuropathy); Dizziness; Tremor; Dysgeusia; Headache	
Common	Lethargy	Syncope; Dizziness; Peripheral

Frequency	All ADRs	Grade 3 and 4 ADRs
		neuropathy; Cerebrovascular accident
Eye Disorders		
Very common	Cataracts; Blurred vision	
Common		Cataracts
Cardiac Disorders		
Common	Atrial fibrillation	Myocardial infarction (including acute); Atrial fibrillation; Tachycardia; Cardiac failure (including congestive)
Vascular Disorders		
Very common	Venous thromboembolic events (predominantly deep vein thrombosis and pulmonary embolism)	Venous thromboembolic events (predominantly deep vein thrombosis and pulmonary embolism)
Common	Hypertension; Hypotension; Haematoma	Hypotension
Respiratory, Thoracic and Mediastinal Disorders		
Very common	Dyspnoea; Epistaxis	
Common		Respiratory distress; Dyspnoea
Gastrointestinal Disorders		
Very common	Diarrhoea; Vomiting; Nausea; Constipation; Abdominal pain; Dyspepsia	
Common	Dry mouth	Diarrhoea; Nausea; Constipation; Toothache
Hepatobiliary Disorders		
Common	Abnormal liver function tests	Cholestasis; Abnormal liver function tests
Skin and Subcutaneous Tissue Disorders		
Very common	Rash[+]; Pruritus, Dry skin; Hyperhidrosis	
Common	Erythema	Rash[+]; Pruritus
Musculoskeletal and Connective Tissue Disorders		
Very common	Musculoskeletal and connective tissue pain and discomfort; Bone pain; Muscle spasms; Arthralgia; Myalgia	
Common	Muscular weakness	Muscular weakness; Musculoskeletal and connective tissue pain and discomfort; Bone pain
Renal Disorders		
Very common	Renal failure (including acute)	
Common		Renal failure
General Disorders and Administration Site Conditions		
Very common	Pyrexia; Oedema (including peripheral); Asthenia; Influenza-like illness syndrome (including pyrexia, cough, rhinitis, myalgia, musculoskeletal pain, headache and rigours); Fatigue	
Common	Chest Pain	Pyrexia; Fatigue; Asthenia
Injury, Poisoning and Procedural Complications		
Common	Fall; Contusion	Fall
Investigations		
Very common	Weight decreased	
Common		Weight decreased

- All PTs under SOC of Infections except for rare infections of Public Health interest will be considered listed.
+ - All PTs under HLT of Rash will be considered listed.
* Algorithm applied for determination of ADRs from clinical trials:

- Phase III studies - MM-009, MM-010, MM-020 and MDS-004 (double-blind safety population)
  - All treatment-emergent adverse events with ≥ 5.0% of subjects in the lenalidomide-containing arm(s) and ≥ 2.0% higher frequency (%) in lenalidomide-containing arm(s) compared to the non-lenalidomide arm
  - All treatment-emergent Grade 3 or 4 adverse events in ≥ 1.0% of subjects in lenalidomide-containing arm(s) and ≥ 1.0% higher frequency (%) in lenalidomide-containing arm(s) compared to the non-lenalidomide arm, occurring in 2 or more subjects unless medically significant
- MDS Phase II study (MDS-003)
  - All treatment-emergent adverse events with ≥ 5.0% of lenalidomide-treated subjects
  - All treatment-emergent Grade 3 or 4 adverse\events in ≥ 1% of lenalidomide-treated subjects
  - If a term met the algorithm for inclusion in Study MDS-004, the highest frequency of the term was used from either Study MDS-004 or MDS-003.

9.4. Post-Marketing Experience

The following adverse reactions have been identified during post-marketing use of Revlimid. Because these reactions are reported voluntarily, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure: Allergic conditions (angioedema, Stevens-Johnson syndrome, and toxic epidermal necrolysis), tumour lysis syndrome (TLS) and tumour flare reaction (TFR), pneumonitis, myocardial infarction, hypothyroidism, hyperthyroidism, transient abnormal liver laboratory tests, hepatic failure, acute hepatic failure, hepatitis toxic, cytolytic hepatitis, cholestatic hepatitis, and mixed cytolytic/cholestatic hepatitis.

In the literature and post-marketing setting, acute graft-versus-host disease has been reported following allogeneic hematopoietic transplant.

9.4.1. Hepatic Disorders

Cases of transient liver laboratory abnormalities (predominantly transaminases) were reported in patients treated with lenalidomide. For such patients, treatment with lenalidomide should be interrupted and restarted once the levels return to baseline. Successful re-challenge with lenalidomide, without recurrence of elevated liver laboratory results, was reported in some patients.

9.4.2. Thyroid Function

Cases of hypothyroidism and hyperthyroidism have been reported. Optimal control of co-morbid conditions is recommended before start of treatment. Baseline and ongoing monitoring of thyroid function is recommended.

10. Dosage and Administration

Treatment must be initiated and monitored under the supervision of a registered Specialist Physician experienced in the management of haematological and oncological malignancies.

10.1. Administration

Revlimid capsules should be taken at about the same time each day. The capsules should not be opened, broken or chewed. The capsules should be swallowed whole, preferably with water and either one hour before or two hours after food. If less than 12 hours have elapsed since missing a dose, the patient can take the dose. If more than 12 hours have elapsed since missing a dose at the normal time, the patient should not take the dose, but take the next dose at the normal time on the following day.

10.2. Haematological Testing

For NDMM patients, CBC should be assessed every 7 days (weekly) for the first 2 cycles, day 1 and day15 of cycle 3, and every 28 days (4 weeks) thereafter.

For patients with previously treated MM, CBC, including white blood cell count with differential count, platelet count, haemoglobin, and haematocrit should be performed at baseline, every week for the first 8 weeks of lenalidomide treatment and monthly thereafter to monitor for cytopenias.

Patients on therapy for del 5q MDS should have their CBCs monitored weekly for the first 8 weeks of therapy and at least monthly thereafter.

10.3. Recommended Dosage

10.3.1. Multiple Myeloma

10.3.1.1. Newly Diagnosed Multiple Myeloma in Patients Not Eligible for ASCT

The recommended starting dose of lenalidomide is 25 mg orally once daily on days 1-21 of repeated 28-day cycles. The recommended dose of low dose dexamethasone is 40 mg orally once daily on days 1, 8, 15 and 22 of repeated 28-day cycles. Patients may continue lenalidomide and dexamethasone therapy until disease progression or intolerance. Dosing is continued or modified based upon clinical and laboratory findings (see Section 7 [Precautions]).

For elderly patients (i.e. > 75 years of age) with NDMM treated with lenalidomide in combination with dexamethasone, the starting dose of dexamethasone is 20 mg/day on Days 1, 8, 15 and 22 of each 28-day treatment cycle.

Lenalidomide treatment in combination with dexamethasone must not be started if the Absolute Neutrophil Count (ANC) < 1.5 x 10^9/L, and platelet count < 50 x 10^9/L.

Recommended dose adjustments for NDMM patients receiving len/dex are found in Section 10.4.1.1.

10.3.1.2. Previously Treated Multiple Myeloma

The recommended starting dose of lenalidomide is 25 mg orally once daily on days 1-21 of repeated 28-day cycles. The recommended dose of dexamethasone is 40 mg orally once daily on days 1-4, 9-12, and 17-20 of each 28-day cycle for the first 4 cycles of therapy and then 40 mg once daily on days 1-4 every 28 days. Treatment should be continued until disease progression or unacceptable toxicity. Dosing is continued or modified based upon clinical and laboratory findings (see Section 7 [Precautions]).

Lenalidomide treatment must not be started if the ANC < 1.0 x 10^9/L, and/or platelet counts < 75 x 10^9/L or, dependent on bone marrow infiltration by plasma cells, platelet counts < 30 x 10^9/L.

Recommended dose adjustments for previously treated MM patients are found in Section 10.4.1.2.

10.3.2. Myelodysplastic Syndrome (MDS)

The recommended starting dose of lenalidomide is 10 mg given orally once a day on days 1 to 21 of repeating 28-day treatment cycles. Dosing is continued or modified based upon clinical and laboratory findings.

Lenalidomide treatment must not be started if the ANC < 0.5 x 10^9/L, and/or platelet counts < 50 x 10^9/L.

Recommended dose adjustments for MDS patients are found in Section 10.4.2.

10.4. Recommended Dose Adjustments During Treatment and Re-initiation of Treatment

Dose adjustments, as summarised below, are recommended to manage Grade 3 or 4 neutropenia or thrombocytopenia, or other Grade 3 or 4 toxicities judged to be related to lenalidomide.

10.4.1. Multiple Myeloma

10.4.1.1. Newly Diagnosed Multiple Myeloma in Patients Not Eligible for ASCT

· *Dose Reduction Levels*

	Lenalidomide	Dexamethasone
Starting dose	25 mg	40 mg
Dose Level 1	20 mg	20 mg
Dose Level 2	15 mg	12 mg
Dose Level 3	10 mg	8 mg
Dose Level 4	5 mg	4 mg
Dose Level 5	2.5 mg	NA

· *Dose Reduction Guidance*

Thrombocytopenia	
When platelets	**Recommended lenalidomide Course**
First fall to < 25 x 10^9/L	Stop lenalidomide dosing for remainder of cycle[a]
Return to ≥ 50 x 10^9/L	Decrease by one dose level when dosing is resumed at next cycle. Do not dose below 2.5 mg daily

Neutropenia[b]	
When neutrophils:	**Recommended lenalidomide Course**
First fall to < 0.5 x 10^9/L or < 1.0 x 10^9/L associated with fever (temperature ≥ 38.5°C)	Interrupt lenalidomide treatment
Return to 1.0 x 10^9/L when neutropenia is the only observed toxicity	Resume lenalidomide at Starting Dose
Return to ≥ 0.5 x 10^9/l when dose-dependent haematological toxicities other than neutropenia are observed	Resume lenalidomide at Dose Level 1 once daily
For each subsequent drop below < 0.5 x 10^9/L Return to ≥ 0.5 x 10^9/L	Interrupt lenalidomide treatment Resume at next lower dose level once daily. Do not dose below 2.5 mg daily

a If dose-limiting toxicity occurs on > Day15 of a cycle, lenalidomide dosing will be interrupted for at least the remainder of the current 28-day cycle.

b At the physician's discretion, if neutropenia is the only toxicity at any dose level, treat the patient with G-CSF and maintain the dose level of lenalidomide.

If the dose of lenalidomide was reduced for a haematologic dose limiting toxicity (DLT), the dose of lenalidomide may be re-increased to the next higher dose level (up to the starting dose) at the discretion of the treating physician if continued lenalidomide / dexamethasone therapy resulted in improved bone marrow function (no DLT for at least 2 consecutive cycles and an ANC ≥1.5 x 10^9/L with a platelet count ≥ 100 x 10^9/L at the beginning of a new cycle at the current dose level).

10.4.1.2. Previously Treated Multiple Myeloma

· *Dose Reduction Levels*

Starting dose	25 mg
Dose Level 1	15 mg
Dose Level 2	10 mg
Dose Level 3	5 mg

· *Dose Reduction Guidance*

Thrombocytopenia	
When platelets	**Recommended lenalidomide Course**
First fall to < 30 x 10^9/L	Interrupt lenalidomide treatment
Return to ≥ 30 x 10^9/L	Resume lenalidomide at Dose Level 1
For each subsequent drop below < 30 x 10^9/L Return to ≥ 30 x 10^9/L	Interrupt lenalidomide treatment Resume at next lower dose level once daily.

Neutropenia[a]	
When neutrophils:	**Recommended lenalidomide Course**
First fall to < 0.5 x 10^9/L	Interrupt lenalidomide treatment
Return to 0.5 x 10^9/L when neutropenia is the only observed toxicity	Resume lenalidomide at Starting Dose
Return to ≥ 0.5 x 10^9/l when dose-dependent haematological toxicities other than neutropenia are observed	Resume lenalidomide at Dose Level 1 once daily
For each subsequent drop below < 0.5 x 10^9/L Return to ≥ 0.5 x 10^9/L	Interrupt lenalidomide treatment Resume at next lower dose level once daily. Do not dose below 5 mg once daily.

a In case of neutropenia, the physician should consider the use of growth factors in patient management.

10.4.2. Myelodysplastic Syndromes (MDS)

· *Dose Reduction Guidance*

For patients with MDS, dose reduction guidelines are divided into 2 sets - for within the first 4 weeks of treatment, and after the first 4 weeks of treatment.

i). For patients who experience thrombocytopenia or neutropenia within the first 4 weeks of treatment:

Thrombocytopenia		
When baseline	**When platelets**	**Recommended Course**
Platelet count ≥ 100 x 10^9/L	Fall to < 50 x 10^9/L	Interrupt lenalidomide treatment
	Return to ≥ 50 x 10^9/L	Resume lenalidomide at 5 mg/day
Platelet count ≥ 60 x 10^9 and < 100 x 10^9/L	Fall by 50% of the baseline value	Interrupt lenalidomide treatment
	Return to ≥ 50 x 10^9/L	Resume lenalidomide at 5 mg/day
Platelet count < 60 x 10^9/L	Fall by 50% of the baseline value	Interrupt lenalidomide treatment
	Return to ≥ 30 x 10^9/L	Resume lenalidomide at 5 mg/day

Neutropenia		
When baseline	**When neutrophils**	**Recommended Course**
ANC ≥ 1 x 10^9/L	Fall to < 0.75 x 10^9/L	Interrupt lenalidomide treatment
	Return to ≥ 1 x 10^9/L	Resume lenalidomide at 5 mg/day
ANC < 1 x 10^9/L	Fall to < 0.5 x 10^9/L	Interrupt lenalidomide treatment
	Return to ≥ 0.5 x 10^9/L	Resume lenalidomide at 5 mg/day

ii). For patients who experience thrombocytopenia after the first 4 weeks of treatment:

Thrombocytopenia	
During treatment at 10 mg/day:	
When platelets	**Recommended Course**
Fall to < 30 x 10^9/L or < 50 x 10^9/L with platelet transfusions	Interrupt lenalidomide treatment
Return to ≥ 30 x 10^9/L (without haemostatic failure)	Resume lenalidomide at 5 mg/day
During treatment at 5 mg/day:	
Fall to < 30 x 10^9/L or < 50 x 10^9/L with platelet transfusions	Interrupt lenalidomide treatment
Return to ≥ 30 x 10^9/L (without haemostatic failure)	Resume lenalidomide at 5 mg/day every other day

Neutropenia	
During treatment at 10 mg/day:	
When neutrophils	**Recommended Course**
Fall to < 0.5 x 10^9/L for ≥ 7 days or to < 0.5 x 10^9/L associated with fever (temperature ≥ 38.5°C)	Interrupt lenalidomide treatment
Return to ≥ 0.5 x 10^9/L	Resume lenalidomide at 5 mg/day
During treatment at 5 mg/day:	
When neutrophils	**Recommended Course**
Fall to < 0.5 x 10^9/L for ≥ 7 days or to < 0.5 x 10^9/L associated with fever (temperature ≥ 38.5°C)	Interrupt lenalidomide treatment
Return to ≥ 0.5 x 10^9/L	Resume lenalidomide at 5 mg every other day

10.4.3. Other Dose Adjustments for MM and MDS

Grade 3/4 Toxicities

For other Grade 3/4 toxicities judged to be related to lenalidomide, stop treatment and restart at next lower dose level when toxicity has resolved to ≤ Grade 2 at the physician's discretion.

Discontinuation of Revlimid

Revlimid interruption or discontinuation should be considered for Grade 2-3 skin rash. Revlimid must be discontinued for angioedema, Grade 4 rash, exfoliative or bullous rash, or if Stevens-Johnson syndrome or toxic epidermal necrolysis is suspected. Revlimid should not be resumed following the discontinuation for these reactions.

10.4.4. Use in Patients with Impaired Renal Function

Lenalidomide is substantially excreted by the kidney. With patients with impaired renal function, care should be taken in dose selection. Monitoring of renal function is advised in patients with renal impairment.

No dose adjustments are required for patients with mild renal impairment. The following dose adjustments are recommended at the start of therapy for patients with moderate or severe impaired renal function, or end stage renal disease.

Renal Function (CLcr)	**Dose Adjustment**	
	MM	**MDS**
Moderate renal impairment (30 £ CLcr < 60 mL/min)	10 mg once daily*	5 mg once daily
Severe renal impairment (CLcr < 30 mL/min, not requiring dialysis)	15 mg every other day	5 mg every other day
End Stage Renal Disease (ESRD) (CLcr < 30 mL/min, requiring dialysis)	5 mg once daily On dialysis days, the dose should be administered following dialysis	5 mg, 3 times a week following each dialysis

* In MM, the dose may be escalated to 15 mg once daily after 2 cycles if patient is not responding to treatment and is tolerating the treatment. CLcr = creatinine clearance.

Monitoring of patients with impaired renal function for signs and symptoms of neutropenia or thrombocytopenia should be done on a weekly basis for the first 8 weeks after the initiation of lenalidomide therapy.

10.4.5. Use in Patients with Impaired Hepatic Function:

Revlimid has not formally been studied in patients with impaired hepatic function and there are no specific dose recommendations.

For further information regarding Revlimid's compatibility with other drugs and monitoring advice, please refer to Section 7 [Precautions].

11. Overdosage

There is no specific experience in the management of lenalidomide overdose in patients with MM or MDS. In dose-ranging studies, healthy subjects were exposed to up to 200 mg (administered 100 mg twice daily) and in single-dose studies, some were exposed to up to 400 mg. Pruritus, urticaria, rash, and elevated liver transaminases were the primary reported adverse events. No clinically significant changes in ECGs, blood pressure, or pulse rate were observed.

While no haematologic events were associated with an overdose, such events may be expected since in clinical trials, the dose limiting toxicity was essentially haematological. In the event of overdose, supportive care is advised. In Australia, contact the Poisons Advisory Centre on 13 11 26 for advice on management. In New Zealand, contact the National Poison Centre on 0800 POISON or 0800 764 766 for advice on management.

12. Presentation and Storage Conditions

12.1. Presentation

Revlimid is available as 2.5 mg, 5 mg, 7.5 mg, 10 mg, 15 mg, 20 mg and 25 mg hard capsules in blister packs containing 21 capsules each. However, not all strengths are being distributed in Australia.

Revlimid 2.5 mg capsules: White/blue-green size 4 capsules marked "2.5 mg REV". Each 2.5 mg capsule contains 2.5 mg lenalidomide. Blister packs containing 21 capsules.

Revlimid 5 mg capsules: White size 2 capsules marked "5 mg REV". Each 5 mg capsule contains 5 mg lenalidomide. Blister packs containing 21 capsules.

Revlimid 7.5 mg capsules: White/pale-yellow size 2 capsules marked "7.5 mg REV". Each 7.5 mg capsule contains 7.5 mg lenalidomide. Blister packs containing 21 capsules.

Revlimid 10 mg capsules: Pale yellow/blue-green size 0 capsules marked "10 mg REV". Each 10 mg capsule contains 10 mg lenalidomide. Blister packs containing 21 capsules.

Revlimid 15 mg capsules: White/powder-blue size 0 capsules marked "15 mg REV". Each 15 mg capsule contains 15 mg lenalidomide. Blister packs containing 21 capsules.

Revlimid 20 mg capsules: Powder-blue/blue-green size 0 capsules marked "20 mg REV". Each 20 mg capsule contains 20 mg lenalidomide. Blister packs containing 21 capsules.

Revlimid 25 mg capsules: White size 0 capsules marked "25 mg REV". Each 25 mg capsule contains 25 mg lenalidomide. Blister packs containing 21 capsules.

12.2. Composition

Active

Lenalidomide

Excipients

See Section 2 [Description] for a list of the excipients.

12.3. Storage Conditions

Store below 25°C. Store in the original package.

12.4. Container Type

Polychlorotrifluoroethylene (PCTFE) / polyvinylchloride (PVC) / Aluminium foil blisters.

13. Name and Address of the Sponsor

Celgene Pty Limited

Level 7, 607 St Kilda Road,
Melbourne, VIC 3004,
Australia.

14. Poison Schedule of the Medicine

Schedule 4 (Prescription Only Medicine)

15. Date of First Inclusion in the Australian Register of Therapeutic Goods (the ARTG)

20 December 2007

16. Date of Most Recent Amendment

11 November 2015