

Information on this form is a guide only. User will be solely responsible for verifying its currency and accuracy with the corresponding BC Cancer treatment protocols located at <a href="https://www.bccancer.bc.ca">www.bccancer.bc.ca</a> and according to acceptable standards of care

## PROTOCOL CODE: UMYLDREL

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Patient RevAid ID:\_\_\_\_

REMINDER: Please ensure drug allergies and previous bleomycin are documented on the Allergy & Alert Form  Risk Category: Female of Childbearing Potential (FCBP) Risk Category: Male or Female of nonChildbearing Potential (NCBP)  START DATE OF THIS CYCLE Cycle # & Cycle # START DATE OF SUBSEQUENT CYCLES Cycle # &	Pharmacy Use for Lenalidomide dispensing:  Part Fill # 1  RevAid confirmation number:  Lenalidomide lot number:  Pharmacist counsel (initial):  Part Fill # 2  RevAid confirmation number:
Allergy & Alert Form  Risk Category:	Part Fill # 1 RevAid confirmation number:  Lenalidomide lot number:  Pharmacist counsel (initial):  Part Fill # 2
Risk Category:	RevAid confirmation number:  Lenalidomide lot number:  Pharmacist counsel (initial):  Part Fill # 2
Risk Category:	RevAid confirmation number:  Lenalidomide lot number:  Pharmacist counsel (initial):  Part Fill # 2
START DATE OF THIS CYCLE  START DATE OF SUBSEQUENT CYCLES  Delay treatment week(s)  May proceed with doses as written if within 7 days  ANC greater than or equal to 1.0 x 10°/L, Platelets greater than or equal to 30 x 10°/L and eGFR as per protocol  Dose modification for:  Hematology Renal Function Other Toxicity  OR Proceed with treatment based on blood work from  LENALIDOMIDE  One cycle = 28 days	Lenalidomide lot number:  Pharmacist counsel (initial):  Part Fill # 2
□ Delay treatment week(s)  May proceed with doses as written if within 7 days  ANC greater than or equal to 1.0 x 10°/L, Platelets greater than or equal to 30 x 10°/L and eGFR as per protocol  Dose modification for: □ Hematology □ Renal Function □ Other Toxicity  OR Proceed with treatment based on blood work from  LENALIDOMIDE  One cycle = 28 days	Pharmacist counsel (initial):  Part Fill # 2
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10 <sup>9</sup> /L and eGFR as per protocol Dose modification for: Hematology Renal Function Other Toxicity  OR Proceed with treatment based on blood work from  LENALIDOMIDE One cycle = 28 days	Part Fill # 2
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OR Proceed with treatment based on blood work from  LENALIDOMIDE One cycle = 28 days	
LENALIDOMIDE One cycle = 28 days	
One cycle = 28 days	RevAid confirmation number:
□ levelide maide* may DO deily, in the examination of device 1 to 24 and off for 7	
☐ lenalidomide*mg PO daily, in the evening, on days 1 to 21 and off for 7	
days	Lenalidomide lot number:
☐ lenalidomide* mg PO	
(*available as 25 mg, 20 mg, 15 mg, 10 mg, 5 mg, 2.5 mg capsules)	
*Note: Use one capsule strength for the total dose; there are cost implications as	Pharmacist counsel (initial):
costing is per capsule and not weight based	
FCBP dispense 21 capsules (1 cycle)	
For Male and Female NCBP:	Part Fill # 3
Mitte: capsules or cycles. Maximum 63 capsules (3 cycles).  Pharmacy to dispense one cycle at a time, maximum 3 cycles if needed	
Thailinacy to dispense one cycle at a time, maximum 5 cycles if needed	RevAid confirmation number:
STEROID*: CHOOSE ONE	<b>1</b>
One cycle = 28 days	
dexamethasone  40 mg or  20 mg PO once weekly, in the morning,	Lenalidomide lot number:
x doses OR number of 28 day cycles (select one)	
dexamethasonemg PO once weekly in the morning, xdoses	Pharmacist counsel (initial):
OR ☐ number of 28 day cycles (select one) ☐ predniSONE mg PO once weekly in the morning, x ☐doses OR	i namaolot oodnoor (mitaar).
number of 28 day cycles (select one)	
No Steroid	
*Refer to Protocol for steroid dosing options	
Physician to assure DVT prophylaxis in place: ASA, Warfarin , low molecular weight	
heparin, direct oral anticoagulant or none Special Instructions	1
ביים וויפנו עכנוטוופ	
DOCTOR'S SIGNATURE:	SIGNATURE:
Physician RevAid ID:	UC:
Physician RevAid ID:	IIC.



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DATE:		
RETURN APPOINTMENT ORDERS		
☐ Return in weeks for Doctor and Cycle ☐ Last cycle. Return inweek(s)		
Laboratory:		
Cycles 1-4: CBC & Diff, Platelets, Creatinine, Calcium every two weeks		
Serum Protein Electrophoresis <u>and/or</u> Serum Free Light Chain Levels (CIRCLE APPROPRIATE) every 4 weeks		
Blood work done prior to next cycle must be done less than or equal to 7 days prior to the start date		
Cycles 5 and subsequent cycles: CBC & Diff, Platelets, Creatinine, Calcium every 4 weeks, less than or equal to 7 days prior to the next cycle		
Serum Protein Electrophoresis <u>and/or</u> Serum Free Light Chain Levels (CIRCLE APPROPRIATE) every 4 weeks		
TSH Every three months		
☐ Pregnancy blood test for FCBP 7-14 days and 24 h prior to cycle 1 and every week for 4 weeks during cycle 1		
☐ <b>Pregnancy blood test for FCBP,</b> every 4 weeks, less than or equal to 7 days prior to the next cycle		
☐ Bilirubin, ALT		
☐ Other tests		
☐ Consults:		
☐ See general orders sheet for additional requests		
DOCTOR'S SIGNATURE:	SIGNATURE:	
	UC:	