

PROTOCOL CODE: MYDARLD (IV Cycle 2+)

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## Patient RevAid #

		ratient Nev	-\iu π		
DOCTOR'S ORDERS	Ht	cm Wt	kg	BSA	m²
REMINDER: Please ensure drug alle	ergies and previous ble	omycin are doc	umented	on the Alle	ergy & Alert Form
DATE:	To be given:		Cycle	e #:	
Date of Previous Cycle:	aring Potential (FCBP)	Rx valid for 7 da	ıys		
*****Ensure Red Blood Cell Phenotype and Group and Screen for all patients prior to Cycle 1****  Delay treatment week(s)  CBC & Diff, platelets day of treatment  Proceed with all medications as written, if within 96 hours of Day 1: ANC greater than or equal to 1 x 10°/L, platelets greater than or equal to 50 x 10°/L, and eGFR or creatinine clearance as per protocol  Dose modification for:  Hematology: Other Toxicity:  Proceed with treatment based on blood work from					
LENALIDOMIDE One cycle = 28 days  • Per physician's clinical judgement, physicially  □ lenalidomide*mg PO daily  □ lenalidomide*mg PO  MITTE: (*available as 25 mg, 20mg, 15 *Note: Use one capsule strength for the per capsule and not weight based  □ FCBP dispense 21 capsules (1 cycle □ For Male and Female NCBP:  Mitte:capsules orcapsules one cycle at a tiphysician to ensure DVT prophylaxis molecular weight heparin, □ direct of the percent	in the evening, on Days of mg, 10 mg, 5 mg and 2 of total dose; there are considered.  e)  ycles. Maximum 63 caps me, maximum 3 cycles is in place:  ASA,	s 1 to 21 and off for the state of the state	or 7 days costing is	Lenalido Part Fill RevAid Pharmac Part Fill RevAid Lenalido Pharmac Part Fill RevAid Lenalido Lenalido Lenalido	confirmation number:  mide lot number:  cist counsel (initial):  # 2  confirmation number:  mide lot number:  cist counsel (initial):
Special Instructions					
DOCTOR'S SIGNATURE: Physician Revaid ID:				SIGNA UC:	TURE:



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DATE:					
☐ CYCLOPHOSPHAMIDE – Cycles 1 to 8 (☐ Cycle 9 onwards optional)					
cyclophosphamide 500 mg PO once weekly in the morning on Days 1, 8, 15 and 22. Dispense cycles.					
cyclophosphamide mg PO once weekly in the morning on Days	Dispense cycles.				
cyclophosphamide 50 mg PO once in the morning every 2 days for doses. Dispense cycles.					
STEROID (select one)* RN to use patient's therapeutic steroid (if applicable) as pre-m  CYCLE # (Cycle 2 onwards)	ed for daratumumab				
□ dexamethasone □ 40 mg or □ 20 mg PO once weekly on Days 1, 8, 15 and 22. Take dose prior to daratumumab and on weeks without daratumumab, take dose in the morning x doses <u>OR</u> number of 28 day cycles					
dexamethasonemg PO once weekly on Days 1, 8, 15 and 22. Take dose prior to daratumumab and on weeks without daratumumab, take dose in the morning x doses OR number of 28 day cycles					
predniSONEmg PO once weekly on Days 1, 8, 15 and 22. Take dose prior to daratumumab and on weeks without daratumumab, take dose in the morning x doses <u>OR</u> number of 28 day cycles					
☐ No Steroid					
*Refer to Protocol for suggested dosing options					
DOCTOR'S SIGNATURE:	SIGNATURE: UC:				



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DATE:				
**Have Hypersensitivity Reaction Tray and Protocol Available**				
DARATUMUMAB				
• Per physician's clinical judgement, physician to ensure prophylaxis with valACYclovir 500 mg PO daily				
DARATUMUMAB PREMEDICATIONS: Patient to take own supply. RN/Pharmacist to confirm				
dexamethasone as ordered in steroid section				
montelukast 10 mg PO prior to each daratumumab				
acetaminophen 650 mg PO prior to each daratumumab. Repeat acetaminophen 650 needed	mg PO every 4 hours when			
Select one of the following:				
☐ <b>Ioratadine 10 mg</b> PO prior to each daratumumab, then <b>diphenhydrAMINE 50 mg</b> I\	/ every 4 hours when needed			
OR				
☐ diphenhydrAMINE 50 mg ☐PO or ☐ IV prior to each daratumumab. Repeat diphen	hydrAMINE 50 mg IV every 4			
hours when needed				
DARATUMUMAB				
CYCLE 2 Days 1 8 15 and 22.				
CYCLE 2, Days 1, 8, 15, and 22:	sioren in line filter			
daratumumab 16 mg/kg x kg = mg IV in 500 mL NS (use 0.2 n	micron m-ime miler)			
☐CYCLE 3 to 6, Days 1 and 15:				
daratumumab 16 mg/kg x kg =mg IV in 500 mL NS (use 0.2 micron in-lir	ne filter) x cycle(s) (max 2 cycles)			
□CYCLE 7 onwards, Day 1:				
daratumumab 16 mg/kg xkg =mg IV in 500 mL NS (use 0.2 micron in-line filter) x cycle(s) (max 3 cycles)				
Infusion rate for cycle 2 onwards: Physician to determine rate of infusion				
If no reaction in the previous infusion or reaction is Grade 2 or less:				
☐ Start at 200 mL/h. If no infusion-related after 30 minutes, infuse the remainder at 450 mL/h (Rapid infusion)				
OR If reaction in the previous infusion is Grade 3:				
Start at 100 mL/h. If no infusion-related reactions after 60 minutes, increase by 50 mL/h maximum rate of 200 mL/h. Refer to protocol for modified starting rate if previous infusion r during infusion rate of greater than or equal to 100 mL/h (Slow infusion)	•			
<b>Vitals monitoring:</b> Vital signs immediately before the start, at the end of the infusion and a	as needed. Observe patient for			
30minutes after infusion (vitals and observation post- infusion not required after 3 treatment				
DOCTOR'S SIGNATURE:	SIGNATURE:			
	UC:			



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DATE:					
RETURN APPOINTMENT ORDERS					
For Cycles 3 to 6, book chemo on Days 1 and 15					
For Cycle 7 onwards, book chemo on Day 1					
Return in four weeks for Doctor and Cycle and Book chemo x 2 cycles.  Return in twelve weeks for Doctor and Cycles, and  Book chemo x 3 cycles, and  Last Cycle. Return in week(s).					
CBC & Diff, platelets, creatinine, urea, sodium, potassium, total bilirubin, ALT, alkaline phosphatase, calcium, albumin, LDH, random glucose, serum protein electrophoresis <u>and</u> serum free light chain levels every 4 weeks  TSH every three months (i.e. prior to Cycles 4, 7, 10, 13, 16 etc)					
☐ Urine protein electrophoresis every 4 weeks					
☐ Immunoglobulin panel (IgA, IgG, IgM) every 4 weeks					
☐ Beta-2 microglobulin every 4 weeks					
CBC & Diff, platelets Days 8, 15, 22					
Creatinine, sodium, potassium Days 8, 15, 22					
Total bilirubin, ALT, alkaline phosphatase Days 8, 15, 22					
Random glucose Days 8, 15, 22					
Calcium, albumin Days 8, 15, 22					
<ul><li>☐ Quantitative beta- hCG blood test for FCBP, every 4 weeks, less than or equal to 7 days prior to the next cycle</li><li>☐ Other tests:</li></ul>					
☐ Consults:					
☐ See general orders sheet for additional requests					
DOCTOR'S SIGNATURE:	SIGNATURE:				
	UC:				