

Standard Pharmaceutical Product and Medical Device Information (Rx Product Only)

Version 2021						Introduction Type	: New Item		x Final Version			Date:	5/17/	2023
			PRODUCT INFORMA	TION					SPECIAL HAN	DLING AND STOR	AGE REQUIF	REMENTS*		
Company Name:	Camber Pharmac	euticals, Inc.				Application	ANDA	a. Temperatu	re - Indicate the USP temp	erature range for th	is product.			
Application Number for NDA/AN	NDA/BLA (drug); PN	MA/510(k)(med device	ce):	21	2414		<u> </u>	1	Temperature Range	Controlled Room -		and 25 C (68	° – 77° F)	
Medical Device Class, if applical									, ,					
DUNS:	82-677-4775							_	Other Temperature Range	Requirement	Excursions p	ermitted to 1	5°C to 30°C (59° – 86° F)
Proprietary Name (If Applicable) a	and Established Na	ame: Lenali	idomide Capsules 15mg					I	(write in)	•				
Selling Unit NDC:	31722-260-21		Unit of Use NDC:	:	31722-260-21	UPC: 33	1722260213		Notes					
UDI			CVX Code:			MVX Code:								
Description:	Lenalidomide Cap	osules 15mg						T	Is this product to be shippe	to customers on ic	e?		No	
									Is this product to be shippe				No	
Active Ingredient(s):		Lenalidomide						1			•			
								b. Contact fo	r temperature excursion qu	estions:				
URL for Additional Product Inform	mation:	www.camberpharma	ı.com						Name:		Soma Raju			
Address:		Ave (and) 800 Center	nnial Ave, Suite 1			Address 2:			Number:		732-529-042			
City:	Piscataway				State:		p: 08854	_	Group E-mail:		somaraju@h	eterousa.cor	<u>n</u>	
Key Contact:	Customer Service	9			Email:	customerservice@ca	mberpharma.com							
Phone Number:	1-866-827-3647				Fax:	732-562-8788		c. Special reg	gulations for product in any				No	
Product Therapeutic Classification	on:	Immunomodulatory	Agent						Special returns requirement	s for this product?			*Yes	
	ADDITI	ONAL PROPUST IN	FORMATION			PRODUCT DEC	ODIDTION INFORMATION							
	ADDITI	ONAL PRODUCT IN	FORMATION			PRODUCT DES	CRIPTION INFORMATION	d. Store prod	luct (unit of sale) upright?				No	
The product is?			Is the Product	Direct And D	Prop-Ship				Protect product (unit of sa	ile) from light?			No	
a legend device?		No	Is the Product	Unit of Use		Size:	21ct	e. Shelf life:					24	Months
if yes, enter class #			Orphan Drug Status						Initial shelf life at launch (if different):				Months
a product kit?		No	FDA 4			Strength:	15mg			ORDER INFORM	ATION			
if yes, list NDCs of			FDA Approval Status				hard gelatin capsule			ORDER INFORM	ATION			
component parts reverse numbered?		No				Dosage Form:	naru gelatiri capsule		Unit of Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present						1 Bottle		1 bottle of 21			
latex-free?		Yes	_				capsule shaped, size '0'		Box/Carton		(Write-in, e.		0 Vials)	
preservative-free?		Yes	Dairy ar	nd Lactose		Product Shape:	.,,,		Ampule		, , , , ,	,	,	
correctional institution block?		No				Product Color:	red opaque cap/white		Glass		Minimum or	der quantity	?	Yes
opioid?		No				Product Color:	opaque body		Tube					
Cannabinoid?		No	Country of Origin	India		Product Imprint:	H' on cap and 'L5' on		Vial Liquid Sgl					
If Unit Dose, is item bar coded to	unit dose for					r roduct imprint.	body		Vial Liquid Multi				ch package t	ype?
hospital scanning?			Is this product covered u						Vial Powder Sql			Each		
If Unit Dose, indicate NDC here:			Trade Agreements Act (TAA)?	No				Vial Power Multi			Inner/Carton	/Pack	
									Other: Write In			Case		
			FOR GENERIC DRUG PR	ODUCTS										
							Authorized Conseis other		DI.	ARMACY ORDER	/ DILL LINET			
					Aut		Authorized Generic, other ction fields are not applicable			ARMACT URDER				
I. Orange Book Rating:	AB					56	ction fields are not applicable	Rec. sell unit	to customer?		Rx billing u		acy:	
II. Generic Equivalent to What Bra	and?:	Revlimid						00/-11- 1	4 \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \			Each		
		DRUG SUBBI	LY CHAIN SECURITY ACT	(Decea) INFO	RMATION			(Write-in, e.g	. 1 Vial)			Gram Milliliter		
		DRUG SUFFE	IT CHAIN SECONITT ACT	(DSCSA) IN O	KMATION							wiiiiiitei		
Does supplier meet DSCSA defini	ition of manufactur	rer?	Yes	\neg	GLN:	0331722000000			ITEN	I AND PACKING IN	FORMATION			
Is product exempt from DSCSA?				_	02.11	0001122000000						•		
			No											Saleable #
If you calact examplian.			No		CCD.					Dimonei	ons (US men	its)		Pieces
If yes, select exemption: Other exemption - Write in:			No		GCP:]	Weight Lbs.		ons (US msn Width	•	Volume (Cube)	0000
Other exemption - Write in:			No			ginal product purchas	sed	Item/Each		Depth	Width	Height	(Cube)	
	s exclusive distribu	utor?				ginal product purchas	sed	Item/Each:	Weight Lbs.		•	•		1
Other exemption - Write in: Is product repackaged?			No		If yes, was ori			Item/Each:	0.072	Depth	Width	Height	(Cube)	1
Other exemption - Write in: Is product repackaged? Is product sold by manufacturer's	on/exemption for pr		No Yes		If yes, was ori	ir?			0.072	Depth	Width	Height	(Cube)	1
Other exemption - Write in: Is product repackaged? Is product sold by manufacturer's Has FDA granted waiver/exceptio	on/exemption for pr	roduct?	No Yes No		If yes, was ori	ir?		Box/Carton/E	0.072 Bundle/	Depth 1.6	Width 1.6	Height 2.9	(Cube) 7.424	
Other exemption - Write in: Is product repackaged? Is product sold by manufacturer's Has FDA granted waiver/exceptio	on/exemption for pr	roduct?	No Yes	NFORMATION	If yes, was ori	ir?		Box/Carton/E	0.072	Depth	Width	Height	(Cube)	24
Other exemption - Write in: Is product repackaged? Is product sold by manufacturer's Has FDA granted waiver/exceptio If yes, attach documentation fro	on/exemption for pr om FDA.	roduct?	No Yes No No	NFORMATION	If yes, was ori direct from m Provide source	r? e manufacturer for re	packaged product	Box/Carton/E	0.072 Bundle/	Depth 1.6	Width 1.6	Height 2.9	(Cube) 7.424	
Other exemption - Write in: Is product repackaged? Is product sold by manufacturer's Has FDA granted waiver/exceptio If yes, attach documentation fro	on/exemption for pr om FDA.	GTI Saleable Quantity	No Yes No	NFORMATION	If yes, was ori direct from mi Provide source	r? e manufacturer for re	packaged product Unit of Use GTIN-14	Box/Carton/E Inner Pack: Case:	0.072 Bundle/	Depth 1.6	Width 1.6	Height 2.9	(Cube) 7.424	
Other exemption - Write in: Is product repackaged? Is product sold by manufacturer's Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure x Item/Each	on/exemption for pr om FDA.	roduct?	No Yes No No	NFORMATION	If yes, was ori direct from mi Provide source	r? e manufacturer for re	packaged product	Box/Carton/E Inner Pack: Case:	0.072 3undle/ 2.758	Depth 1.6	Width 1.6 6.5	4.13	7.424 264.1548	24
Other exemption - Write in: Is product repackaged? Is product sold by manufacturer's Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure X tem/Each Box/Carton/Bundle/Inner Pack	on/exemption for pr om FDA.	Galeable Quantity	No Yes No No	NFORMATION	If yes, was ori direct from mt Provide source GTIN 0033	F. P.	packaged product Unit of Use GTIN-14	Box/Carton/E Inner Pack: Case:	0.072 Bundle/	Depth 1.6	Width 1.6 6.5	4.13	(Cube) 7.424	24
Other exemption - Write in: Is product repackaged? Is product sold by manufacturer's Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure X	on/exemption for pr om FDA.	GTI Saleable Quantity	No Yes No No	NFORMATION	If yes, was ori direct from mt Provide source GTIN 0033	r? e manufacturer for re	packaged product Unit of Use GTIN-14	Box/Carton/E Inner Pack: Case: Pallet:	0.072 3undle/ 2.758 COST INFORMATION	Depth 1.6	Width 1.6 6.5	4.13	7.424 264.1548	24
Other exemption - Write in: Is product repackaged? Is product sold by manufacturer's Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure X tem/Each Box/Carton/Bundle/Inner Pack	on/exemption for pr om FDA.	Galeable Quantity	No Yes No No	NFORMATION	If yes, was ori direct from mt Provide source GTIN 0033	F. P.	packaged product Unit of Use GTIN-14	Box/Carton/E Inner Pack: Case: Pallet: Regular Cost	0.072 3undle/ 2.758 COST INFORMATION	Depth 1.6 9.84	Width 1.6 6.5 Vendor #:	Height 2.9 4.13	7.424 264.1548	24
Other exemption - Write in: Is product repackaged? Is product sold by manufacturer's Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure X	on/exemption for pr om FDA.	Galeable Quantity	No Yes No No	NFORMATION	If yes, was ori direct from mt Provide source GTIN 0033	F. P.	packaged product Unit of Use GTIN-14	Box/Carton/E Inner Pack: Case: Pallet:	0.072 3undle/ 2.758 COST INFORMATION	Depth 1.6	Width 1.6 6.5 Vendor #:	Height 2.9 4.13	7.424 264.1548	24
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Other exemption - Write in: Is product repackaged? Is product sold by manufacturer's Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure x	on/exemption for pr om FDA.	Galeable Quantity	No Yes No No	NFORMATION	If yes, was ori direct from mt Provide source GTIN 0033	F. P.	packaged product Unit of Use GTIN-14	Box/Carton/E Inner Pack: Case: Pallet: Regular Cost Invoice Cost	0.072 3undle/ 2.758 COST INFORMATION (WAC) (\$)	Depth 1.6 9.84	Width 1.6 6.5 Vendor #:	Height 2.9 4.13	7.424 264.1548	24
Other exemption - Write in: Is product repackaged? Is product sold by manufacturer's Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure x	on/exemption for pr om FDA.	Galeable Quantity	No Yes No IN AND HIBCC PRODUCT I		If yes, was ori direct from mr Provide source GTIII 0033	I-14 I-1722260213	packaged product Unit of Use GTIN-14	Box/Carton/E Inner Pack: Case: Pallet: Regular Cost Invoice Cost As of date:	0.072 3undle/ 2.758 COST INFORMATION (WAC) (\$) 4/20/2023	Depth 1.6 9.84	Width 1.6 6.5 Vendor #:	Height 2.9 4.13	7.424 264.1548	24



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For Designated Drop Ship Only Products, Please Use Page 3

MATI	ERIAL HAZ	ARD CLASSIFICATION and TRANSPORTATION					
Is this product (check all that apply):							
a. Cytotoxic? No		SDS Hazard Classification					
b. CA Prop. 65 Carcinogen or Reproductive Toxicant?							
Is the product a CA Prop 65 carcinogen?		x Organic	Corrosive				
Is the product a CA Prop 65 reproductive toxicant?		Inorganic	Oxidizer				
Does the product label bear a CA Prop 65 warning?	No	Steroid/Androgen	Contact Hazard				
c. Contact Hazard?	Yes	Does the product have an Aerosol class? If yes,					
d. Does this product require special clean-up instructions?	Yes	identify NFPA Storage Level:					
(If yes, attach SDS with special instructions.)		NFPA Storage Level:					
e. Does the product contain DEHP?	No						
Is this product regulated for shipment by DOT?	No	Is the product a NIOSH hazardous drug?	Yes				
(if yes, answer a-e below and provide SDS)		If yes, indicate which:	Group 1 items (antineoplastic)				
a. UN/Identification Number		, ,					
b. Proper Shipping Name							
c. DOT Hazard Class		Haza	rdous Waste Identification				
d. Packing Group							
e. Inhalation Hazard?	No	EPA Hazardous Waste Code: WT02, MN01, R006	Waste Characteristics D004				
Is this product regulated for shipment by IATA?	No						
(if yes, answer a-e below and provide SDS)		REMS o	r REGISTRY RESTRICTIONS				
a. UN/Identification Number							
b. Proper Shipping Name		Is there a REMS on this product?	Yes				
c. DOT Hazard Class		If Yes, is it managed with a pharmacy registry?	Yes				
d. Packing Group		Website URL:	www.lenalidomiderems.com				
e. Inhalation Hazard?	No						
Is the product restricted for air shipment? If so, indicate restriction:	No	Med Guide Required	Yes				
Passenger		Limited Distribution Requirement	Yes				
Cargo		Comments / Details: (For example, iPledge program?)	Must be a certified Lenalidomide REMS Program Location				
Passenger & Cargo							
Is this a reportable quantity? No		REMS:	Yes				
RQ Threshold:		REMS Program Manager Name:	Bristol Myers Squibb Phone: 1-888-423-5436				
Is this a marine pollutant? Yes		Supplier Manages REMS registry exclusively:	No				
Is this product shipped utilizing an authorized DOT exception or Special Permit?		Wholesale distributor support:	No				
No (if yes, identify method below)		Provider Name:	DEA #:				
Limited Quantity		Site Enrollment Number assigned	NCPDP#:				
Consumer Commodity, ORM-D		by Supplier:	NPI #:				
Small Quantity (49 CFR 173.4)							
Special Permit; DOT-SP		Comments					
Special Provision (listed in Column 7 of 49 CFR 172.101);							
SP#		Registry:	Yes				
		Registry Program Contact Name:	REMS Call Center Phone: 1-888-423-5436				
ADD'L STORAGE INFORMATION		Comments Lenalidomide REMS	is a shared REMS program				
Is the Product							
Controlled Substance? No Controlled Substance Code			ETURN INSTRUCTIONS				
Controlled by State(s)? No Listed Chemical (List I or II)	No						
ARCOS Reportable? No If yes, indicate which:		Contact tel. # if product received damaged:	1-888-423-5436				
Schedule No. Is it a scheduled listed chemical product?:	No	Is product returnable for credit:	No				
CLASS OF TRADE RESTRICTION:		URL/Link to returns policy:					
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices	Yes	contact - www.lenali	domiderems com				
			dominations.com				
Restricted to retail pharmacy only:	No	Special regulations or returns requirements for this					
Restricted to hospital, clinics, and physician offices only:	No	product in certain states?	Yes				
Restricted from US territories? (explain in comments)	No	If so, which states? Other requirements? Comments?					
Comments:			1-888-423-5436 to return directly to Lenalidomide REMS program. (All States)				
		Non-Dispensed Product Returns: Return directly to Camber's third Damaged in Transit Returns (by carrier): Return to Camber Distribution	party return goods processor. (All States)				
MIS	CELLANEC	OUS NOTES and/or Image of Product Barcode:	Autori Goritor, trait Glatest				
- WIGH							



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FOR DESIGNATED DROP SHIP PRODUCT ONLY - if not a designated drop ship, do not complete.

Order Method for Desig	nated Drop Ship Product	Standard Order Receipt and Processing
Contracted 3PL company / contact #: Name: Phone:	Fax Number: 732-562-8788 Fax Number: 732-562-8788	Purchase order daily receipt cut off time by supplier Cut off time: 11:00 AM Monday - Thursday
Expedited freight fees billed with each order:	Yes	Overnight receipt available: No
Drop Ship service fee billed with each order:	No	PO Receipt cut off time:
Drop Ship miscellaneous fees billed: Comments:	No	Days of week overnight is available: Monday Tuesday Wednesday Thursday Friday Priority Overnight receipt available:
Class of Tra	de Restriction:	PO Receipt Cut off time:
No restriction: Select YES if sold to retail pharmacy, Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices o Restricted from US territories? (explain in comments	hospitals, clinics and physician offices No No No	Saturday Overnight receipt available: PO Receipt Cut off time: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:
Other Data Information	Required to Process PO:	Return Instructions
Patient Procedure Date: Physician Name: Physician/Clinic Phone # Physician State License # Physician/Clinic DEA #: Physician/Clinic Specialty: Miscelland	eous Notes:	Contact # if product is received damaged: Is product returnable for credit: URL/Link to returns policy: https://www.camberpharma.com/partner-resources/#returned-goods-policy Special regulations or returns requirements for this product in certain states? If so, which states? Other requirements? Comments? Dispensed Product Returns: Return to prescriber, pharmacy or call 1-888-423-5436 to return directly to Lenalidomide REMS program. (All States)
		Non-Dispensed Product Returns: Return directly to Camber's third party return goods processor. (All States) Damaged in Transit Returns (by carrier): Return to Camber Distribution Center. (All States)
		ADDITIONAL INFORMATION
		Is product order for scheduled patient procedure? Is product order for restocking purposes? Yes Yes