

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

Version 2021						Introduction Type:	Post Launch Change		x Final Version			Date:	11/29	9/2024
			PRODUCT INFORMA	TION					SPECIAL HAN	DLING AND STOR	RAGE REQUI	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc. Application: ANDA						a. Temperature – Indicate the USP temperature range for this product.								
Application Number for NDA/AN	IDA/BLA (drug); PN	IA/510(k)(med devi	ice):	21	0500				Temperature Range	Controlled Room		and 25 C (68	3° – 77° F)	
Medical Device Class, if applica									· -					
DUNS:	11-856-3719								Other Temperature Range F	Requirement				
Proprietary Name (If Applicable) a	and Established Na	me: Dime	thyl Fumarate Delayed-Relea	se Capsules 12	0 mg				(write in)					
Selling Unit NDC:	31722-657-31		Unit of Use NDC		31722-657-31		22657310		Notes					
UDI			CVX Code:			MVX Code:								
Description:	Dimethyl Fumarate	Delayed-Release	Capsules 120 mg						Is this product to be shipped	I to customers on i	ce?		No	1
	-								Is this product to be shipped				No	1
Active Ingredient(s):		Dimethyl fumarate	•											
								b. Contact for temperature excursion questions:						
URL for Additional Product Inform		www.camberpharn	na.com						Name:		Soma Raju			
Address:	800 Centennial Av	e, Suite 1			State:	Address 2: NJ Zip	00054		Number:		732-529-042			
City: Key Contact:	Piscataway Customer Service				Email:	customerservice@cam	: 08854		Group E-mail:		somarajuei	neterousa.com	<u>11</u>	
Phone Number:	1-866-827-3647				Fax:	732-562-8788	berpriama.com	c Special rec	ulations for product in any	etatos?			No	٦
Product Therapeutic Classification		Fumaric acid deriv	rative (NRF2 activator)		- I ux.	732 302 0700		c. Special reg	Special returns requirement				No	-
Troduct Therapeutic Glassification	//··	T dillatic acid acity	rative (rara 2 delivator)						opecial returns requirement	s for this product:			140	_
	ADDITIO	ONAL PRODUCT IN	NEORMATION			PRODUCT DESC	RIPTION INFORMATION	d Store prod	uct (unit of sale) upright?				No	٦
The was dead in 0	7,55,111	J.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		Direct-Ship C)nlv	1 1105001 5200		u. otore prou		I-) (l'-l-10				4
The product is? a legend device?		No	Is the Product Is the Product	Unit of Use	niiy		14 ct	e. Shelf life:	Protect product (unit of sa	le) from light?			No 24	Mantha
if yes, enter class #		INO	Orphan Drug Status	Offic of Ose		Size:	14 Ct	e. Shell life:	Initial shelf life at launch (f different):			24	Months Months
a product kit?		No	Orphan Drug Status				120 mg		illitiai Sileli ille at laulicii (i dillerentj.				Wionins
if yes, list NDCs of		140	FDA Approval Status			Strength:	125 mg			ORDER INFORM	MATION			
component parts						B	Hard gelatin, delayed-							
reverse numbered?		No				Dosage Form:	release capsule		Unit of Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present						x Bottle		1 Bottle of 1	4 Capsules		
latex-free?		Yes	Dye, Corn, Alcohol, A		s, Sugar,	Product Shape:	Capsule		Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free?		Yes	Oats	, Spelt		i roduct onapc.			Ampule					
correctional institution block?		No				Product Color:	Opaque light blue cap		Glass		Minimum o	rder quantity	/?	Yes
opioid?		No					and opaque light blue		Tube					
Cannabinoid?		No	Country of Origin	India		Product Imprint:	Imprinted with 'H' on cap		Vial Liquid Sgl					
If Unit Dose, is item bar coded to	unit dose for		In this was don't account to	and an the		•	and 'D12' on body		Vial Liquid Multi Vial Powder Sql			many of whi	ich package	type?
hospital scanning? If Unit Dose, indicate NDC here:			Is this product covered Trade Agreements Act (No				Vial Powder Sgi Vial Powder Multi		24	Inner/Cartor	/Pook	
II Offit Dose, indicate NDC fiele.			Trade Agreements Act (IAA):	INU				Other: Write In			Case	I/Fack	
			FOR GENERIC DRUG PF	ODUCTS					Other: Write III			Ousc		
			TOR GENERIC DROG FI	000013										
					Aut	horized Generic *If A	uthorized Generic, other		PH	ARMACY ORDER	/ BILL UNIT			
I Oronno Book Betimer	AB				7.0		on fields are not applicable	Pac call unit	to customer?					
I. Orange Book Rating: II. Generic Equivalent to What Bra		Tecfidera						itec. sen unit	to customer:	1	KX billing u	nit to pharm Each	acy:	
II. Generic Equivalent to What Bra	anur.	Techidera						(Write-in, e.g.	1 Vial)	I		Gram		
		DRUG SUPP	PLY CHAIN SECURITY ACT	(DSCSA) INFO	RMATION			(**************************************	· viai,			Milliliter		
				,										
Does supplier meet DSCSA defin	ition of manufacture	er?	Yes		GLN:	0331722498975			ITEN	AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			No											
If yes, select exemption:					GCP:			1		Dimens	ions (US msr	nts.)	Volume	Saleable #
Other exemption - Write in:								· [Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		If yes, was or	iginal product purchase	d	Item/Each:	0.07	1.8	1.8	3.6	11.66	1
Is product sold by manufacturer's			Yes	_	direct from m					1.0	1.0	3.0	11.00	'
Has FDA granted waiver/exception		oduct?	No		Provide source	e manufacturer for repa	ckaged product	Box/Carton/B	undle/					
If yes, attach documentation fro	m FDA.							Inner Pack:						
		61	TIN AND HIBCC PRODUCT I	NEODMATION				Case:	2	11.5	8.75	4.5	452.81	24
		GI	TIN AND RIBCC PRODUCT	NFORMATION				Pallet:			-			
Saleable Unit of Measure	c	aleable Quantity	HIBCC		GTII	1.14	Unit of Use GTIN-14	Pallet:						
X Item/Each	3	1	TIIDOO			31722657310	00331722657310							
Box/Carton/Bundle/Inner Pack					0000		200022007070		COST INFORMATION			WHOLESAL	ER USE ONL	LY:
X Case		24			2033	31722657314								
Pallet								Regular Cost			Vendor #:			
								Invoice Cost	(WAC) (\$)	\$37.50	Whsl. Code	#:		
											Fineline Co	de:		
								As of date:	12/1/2024		ļ.			
ļ.								Ц			<u> </u>			
			Attach copy of SAFETY D	ATA SHEET (SE	S) or non hazaı	d letter, PACKAGE INSE	RT, LABEL AND PHOTO OF F	PRODUCT PACKA	AGING and BARCODE.					
*Please provide any additional in			/	(01	-,		nated Drop Ship Only.		Signature:					



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For Designated Drop Ship Only Products, Please Use Page 3

MATERIAL HAZARD 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? Is the product a CA Prop 65 reproductive toxicant? No Does the product label bear a CA Prop 65 warning? No	x Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard						
c. Contact Hazard? d. Does this product require special clean-up instructions? (If yes, attach SDS with special instructions.) e. Does the product contain DEHP?	Does the product have an Aerosol class? If yes, identify NFPA Storage Level: NFPA Storage Level:						
Is this product regulated for shipment by DOT? (if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name	Is the product a NIOSH hazardous drug? If yes, indicate which:						
c. DOT Hazard Class d. Packing Group	Hazardous Waste Identification						
e. Inhalation Hazard? Is this product regulated for shipment by IATA? No	EPA Hazardous Waste Code: Waste Characteristics						
(if yes, answer a-e below and provide SDS) a. UN/Identification Number	REMS or REGISTRY RESTRICTIONS						
b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	Is there a REMS on this product? If Yes, is it managed with a pharmacy registry? Website URL:						
Is the product restricted for air shipment? If so, indicate restriction: Passenger Cargo Passenger & Cargo	Med Guide Required Limited Distribution Requirement Comments / Details: (For example, iPledge program?)						
Is this a reportable quantity? No RQ Threshold: Is this a marine pollutant? No Is this product shipped utilizing an authorized DOT exception or Special Permit? No (if yes, identify method below) Limited Quantity Consumer Commodity, ORM-D Small Quantity (49 CFR 173.4)	REMS: REMS Program Manager Name: Supplier Manages REMS registry exclusively: Wholesale distributor support: Provider Name: Site Enrollment Number assigned by Supplier: No Phone: DEA #: NCPDP#: NPI #:						
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101); SP#	Comments						
ADD'L STORAGE INFORMATION	Registry: Registry Program Contact Name: Comments No Phone:						
Is the Product							
Controlled Substance Code Controlled by State(s)? ARCOS Reportable? Schedule No. No Controlled Substance Code Listed Chemical (List I or II) No If yes, indicate which: Is it a scheduled listed chemical product?: No CLASS OF TRADE RESTRICTION:	RETURN INSTRUCTIONS Contact tel. # if product received damaged: I-866-827-3647 Is product returnable for credit: Yes						
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Yes	URL/Link to returns policy: contact - customerservice@camberpharma.com						
Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: No	Special regulations or returns requirements for this product in certain states?						
Restricted from US territories? (explain in comments) No Comments:	If so, which states? Other requirements? Comments?						
MISCELLANE	DUS NOTES and/or Image of Product Barcode:						



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

Order Method for Designated Drop S	nip Product	Standard Order Receipt and Processing
Purchase orders may be accepted by: a. EDI		Purchase order daily receipt cut off time by supplier Cut off time:
b. Autofax c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Phone:	per:	Shipping lead time of PO: Hours Days Ships same day for next day receipt: Ships for second day receipt: Ships regular ground for 3-10 days receipt:
Expedited Freight Charges or Other Designa	ed Drop Ship Fees:	Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each order: Drop Ship service fee billed with each order:		Overnight receipt available: PO Receipt cut off time:
Drop Ship miscellaneous fees billed: Comments:		Days of week overnight is available: Monday Tuesday Wednesday Thursday Friday
		Priority Overnight receipt available:
Class of Trade Restriction		PO Receipt Cut off time:
No restriction: Select YES if sold to retail pharmacy, hospitals, clinic Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	s and physician offices	Saturday Overnight receipt available: PO Receipt Cut off time: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:
Other Data Information Required to F	rocess PO:	Return Instructions
Patient Procedure Date: Physician Name: Physician/Clinic Phone # Physician State License # Physician/Clinic DEA #: Physician/Clinic Specialty:		Contact # if product is received damaged: Is product returnable for credit: URL/Link to returns policy: Special regulations or returns requirements for this product in certain states? If so, which states? Other requirements? Comments?
Miscellaneous Notes:		
		ADDITIONAL INFORMATION
		Is product order for scheduled patient procedure? Is product order for restocking purposes?