

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

Version 2021						Introduction T	уре:	Post Launch Change	]	x Final Version			Date:	10/22	2/2025
			PRODUCT INFORMA	TION						SPECIAL H.	ANDLING AND STO	RAGE REQU	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc. Application: ANDA						a. Temperature – Indicate the USP temperature range for this product.									
Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): 214603 Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)															
Medical Device Class, if applical	ble:								I						
DUNS:	11-856-3719									Other Temperature Rang	e Requirement				
Proprietary Name (If Applicable) a		ame: Gem	fibrozil Tablets, USP 600 mg						1	(write in)					
Selling Unit NDC: UDI	31722-128-05		Unit of Use NDC: CVX Code:			UPC: MVX Code:	33172212	8056	+	Notes					
			CVA Code.			INVX Code.			-						
Description:	Gemfibrozil Table	ts, USP 600 mg								Is this product to be ship				No No	-
Active Ingredient(s): Gemfibrozil, USP							+	Is this product to be shipped to customers on dry ice?							
Active ingredient(s).		CCITIIDIOZII, CCI							b. Contact for temperature excursion questions:						
URL for Additional Product Inforn	nation:	www.camberpharr	na.com							Name: Soma Raju					
Address:	800 Centennial A	ve, Suite 1				Address 2:			]	Number:		732-529-04			
City:	Piscataway				State:	NJ	Zip: 08		Group E-mail: somaraju@heterousa.com						
Key Contact:	1-866-827-3647	)			Email: Fax:	customerservice@camberpharma.com 732-562-8788			. Consider		4-42			No	1
Phone Number: Product Therapeutic Classificatio		Lipid regulating ag	ront		гах:	132-302-0100			c. Special re	gulations for product in a Special returns requirem	-	,		No	-
Product Therapeutic Classificatio	on:	Lipid regulating ag	Jent .							Special returns requirem	ents for this product			NO	
	ADDITI	ONAL PRODUCT I	NFORMATION			PRODUCT	DESCRIPTI	ION INFORMATION	d Store proc	duct (unit of sale) upright	,			No	1
The product is?			Is the Product	Direct-Ship C	nly				an otono proc	Protect product (unit of				No	1
a legend device?		No	Is the Product	Neither	Jilly		500	n et	e. Shelf life:	Protect product (unit of	sale) from light?			24	Months
if yes, enter class #		140	Orphan Drug Status			Size:	000	0 01	C. Onen me.	Initial shelf life at launc	n (if different):			2-1	Months
a product kit?		No				Ctue w mth.	600	0 mg			(				
if yes, list NDCs of			FDA Approval Status			Strength:					ORDER INFOR	MATION			
component parts						Dosage Form	n: Filn	m coated tablet							
reverse numbered?		No	All B							Unit of Sale			NDC selling	unit?	
co-licensed?		No Yes	Allergens Present				Ova	al		x Bottle Box/Carton		1 Bottle of 5	g. 1 Box of 1	0 Viale)	
preservative-free?		Yes	Corn, Alco	hol, Animal		Product Sha	pe:	ai		Ampule		(vviite-iii, e	.g. I Dox of I	U Viais)	
correctional institution block?		No				Burnet Call	Wh	nite to off white		Glass		Minimum o	rder quantity	<b>y</b> ?	Yes
opioid?		No				Product Cold	or:			Tube				•	
Cannabinoid?		No	Country of Origin	USA		Product Imp	rint Debo	ossed with '1' with two partial cts on one side and two partial		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	unit dose for							cts on other side		Vial Liquid Mult				ich package	type?
hospital scanning?			Is this product covered u Trade Agreements Act (1		Vaa					Vial Powder Sg		12	Each	-/DI/	
If Unit Dose, indicate NDC here:			Trade Agreements Act (1	AA)!	Yes					Vial Powder Mu Other: Write In	IU		Inner/Cartor Case	I/Pack	
			FOR GENERIC DRUG PR	ODUCTS		<u> </u>			1	Outer: Write iii			Ousc		
			TOR GENERIO BROOT R	000010											
					Au	uthorized Generic	*If Authoria	zed Generic, other			PHARMACY ORDE	R / BILL UNIT			
I. Orange Book Rating:	AB				section fields are not applicable				Rec. sell unit to customer?			Rx billing ı	Rx billing unit to pharmacy:		
II. Generic Equivalent to What Bra		Lopid							T				Each	,	
						(Write-in, e.g. 1 Vial) Gram									
		DRUG SUPF	PLY CHAIN SECURITY ACT (	DSCSA) INFOR	RMATION								Milliliter		
Dana aumuliau maast DCCCA dafini	:tian af manufaat	2	Voc	_	CI N.	0224722400075				IT.	EM AND PACKING	INFORMATIO	M		
Does supplier meet DSCSA defini Is product exempt from DSCSA?	ition of manufactul	lei r	Yes No	+	GLN:	0331722498975					LM AND PACKING	INFORMATIC	N-		
If yes, select exemption:					GCP:				1		Dimon	sions (US ms	mte \	Volume	Saleable #
Other exemption - Write in:					GCP:				1	Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		If ves. was o	riginal product pure	chased		Item/Each:					<u> </u>	
Is product sold by manufacturer's	s exclusive distribu	itor?	Yes		direct from n					1.1	3.5	3.5	6.75	82.69	1
Has FDA granted waiver/exceptio		roduct?	No		Provide sour	ce manufacturer fo	r repackag	jed product	Box/Carton/E	Bundle/					
If yes, attach documentation from	m FDA.								Inner Pack:						
		67	TIN AND HIBCC PRODUCT II	JEODMATION					Case:	14	14.25	11	7.5	1175.63	12
		G	TIN AND RIBCC PRODUCT IF	NFORMATION					Pallet:						
Saleable Unit of Measure	8	Saleable Quantity	HIBCC		GTI	IN-14	U	Init of Use GTIN-14	r allet.						
X Item/Each	_	1				31722128056									
A Monte Edon						COST INFORMATION WHOLESALER USE ONLY:					_Y:				
Box/Carton/Bundle/Inner Pack					103	331722128053									
Box/Carton/Bundle/Inner Pack X Case		12							Regular Cos	t		Vendor #:			
Box/Carton/Bundle/Inner Pack		12											. и.		
Box/Carton/Bundle/Inner Pack X Case		12							Invoice Cost		\$59.8	Whsl. Code			
Box/Carton/Bundle/Inner Pack X Case		12							Invoice Cost		\$59.8				
Box/Carton/Bundle/Inner Pack X Case		12								(WAC) (\$)	\$59.8	Whsl. Code			
Box/Carton/Bundle/Inner Pack X Case		12							Invoice Cost	(WAC) (\$)	\$59.8	Whsl. Code			
Box/Carton/Bundle/Inner Pack X Case		12	Attach copy of SAFETY DA	TA SHEET (SC	DS) or non haza	ard letter, PACKAGE	INSERT, L	ABEL AND PHOTO OF F	As of date:	(WAC) (\$) 3/8/2021	\$59.8	Whsl. Code			



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

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For Designated Drop Ship Only Products, Please Use Page 3

MATERI.	AL HAZARD CLASSIFICATION and TRANSPORTATION					
Is this product (check all that apply): a. Cytotoxic? b. CA Prop. 65 Carcinogen or Reproductive Toxicant?	SDS Hazard Classification					
Is the product a CA Prop 65 carcinogen? Is the product a CA Prop 65 reproductive toxicant?	x     Organic     Corrosive       /es     Inorganic     Oxidizer       No     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?	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:  Phone:  DEA #: NCPDP#: NCPDP#: NPI #:					
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101);	Comments					
SP# ADD'L STORAGE INFORMATION	Registry:  Registry Program Contact Name:  Comments  No  Phone:					
Is the Product	Comments					
Controlled Substance?  Controlled by State(s)?  ARCOS Reportable?  No  Controlled Substance Code  Listed Chemical (List I or II)  If yes, indicate which:	RETURN INSTRUCTIONS  Contact tel. # if product received damaged:  Is product returnable for credit:  Yes					
	URL/Link to returns policy:  'es contact - customerservice@camberpharma.com					
	- J					
Restricted to hospital, clinics, and physician offices only:	Special regulations or returns requirements for this product in certain states?  If so, which states? Other requirements? Comments?					
Comments:	No If so, which states? Other requirements? Comments?					
MISGEL	LANEOUS 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 Ship 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 #:	Fax Number: Fax Number: Phone No.: Site Address:	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:
	Phone:	
Expedited Freight Cha	arges or Other Designated Drop Ship Fees:	Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each orde	er:	Overnight receipt available:
Drop Ship service fee billed with each orde	er:	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:
Cla	ass of Trade Restriction:	PO Receipt Cut off time:
No restriction: Select YES if sold to retail p Restricted to retail pharmacy only: Restricted to hospital, clinics, and physicia Restricted from US territories? (explain in of Comments:		Saturday Overnight receipt available:  PO Receipt Cut off time: Phone: Fax: EDI:  Overnight Fees apply: Other fees apply:
Other Data In	formation Required to Process PO:	Return Instructions
Patient Procedure Date: Physician Name: Physician/Clinic Phone # Physician State License # Physician/Clinic DEA #: Physician/Clinic Specialty:	Miscellaneous Notes:	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?
	misochaneous notes.	
		ADDITIONAL INFORMATION
		Is product order for scheduled patient procedure? Is product order for restocking purposes?