

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

Version 2021						Introduction	Туре:	Post Launch Change)	Final Version			Date:	6/23/	/2024
			PRODUCT INFORMA	TION						SPECIAL HAN	NDLING AND STOP	RAGE REQUI	REMENTS*	m	
Company Name:	Camber Pharmaceutic	als. Inc.				Applica	ation:	ANDA	a. Temperature – Ir	dicate the USP temp	erature range for t	his product.			
Application Number for NDA/AN			e):	202	2438					perature Range	Controlled Room		and 25 C (68	° – 77° F)	
Medical Device Class, if applicat			,							Ū.					
DUNS:	11-856-3719								Othe	r Temperature Range	Requirement				
Proprietary Name (If Applicable) a	nd Established Name:	Famcicl	lovir Tablets 125 mg						I	(write in)					
Selling Unit NDC:	31722-706-30		Unit of Use NDC:	-	31722-706-30		33172270	6308	Note	s					
UDI			CVX Code:			MVX Code:									
Description:	Famciclovir Tablets 12	25 mg							Is thi	s product to be shippe	ed to customers on i	ce?		No	1
Is this product to be shipped to customers on dry ice? No															
Active Ingredient(s): Famciclovir, USP															
										erature excursion qu	lestions:				
URL for Additional Product Inform Address:		w.camberpharma.	.com			Address 2:			Nam			Soma Raju	0		
City:	800 Centennial Ave, S Piscataway	suite i			State:	NJ	Zip: 0	00E1	Num	ip E-mail:		732-529-042	is ieterousa.cor		
Key Contact:	Customer Service					customerservice@camberpharma.com					<u>somaraju er</u>	leterousa.con	<u>u</u>		
Phone Number:	1-866-827-3647				Fax:	732-562-8788	<u> </u>		c. Special regulation	ns for product in any	/ states?			No	1
Product Therapeutic Classification		cleoside analog D	NA polymerase inhibitor							ial returns requiremen				No	1
			1.7												1
	ADDITIONA	L PRODUCT INF				PRODUCT	DESCRIPT	ION INFORMATION	d. Store product (u	nit of sale) upright?				No	1
The product is?			Is the Product	Direct-Ship C	nlv					ect product (unit of s	ale) from light?			No	1
a legend device?	No		Is the Product	Unit of Use			30	ct	e. Shelf life:	Sor product (unit of S	ale, nom nymr			24	Months
if yes, enter class #	110		Orphan Drug Status			Size:	50			I shelf life at launch	(if different):				Months
a product kit?	No					Strength:	12	5 mg			(
if yes, list NDCs of			FDA Approval Status			Suengui.					ORDER INFORM	NATION			
component parts						Dosage For	m: Fili	m coated tablet							
reverse numbered?	No									of Sale			NDC selling	unit?	
co-licensed?	No		Allergens Present				-		2			1 Bottle of 3			
latex-free?	Yes	-	Lao	ctose		Product Sha	ape: Ro	ound, biconvex		Box/Carton		(Write-in, e.	g. 1 Box of 10) Vials)	
preservative-free? correctional institution block?	Yes						0	f white		Ampule Glass		Minimum or	der quantity	,	Yes
opioid?	No					Product Col	lor:	i white		Tube		Minimum of	uer quantity	r	Tes
Cannabinoid?	No		Country of Origin	India			Del	bossed with 'I' on one side		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u			jj			Product Imp		d '50' on the other side		Vial Liquid Multi		If Yes, how	many of whi	ch package t	type?
hospital scanning?			Is this product covered u	inder the						Vial Powder Sgl			Each		
If Unit Dose, indicate NDC here:			Trade Agreements Act (TAA)?	No					Vial Powder Multi			Inner/Carton	/Pack	
										Other: Write In			Case		
			FOR GENERIC DRUG PR	ODUCTS											
					Au	thorized Generic		ized Generic, other		PI	HARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AB						section fie	elds are not applicable	Rec. sell unit to cu	stomer?	_	Rx billing u	nit to pharma	acy:	
II. Generic Equivalent to What Bra	nd?: Far	mvir											Each		
			CHAIN SECURITY ACT						(Write-in, e.g. 1 Via)			Gram Milliliter		
		DRUG SUPPLY	CHAIN SECURITY ACT	(DSCSA) INFOR	MATION								Milliliter		
Does supplier meet DSCSA definit	tion of manufacturer?		Yes	_	GLN:	0331722498975				ITE	M AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			No	-		100000									
If yes, select exemption:					GCP:						Dimensi	ions (US msm	nts.)	Volume	Saleable #
Other exemption - Write in:									1	Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		If ves, was or	iginal product pur	rchased		Item/Each:		1				
Is product sold by manufacturer's	exclusive distributor?	2	Yes	_	direct from m					0.07	1.5	1.5	3	6.75	1
Has FDA granted waiver/exception		ct?	No		Provide source	ce manufacturer fo	or repackag	ged product	Box/Carton/Bundle	/					
If yes, attach documentation from	m FDA.								Inner Pack:						
									Case:	3.9	12.5	9.5	4.5	534.38	48
		GTIN	I AND HIBCC PRODUCT I	NFORMATION					Pallet:						
Saleable Unit of Measure	Salaa	ble Quantity	HIBCC		CTU	N-14		Jnit of Use GTIN-14	Pallet:						
x Item/Each	Salea	1	півсс			31722706308		0331722706308	L						
Box/Carton/Bundle/Inner Pack			00001722700000					(WHOLESALER USE ONLY:						
X Case		48			203	31722706302									
Pallet									Regular Cost			Vendor #:			
									Invoice Cost (WAC	(\$)	\$10.95	Whsl. Code			
							_			= = 0.0		Fineline Co	de:		
							_		As of date:	7/7/2017		4			
μ			August and a second second second		0)					I DADCODE					
	ormation on page 2.		Attach copy of SAFETY DA	ATA SHEET (SD	S) or non haza										
						See new p. 3 for	r Designate	d Drop Ship Only.	Sign	ature:					

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

Version 2021 For Designated Drop Ship Only Products, Please Use Page 3							
MATERIAL HA	ZARD CLASSIFICATION and TRANSPORTATION						
Is this product (check all that apply): a. Cytotoxic? 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? 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? No Is this product regulated for shipment by DOT? (if yes, answer a-e below and provide SDS) a. UN/Identification Number No No No No No No No N	Does the product have an Aerosol class? If yes, identify NFPA Storage Level: No NFPA Storage Level: Is the product a NIOSH hazardous drug? Is the product a NIOSH hazardous drug? No If yes, indicate which: If yes, indicate which:						
a. On/definition for holder b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard? Is this product regulated for shipment by IATA? No	Hazardous Waste Identification EPA Hazardous Waste Code: Waste Code:						
(if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? No 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) Special Permit; DOT-SP	REMS: No REMS Program Manager Name: Phone: Supplier Manages REMS registry exclusively: Phone: Wholesale distributor support: DEA #: Provider Name: DEA #: Site Enrollment Number assigned NCPDP#: by Supplier: NPI #:						
Special Provision (listed in Column 7 of 49 CFR 172.101); SP#ADD'L STORAGE INFORMATION	No Registry Program Contact Name: Comments						
Is the Product Controlled Substance? Controlled Substance? No Listed Chemical (List I or II) No ARCOS Reportable? No If yes, indicate which: Schedule No. Is it a scheduled listed chemical product?: No	RETURN INSTRUCTIONS Contact tel. # if product received damaged: 1-866-827-3647 Is product returnable for credit: Yes						
CLASS OF TRADE RESTRICTION: 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: No Restricted to hospital, clinics, and physician offices only: No Restricted from US territories? (explain in comments) No Comments: No	Special regulations or returns requirements for this product in certain states? No If so, which states? Other requirements? Comments?						
MISCELLANE	OUS NOTES and/or Image of Product Barcode:						



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

Version 2021 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 b. Autofax c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Name: Phone:	Purchase order daily receipt cut off time by supplier Cut off time: 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 Designated Drop Ship Fees:	Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each order: Drop Ship service fee billed with each order: Drop Ship miscellaneous fees billed: Comments:	Overnight receipt available: Image: Comparison of the co
Class of Trade Restriction:	PO Receipt Cut off time:
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	Saturday Overnight receipt available: PO Receipt Cut off time: Order receipt method: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:
Other Data Information Required to Process PO:	Return Instructions
Patient Procedure Date:	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?