

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

Version 2021							Introduction	Type: Post Launch Cha	ange	x	Final Version			Date:	6/1/2	2024
				PRODUCT INFORMA	TION						SPECIAL HAN	DLING AND STOP	RAGE REQUIR	REMENTS*	m	
Company Name: Camber Pharmaceuticals, Inc. Application:						a. Temperature – Indicate the USP temperature range for this product.										
Application Number for NDA/AN			device):		203	3834					ature Range	Controlled Room		and 25 C (68	° – 77° F)	
Medical Device Class, if applicat	ole:										-					
DUNS:	11-856-3719									Other Te	mperature Range F	Requirement	Store betwee	en 15° and 25	i° C.	
Propriet Is the product a CA Prop 6		cant?	Acyclovir	Tablets, USP 800 mg							ite in)					
Selling Unit NDC:	31722-778-01			Unit of Use NDC:			UPC:	331722778015		Notes						
UDI				CVX Code:			MVX Code:									
Description:	Acyclovir Tablets,	, USP 800 mg									oduct to be shipped				No	
Is this product to be shipped to customers on dry ice? No																
Active Ingredient(s): Acyclovir, USP b. Contact for temperature excursion questions:																
URL for Additional Product Information: www.camberpharma.com							b. Contact for tempera Name:	ture excursion que	estions:	Soma Raju						
Address:	800 Centennial A			<u>.</u>			Address 2:			Number	:		732-529-042	3		
City:	Piscataway State:				NJ	NJ Zip: 08854 Group E-mail:			-mail:		somaraju@he					
Key Contact:	Customer Service						camberpharma.com									
Phone Number:	1-866-827-3647	1-866-827-3647 Fax: 75				732-562-8788					states?			No		
Product Therapeutic Classification	Product Therapeutic Classification: Synthetic nucleoside analogue antiviral Special returns requirements for this product? No															
	ADDITI	ONAL PRODU					PRODUCT	DESCRIPTION INFORMATIO	ON	d. Store product (unit					No	
The product is?				Is the Product	Direct-Ship C	Only					product (unit of sa	le) from light?			No	
a legend device?		No		Is the Product	Neither		Size:	100 ct		e. Shelf life:					24	Months
if yes, enter class # a product kit?		No		Orphan Drug Status				800		Initial sl	nelf life at launch (	if different):				Months
if yes, list NDCs of		INO		FDA Approval Status			Strength:	800 mg				ORDER INFORM	ATION			
component parts			1	i DA Appioval otatus				Tablet				ONDER IN ON				
reverse numbered?		No	I.				Dosage For	n:		Unit of S	Sale		What is the	NDC selling	unit?	
co-licensed?		No		Allergens Present						x	Bottle		1 Bottle of 10	00 Tablets		
latex-free?		Yes					Product Sha	Oval, biconxex			Box/Carton		(Write-in, e.	g. 1 Box of 1	) Vials)	
preservative-free?		Yes					. roudor one	•			Ampule					
correctional institution block?		No					Product Col	Blue			Glass		Minimum or	der quantity	?	Yes
opioid? Cannabinoid?		No		Country of Origin	India			Debossed with 'J' on one	nido		Tube Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	init doce for	No		Country of Origin	India		Product Imp	rint: and '50' on the other side	side		Vial Liquid Sgi		If Yes how	many of whi	ch package t	vne?
hospital scanning?	init dose toi			Is this product covered u	under the						Vial Powder Sol			Each	cii package i	ype:
If Unit Dose, indicate NDC here:				Trade Agreements Act (		No					Vial Powder Multi			Inner/Carton	/Pack	
											Other: Write In			Case		
			FC	DR GENERIC DRUG PR	ODUCTS											
						Au	thorized Generic	*If Authorized Generic, othe				ARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AB							section fields are not applic	cable	Rec. sell unit to custor	ner?	-	Rx billing u		acy:	
II. Generic Equivalent to What Brand?: Zovirax							Allelia in a stabiliation				Each Gram					
		DRUG		HAIN SECURITY ACT						(Write-in, e.g. 1 Vial)				Gram Milliliter		
		DIGOG	DOITEIC		BOOCK) IN OF									Willinger		
Does supplier meet DSCSA definit	tion of manufactu	rer?		Yes		GLN:	0331722498975				ITEN	I AND PACKING II	NFORMATION	N		
Is product exempt from DSCSA?				No												
If yes, select exemption:						GCP:						Dimensi	ions (US msm	nts.)	Volume	Saleable #
Other exemption - Write in:											Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?				No	_		riginal product pur	chased		Item/Each:	0.31	2.6	2.6	4.5	30.42	1
Is product sold by manufacturer's				Yes No	_	direct from m		a several sever		Box/Carton/Bundle/						
Has FDA granted waiver/exception If yes, attach documentation from		roduct?		INU		Provide sour	ce manufacturer to	r repackaged product		Inner Pack:						
										Case:						10
			GTIN A	ND HIBCC PRODUCT I	NFORMATION						4.5	10.75	8.5	5.5	502.56	12
										Pallet:						
Saleable Unit of Measure	S	Saleable Quant	ity	HIBCC			N-14	Unit of Use GTIN-1	14							
X Item/Each		1				003	31722778015						_			×
Box/Carton/Bundle/Inner Pack		40				0.00	04700770040	-		COS	T INFORMATION			WHOLESAL	ER USE ONL	r:
X Case Pallet		12				203	31722778019	-		Regular Cost			Vendor #:			
										Invoice Cost (WAC) (\$		\$21.19	Whsl. Code	#:		
												ψ21.10	Fineline Cod			
	1									As of date:	4/15/2024		1			
													1			
μ													I			
			At	tach copy of SAFETY DA	ATA SHEET (SD	S) or non haza		INSERT, LABEL AND PHOT								
*Please provide any additional information on page 2.     See new p. 3 for Designated Drop Ship Only.     Signature:																

## HDA🔾

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

Version 2021 For De	signated Drop Ship Only Products, Please Use Page 3						
MATERIA	L HAZARD 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? N	X     Organic     Corrosive       Inorganic     Oxidizer       Steroid/Androgen     Contact Hazard						
c. Contact Hazard?     A. Does this product require special clean-up instructions?     (If yes, attach SDS with special instructions.)     e. Does the product contain DEHP?     Is this product regulated for shipment by DOT?     (if yes, answer a-e below and provide SDS)     a. UN/Identification Number	o identify NFPA Storage Level: NFPA Storage Level:						
b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	Hazardous Waste Identification EPA Hazardous Waste Code: Waste Characteristics						
Is this product regulated for shipment by IATA? (if yes, answer a-e below and provide SDS) a. UN/Identification Number	REMS or REGISTRY RESTRICTIONS						
a. Orviterinitation Name b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	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	o     Med Guide Required     No       Limited Distribution Requirement     Image: Comments / Details: (For example, iPledge program?)     Image: Comment is a comment in the comment is a comment is a comment is a comment is a comment in the comment in the comment is a comment in the comment in the comment in the comment in the comment is a comment in the comme						
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:     No       REMS Program Manager Name:     Phone:       Supplier Manages REMS registry exclusively:     Phone:       Wholesale distributor support:     Provider Name:       Provider Name:     DEA #:       Site Enrollment Number assigned     NCPDP#:       by Supplier:     NPI #:						
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101);	Comments						
ADD'L STORAGE INFORMATION	Registry:     No       Registry Program Contact Name:     Phone:       Comments						
Is the Product Controlled Substance? No Controlled Substance Code	RETURN INSTRUCTIONS						
Controlled by State(s)?       No       Listed Chemical (List I or II)       N         ARCOS Reportable?       No       If yes, indicate which:       If yes, indicate which:       If yes, indicate which:         Schedule No.       Is it a scheduled listed chemical product?:       N         CLASS OF TRADE RESTRICTION:	Contact tel. # if product received damaged: 1-866-827-3647						
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices	contact - customerservice@camberpharma.com						
Restricted to retail pharmacy only:       N         Restricted to hospital, clinics, and physician offices only:       N         Restricted from US territories? (explain in comments)       N         Comments:       Image: Comments in the second seco	Special regulations or returns requirements for this product in certain states? No If so, which states? Other requirements? Comments?						
	ANEOUS 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?