

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

Version 2021						Introduction 1	Type: P	ost Launch Change	x	Final Version			Date:	6/23/	2024	
			PRODUCT INFORMA	TION						SPECIAL HAN	DLING AND STOR	AGE REQUIR	REMENTS*			
Company Name: Camber Pharmaceuticals, Inc.				Application: ANDA			a. Temperature – Indicate the USP temperature range for this product.									
	/ANDA/BLA (drug); PMA/510(k)(med device): 203053 Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)															
Medical Device Class, if applicab	ole:															
DUNS:	11-856-3719								Other	Temperature Range F	Requirement					
Proprietary Name (If Applicable) and		e: Efavirer	nz, Emtricitabine and Tenol							write in)						
Selling Unit NDC:	31722-736-30		Unit of Use NDC:		31722-736-30		3317227363	605	Notes							
UDI			CVX Code:			MVX Code:										
Description:	Efavirenz, Emtricitabi	ine and Tenofovir I	Disoproxil Fumarate Tablets	s 600 mg/200 mg	g/300 mg					product to be shipped				No		
Is this product to be shipped to customers on dry ice? No																
Active Ingredient(s): Efavirenz, USP, emtricitabine, and tenofovir disoproxil fumarate b. Contact for temperature excursion questions:																
URL for Additional Product Inform	r Additional Product Information: www.camberpharma.com								b. Contact for tempe Name:		estions:	Soma Raiu				
Address:	300 Centennial Ave, Suite 1				Address 2:			Number:			732-529-0423					
City:					NJ	Zip: 088	54	Group E-mail: somaraju@heterousa.com					<u>n</u>			
Key Contact:	Customer Service	Customer Service Email:				customerservice	erservice@camberpharma.com									
Phone Number:	1-866-827-3647	827-3647 Fax: 732				732-562-8788	32-562-8788			c. Special regulations for product in any states?				No		
Product Therapeutic Classification	n: Co	ombination HIV-1 a	antiviral						Specia	I returns requirement	s for this product?			No		
	ADDITION					DRODUCE										
	ADDITION	AL PRODUCT INF				PRODUCT	DESCRIPTIO	N INFORMATION	d. Store product (uni					No		
The product is?			Is the Product	Direct-Ship O	niy		0.5			t product (unit of sa	le) from light?			No		
a legend device?	N	0	Is the Product	Unit of Use		Size:	30 ct		e. Shelf life:	obolf life at launch (if different's			24	Months Months	
if yes, enter class # a product kit?	N	0	Orphan Drug Status				600 r	ng/200 mg/300 mg	initiai	shelf life at launch (if different):				Months	
if yes, list NDCs of		0	FDA Approval Status			Strength:	0001	11g/200 11g/300 11g			ORDER INFORM	IATION				
component parts						D	Film-	coated tablet								
reverse numbered?	N	0				Dosage Forr	n:		Unit o	Sale		What is the	NDC selling	unit?		
co-licensed?	N	0	Allergens Present						X	Bottle		1 Bottle of 30				
latex-free?		es	N	uts		Product Sha	Caps	ule		Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)		
preservative-free?	Ye									Ampule				•	No. 1	
correctional institution block? opioid?	N					Product Col	or: White	e to off-white		Glass Tube		Minimum or	der quantity	?	Yes	
Cannabinoid?	N		Country of Origin	India			Debos	sed with 'H' on one side		Vial Liquid Sgl						
If Unit Dose, is item bar coded to u		0	j			Product Imp	rint: and '1	28' 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		24	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												
Authorized Generic *If Authorized Generic, other								PHARMACY ORDER / BILL UNIT								
	40			_	Au	thorized Generic		s are not applicable	Dee cell unit to quet							
I. Orange Book Rating: AB Section fields are not applicable II. Generic Equivalent to What Brand?: Atripla								Rec. sell unit to cust	omerr	1	Rx billing u	Each	acy:			
II. Generic Equivalent to what Brai	iur.	пра							(Write-in, e.g. 1 Vial)		_		Gram			
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION																
Does supplier meet DSCSA definit	tion of manufacturer?	?	Yes		GLN:	0331722498975			-	ITEN	I AND PACKING I	NFORMATION	N			
Is product exempt from DSCSA?			No													
If yes, select exemption:					GCP:					Weight Lbs.		ons (US msm	'	Volume	Saleable	
Other exemption - Write in:			No		K	dalaal aas doot o	ahaaad 📃		How /Fast		Depth	Width	Height	(Cube)	#Pieces	
Is product repackaged? Is product sold by manufacturer's	exclusive distributor	2	Yes		If yes, was or direct from m	riginal product pur	cnased		Item/Each:	0.2	2	2	3.6	14.40	1	
Has FDA granted waiver/exception			No			ce manufacturer fo	or repackage	d product	Box/Carton/Bundle/							
If yes, attach documentation from		1							Inner Pack:							
									Case:	5.5	12.5	8.75	4.5	492.19	24	
		GTIN	I AND HIBCC PRODUCT I	NFORMATION												
Saleable Unit of Measure	0-1-				OT1	N-14	11-2	t of Use GTIN-14	Pallet:							
x Item/Each	Sale	able Quantity	HIBCC			N-14 31722736305		31722736305								
Box/Carton/Bundle/Inner Pack	00331722736305							CC	ST INFORMATION			WHOL <u>ESAL</u>	ER USE ONL	Y:		
X Case		24			203	31722736309										
Pallet									Regular Cost			Vendor #:				
									Invoice Cost (WAC) (\$)	\$120.00	Whsl. Code				
	_						-			0/40/0000		Fineline Co	de:			
	-						-		As of date:	2/18/2022						
 			Attach copy of SAFETY DA		S) or non haza		INSERT 1 A					!				
*Please provide any additional info	ormation on page 2		Auton copy of OAI ETT DA	(30	o, or non ndzd	See new p. 3 for			Signal							
. isase provide any additional line						556 new p. 5 10	Designated	and and and	Signa							

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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 H/	AZARD CLASSIFICATION and TRANSPORTATION						
Is this product (check all that apply): a. Cytotoxic? No 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? No d. Does this product require special clean-up instructions? No (If yes, attach SDS with special instructions.) e. Does the product contain DEHP? No Is this product regulated for shipment by DOT? No	Does the product have an Aerosol class? If yes, identify NFPA Storage Level: No NFPA Storage Level: Image: Storage Level: Is the product a NIOSH hazardous drug? No						
(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?	If yes, indicate which: Hazardous Waste Identification EPA Hazardous Waste Code: Waste Characteristics						
Is this product regulated for shipment by IATA? No							
(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?	No If Yes, is it managed with a pharmacy registry? Website URL:						
Is the product restricted for air shipment? If so, indicate restriction: No 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: Provider Name: Provider Name: DEA #: Site Enrollment Number assigned NCPDP#: by Supplier: NPI #:						
Special Provision (listed in Column 7 of 49 CFR 172.101);	Devision						
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) No ARCOS Reportable? No If yes, indicate which:	Contact tel. # if product received damaged: 1-866-827-3647						
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 - customerservice@camberpharma.com						
Restricted to retail pharmacy only: No							
No No Restricted from US territories? (explain in comments) No	Special regulations or returns requirements for this product in certain states? No If so, which states? Other requirements? Comments?						
Comments:	EOUS NOTES and/or Image of Product Barcode:						
	rz (EFV), a non-nucleoside reverse transcriptase inhibitor, and emtricitabine (FTC) and tenofovir disoproxil fumarate (TDF), both HIV-1						



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?