

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

Version 2021						Introduction	Туре:	New Item	x	Final Version			Date:	3/20/	2024	
			PRODUCT INFORMA	TION						SPECIAL HAN	DLING AND STOP	RAGE REQUIF	REMENTS*	•		
Company Name: Camber Pharmaceuticals, Inc.				Application: ANDA		a. Temperature – Indicate the USP temperature range for this product.										
Application Number for NDA/ANI	DA/BLA (drug); PMA/510	k)(med devic	e):	21	7748					rature Range	Controlled Room		and 25 C (68	8° – 77° F)		
Medical Device Class, if applicable:																
	11-856-3719									Temperature Range F	Requirement					
Proprietary Name (If Applicable) and		Allopur	inol Tablets, USP 300 mg		1					write in)						
Selling Unit NDC: UDI	31722-253-10		Unit of Use NDC: CVX Code:			UPC: MVX Code:	33172225	3109	Notes							
-			CVA Code.			WIVA Code.						-				
Description:	Allopurinol Tablets, USP	300 mg								product to be shipped				No No		
Active Ingredient(s): Allopurinol, USP																
b. Contact for temperature excursion questions:																
URL for Additional Product Inform		amberpharma.							Name			Soma Raju				
Address:	1031 Centennial Ave (and) 800 Centennial Ave, Suite 1			Address 2:				Number:			732-529-0423					
City:	Piscataway				State:	NJ				E-mail:	somaraju@heterousa.com					
Key Contact: Phone Number:	Customer Service Email: 1-866-827-3647 Fax:				customerservice@camberpharma.com 732-562-8788			c. Special regulations for product in any states?				No				
Product Therapeutic Classification		ine oxidase inl	hibitor		Tax.	132-302-0100	102 002 0100			Special returns requirements for this product?				No		
Froduct merapeutic classification	. Xana								Opecia	i returns requirement	s for this product?			INO		
	ADDITIONAL I	PRODUCT INF				PRODUCT	DESCRIPTI	ION INFORMATION	d. Store product (uni	t of sale) upright?				No		
The product is?			Is the Product	Direct-Ship C	Dnly					t product (unit of sa	le) from light?			No		
a legend device?	No		Is the Product	Neither		Size:	100	00 ct	e. Shelf life:		,			24	Months	
if yes, enter class #			Orphan Drug Status			Size:				shelf life at launch (if different):				Months	
a product kit?	No					Strength:	300	0 mg								
if yes, list NDCs of			FDA Approval Status				T-1	h l a c				MATION				
component parts reverse numbered?	No					Dosage For	m: ^{Tar}	blet	Unit o	Sala		What is the	NDC selling	unit?		
co-licensed?	No		Allergens Present						x	Bottle		1 Bottle of 10		unit.		
latex-free?	Yes					Product Sha	Ro	und		Box/Carton		(Write-in, e.g		0 Vials)		
preservative-free?	Yes					FIGUUCESIN				Ampule						
correctional institution block?	No					Product Col	or: Wh	nite to off white		Glass		Minimum or	der quantity	?	Yes	
opioid?	No			La d'a			Dut	bossed with "U" and "6" on		Tube						
Cannabinoid? If Unit Dose, is item bar coded to u	No		Country of Origin	India		Product Imp		e side and functional		Vial Liquid Sgl Vial Liquid Multi		If Yes, how I	many of whi	ich nackade f	wne?	
hospital scanning?			Is this product covered u	nder the			sco	ored line with "H" on the		Vial Powder Sql			Each	ich package i	iype :	
If Unit Dose, indicate NDC here:			Trade Agreements Act (1		No		oth	er side.		Vial Power Multi			Inner/Cartor	n/Pack		
										Other: Write In			Case			
			FOR GENERIC DRUG PR	ODUCTS												
						uthorized Generic	*16. 4. 44	zed Generic, other		DL	ARMACY ORDER					
	10					unonzed Generic		elds are not applicable	Rec. sell unit to cust		IARIMACT ORDER					
	I. Orange Book Rating.								Rec. sell unit to cust	omer ?	1	Rx billing ur		acy:		
II. Generic Equivalent to What Brand?: Zyloprim							(Write-in, e.g. 1 Vial) Gram									
		ORUG SUPPL	Y CHAIN SECURITY ACT (DSCSA) INFOR	RMATION				(,				Milliliter			
Does supplier meet DSCSA definit	ion of manufacturer?		Yes	_	GLN:	0331722000000				ITEN	I AND PACKING II	NFORMATION	N			
Is product exempt from DSCSA?			No													
If yes, select exemption:					GCP:					Weight Lbs.		ions (US msm		Volume	Saleable #	
Other exemption - Write in: Is product repackaged?			No		If yoe was a	riginal product pur	chased		Item/Each:	-	Depth	Width	Height	(Cube)	Pieces	
Is product repackaged?	exclusive distributor?		Yes	-	direct from n		ulaseu		nenveach:	1.4	3.35	3.35	7	78.5575	1	
Has FDA granted waiver/exception			No			rce manufacturer fo	or repackag	jed product	Box/Carton/Bundle/							
If yes, attach documentation from	n FDA.								Inner Pack:							
									Case:	17.5	14	10.5	8	1176	12	
		GTI	N AND HIBCC PRODUCT I	NFORMATION					Dellet							
Saleable Unit of Measure	Saleable	Quantity	HIBCC		CT.	IN-14		Init of Use GTIN-14	Pallet:							
x Item/Each		1	Півсо			331722253109										
Box/Carton/Bundle/Inner Pack									CC	ST INFORMATION		<u> </u>	WHOLESAL	ER USE ONL	Y:	
X Case	1	2			203	331722253103										
Pallet							-		Regular Cost	•		Vendor #:				
	-						-		Invoice Cost (WAC) (\$)	\$177.56	Whsl. Code Fineline Cod				
									As of date:	2/23/2024		ineime coo	u c .			
							-					1				
							-									
			Attach copy of SAFETY DA	TA SHEET (SE	S) or non haza	ard letter, PACKAGE	E INSERT, L	ABEL AND PHOTO OF P	RODUCT PACKAGING	nd BARCODE.						
*Please provide any additional info	ormation on page 2.							d Drop Ship Only.	Signa							
L																

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

·	nated Drop Ship Only Products, Please Use Page 3
MATERIAL H	HAZARD 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? No Is the product a CA Prop 65 reproductive toxicant? No Does the product label bear a CA Prop 65 warning? C. Contact Hazard? (If yes, attach SDS with special instructions? (If yes, attach SDS with special instructions.) e. Does the product contain DEHP? No	x Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard Does the product have an Aerosol class? If yes, identify NFPA Storage Level: No NFPA Storage Level: Image: Contact Hazard
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 c. DOT Hazard Class	Is the product a NIOSH hazardous drug? No If yes, indicate which: Hazardous Waste Identification
d. Packing Group e. Inhalation Hazard?	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 b. Proper Shipping Name	REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? No
c. DOT Hazard Class d. Packing Group e. Inhalation Hazard? Is the product restricted for air shipment? If so, indicate restriction: Passenger	If Yes, is it managed with a pharmacy registry? Website URL: Med Guide Required Limited Distribution Requirement
Cargo Passenger & Cargo Is this a reportable quantity? No RQ Threshold:	REMS: No REMS Program Manager Name: Phone:
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)	Supplier Manages REMS registry exclusively:
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
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: Image: Control of the state of	Contact tel. # if product received damaged: Is product returnable for credit: 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 Restricted to hospital, clinics, and physician offices only: 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:	
MISCELLAN	IEOUS NOTES and/or Image of Product Barcode:



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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?