

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

Version 2024						Introduction Type:	New Item		x Final Version			Date:	11/20	0/2024
			PRODUCT INFORMA	TION					SPECIAL HAN	IDLING AND STOP	RAGE REQUI	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc.				Application: ANDA		a. Temperature – Indicate the USP temperature range for this product.								
Application Number for NDA/AN	IDA/BLA; PMA/510	(k): 217748				NDA 505(b) Type:	NOT APPLICABLE		Temperature Range	Controlled Room		and 25 C (68	° – 77° F)	
Medical Device Class, if applicat	ble:							T .						
DUNS:	11-856-3719							_	Other Temperature Range	Requirement				
Proprietary Name (If Applicable) a		ame: Allopurir	nol Tablets, USP 300 mg		I				(write in)					
Selling Unit NDC:	31722-253-10		Unit of Use NDC: CVX Code:			UPC: 331 MVX Code:	722253109		Notes					
UDI			CVX Code:			WIVA Code.		1						
Description:	Allopurinol Tablet	s, USP 300 mg							Is this product to be shippe				No	-
Active Ingredient(s):		Allopurinol, USP							Is this product to be shippe	a to customers on a	ary ice?		No	
Active ingredient(s).		Allopulliloi, 001						b. Contact for	temperature excursion qu	estions:				
URL for Additional Product Inform	nation:	www.camberpharma.	com						Name:		Soma Raju			
Address:	800 Centennial Av	ve, Suite 1				Address 2:		1	Number:		732-529-042	23		
City:	Piscataway				State:		p: 08854	Group E-mail: somaraju@heterousa.com				<u>n</u>		
Key Contact:	Customer Service	9			Email:	customerservice@car	mberpharma.com					7		
Phone Number:	1-866-827-3647	Marath la sa shi da sa la b	11. 11		Fax:	732-562-8788			ulations for product in any				No	-
Product Therapeutic Classification	n:	Xanthine oxidase inh	IDITOF						Special returns requirement	ts for this product?			No	_
		ONAL PRODUCT INFO	ORMATION			PRODUCT DESC	CRIPTION INFORMATION		ct (unit of sale) upright?				No	Т
The preduct is 2	Abbitt			Direct Ship C	nlu	TROBOOT BED				ala) faam links?				-
The product is? a legend device?		No	Is the Product Is the Product	Direct-Ship C Neither	riiy		1000 ct	e. Shelf life:	Protect product (unit of s	ale) from light?			No 24	Months
if yes, enter class #		INU	Orphan Drug Status	T Clurici		Size:	1000 Cl		Initial shelf life at launch	if different):			24	Months
a product kit?		No		1		Strongth	300 mg			,				1
if yes, list NDCs of			FDA Approval Status			Strength:				ORDER INFOR	MATION			
component parts						Dosage Form:	Tablet							
reverse numbered?		No				•			Unit of Sale			NDC selling	unit?	
co-licensed? latex-free?		No Yes	Allergens Present				Round		x Bottle Box/Carton		1 Bottle of 1	g. 1 Box of 1) \/iale)	
preservative-free?		Yes				Product Shape:	Round		Ampule		(11111111111111111111111111111111111111	g. I Dox of h	J viais)	
correctional institution block?		No				Product Color:	White to off-white		Glass		Minimum o	rder quantity	?	Yes
opioid?		No				Product Color:			Tube					
Cannabinoid?		No	Country of Origin	India		Product Imprint:	Debossed with 'U' and '6' on one side and functional scored line with 'H' on		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	unit dose for						the other side		Vial Liquid Multi				ch package t	type?
hospital scanning? If Unit Dose, indicate NDC here:			Is this product covered u Trade Agreements Act (1		No				Vial Powder Sgl Vial Powder Multi		12	Each Inner/Carton	/Dook	
Il Offit Dose, indicate NDC fiele.				~~):	NU				Other: Write In			Case	Fack	
			FOR GENERIC DRUG PR	ODUCTS										
				000010										
					Au	uthorized Generic *If A	Authorized Generic, other		PI	ARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AB					sec	tion fields are not applicable	Rec. sell unit to customer?			Rx billing unit to pharmacy:			
II. Generic Equivalent to What Bra	and?:	Zyloprim									Each			
								(Write-in, e.g.				Gram		
		DRUG SUPPLY	CHAIN SECURITY ACT (DSCSA) INFOR	MATION			HCPCS J-Cod	e:			Milliliter		
Does supplier meet DSCSA defini	tion of monufactur	ror?	Yes	_	GLN:	0331722498975			ITE	AND PACKING I		M		
Is product exempt from DSCSA?			No	-	GLN.	0331722490975			1161	AND FACKING I		N		
			-		CCP.			1		Dimens	ions (US msr	nte)	Volume	Coloriste #
If yes, select exemption: Other exemption - Write in:					GCP:			1	Weight Lbs.	Dimens	Width	nts.) Height	Volume (Cube)	Saleable # Pieces
Is product repackaged?			No		If yes, was o	riginal product purchas	ed	Item/Each:						
Is product sold by manufacturer's	exclusive distribute	utor?	Yes	-	direct from m				1.4	3.35	3.35	7	78.56	1
Has FDA granted waiver/exception	n/exemption for pr	roduct?	No		Provide sour	rce manufacturer for rep	backaged product	Box/Carton/Bu	undle/					
If yes, attach documentation from	m FDA.							Inner Pack:						
		CTIN	AND HIBCC PRODUCT I	FORMATION				Case:	17.5	14	10.5	8	1176.00	12
		GTIN	AND HIBCC PRODUCT IN	NFORMATION				Pallet:						
Saleable Unit of Measure	RFID tag(Y/N)	Saleable	HIBCC		GTI	IN-14	Unit of Use GTIN-14	r allet.						
		Quantity						1		1				1
x Item/Each	N	1			003	331722253109								
Box/Carton/Bundle/Inner Pack									COST INFORMATION			WHOLESAL	ER USE ONL	LY:
X Case	N	12			203	331722253103								
Pallet								Regular Cost Invoice Cost (\$477 FO	Vendor #: Whsl. Code	#-		
								invoice cost (WAC) (\$)	06.111¢	Fineline Co			
								As of date:	3/18/2024					
											1			
			Attach copy of SAFETY DA	TA SHEET (SD	S) or non haza		ERT, LABEL AND PHOTO OF F							
*Please provide any additional info	formation on page		Attach copy of SAFETY DA	TA SHEET (SD	S) or non haza		ERT, LABEL AND PHOTO OF F ignated Drop Ship Only.		GING and BARCODE. Signature:					

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

Version 2024 For Design	ated Drop Ship Only Products, Please Use Page 3				
MATERIAL H	AZARD CLASSIFICATION and TRANSPORTATION				
MATERIAL H Is this product (check all that apply): a. Cytotoxic? No b. CA Prop. 65 Carcinogen or Reproductive Toxicant? No 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? No c. Contact Hazard? No d. Does this product require special clean-up instructions? No (If yes, attach SDS with special instructions.) No e. Does the product contain DEHP? No Is this product regulated for shipment by DOT? Yes (if yes, answer a-e below and provide SDS) UN3077 b. Proper Shipping Name Environmentally hazardous substances, solid, n.o.s. (ALLOPURNOL), MARINE POLLUTANT	AZARD CLASSIFICATION and TRANSPORTATION				
c. DOT Hazard Class 9	Hazardous Waste Identification				
d. Packing Group					
e. Inhalation Hazard? No	EPA Hazardous Waste Code: Waste Characteristics				
Is this product regulated for shipment by IATA? Yes					
(if yes, answer a-e below and provide SDS)	REMS or REGISTRY RESTRICTIONS				
a. UN/Identification Number UN3077					
b. Proper Shipping Name Environmentally hazardous substance, solid, n.o.s. (ALLOPURINOL) c. DOT Hazard Class 9	Is there a REMS on this product? No				
c. DOT Hazard Class 9 d. Packing Group III	If Yes, is it managed with a pharmacy registry? Website URL:				
e. Inhalation Hazard?	Website ONL.				
	Med Guide Required				
Is the product restricted for air shipment? If so, indicate restriction: No Passenger	Med Guide Required No Limited Distribution Requirement Image: Comparison of the second seco				
Cargo	Common Drains (For example, iPledge program?)				
Passenger & Cargo					
Is this a reportable quantity? No	REMS: No				
RQ Threshold:	REMS Program Manager Name: Phone:				
Is this a marine pollutant? Yes	Supplier Manages REMS registry exclusively:				
Is this product shipped utilizing an authorized DOT exception or Special Permit?	Wholesale distributor support:				
No (if yes, identify method below)	Provider Name: DEA #:				
Limited Quantity	Site Enrollment Number assigned NCPDP#:				
Consumer Commodity, ORM-D	by Supplier: NPI #:				
Small Quantity (49 CFR 173.4)					
Special Permit; DOT-SP	Comments				
Special Provision (listed in Column 7 of 49 CFR 172.101); SP#	Registry: No				
3F#					
ADD'L STORAGE INFORMATION	Registry Program Contact Name: Phone: Comments				
	Cumments				
Is the Product	RETURN INSTRUCTIONS				
Controlled Substance? No Controlled Substance Code Controlled by State(s)? No Listed Chemical (List I or II) No	KETOKA INSTRUCTIONS				
ARCOS Reportable? No If yes, indicate which:	Contact tel. # if product received damaged: 1-866-827-3647				
Schedule No. Is it a scheduled listed chemical product?: No					
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	Special regulations or returns requirements for this				
Restricted to hospital, clinics, and physician offices only: No	product in certain states? No				
Restricted from US territories? (explain in comments) No	If so, which states? Other requirements? Comments?				
Comments:					
MISCELLAN	EOUS NOTES and/or Image of Product Barcode:				



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Version 2024	FOR DESIGNATED DROP SHIP PRODUCT ONLY - if r	ot a designated drop ship, do not complete.					
Order Method for Des	signated 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 Expedited Freight Charges o		Purchase order daily receipt cut off time by supplier Cut off time: Shipping lead time of PO: Hours Da Ships same day for next day receipt: Ships for second day receipt: Ships regular ground for 3-10 days receipt: Overnight and Priority Overnight PO Processing	ays				
Expedited freight fees billed with each order:		Overnight receipt available:					
Drop Ship service fee billed with each order:		PO Receipt cut off time:					
Drop Ship miscellaneous fees billed:		Days of week overnight is available: Monday					
Comments:	s only:	Days of week overnight is available: Tuesday Tuesday Wednessi Priority Overnight receipt available: Friday PO Receipt Cut off time: PO Receipt Cut off time: Order receipt method: Phone: Fax: EDI: Overnight Fees apply: Policity	/ day				
Other Data Informati	ion Required to Process PO:	Return Instructions					
Patient Procedure Date: Physician Name: Physician/Clinic Phone # Physician State License # Physician/Clinic DEA #: Physician/Clinic Specialty:		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?					
Miscell	aneous Notes:						
		ADDITIONAL INFORMATION					
		Is product order for scheduled patient procedure? Is product order for restocking purposes?					