

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

Version 2024						Introduction Type:	New Item		x Final Version			Date:	11/15	5/2024
			PRODUCT INFORMA	TION					SPECIAL HAN	DLING AND STOP	RAGE REQUI	REMENTS*		
Company Name:	Camber Pharmac	euticals, Inc.				Application:	ANDA	a. Temperature -	Indicate the USP tempe	erature range for t	his product.			
Application Number for NDA/AN	IDA/BLA; PMA/510	(k): 217748				NDA 505(b) Type:	NOT APPLICABLE		mperature Range	Controlled Room		and 25 C (68	° – 77° F)	
Medical Device Class, if applicat	ble:							T						
DUNS:	11-856-3719							01	her Temperature Range I	Requirement				
Proprietary Name (If Applicable) a		ame: Allopuri	nol Tablets, USP 300 mg						(write in)					
Selling Unit NDC:	31722-253-01		Unit of Use NDC: CVX Code:			UPC: 331 MVX Code:	722253017	No	otes					
UDI			CVX Code:			MIVA Code.					-			1
Description:	Allopurinol Tablet	s, USP 300 mg							this product to be shipped				No No	-
Active Ingredient(s):		Allopurinol, USP						IS	this product to be shipped	a to customers on t	iry ice?		INO	]
Active ingredient(s).								b. Contact for ter	nperature excursion qu	estions:				
URL for Additional Product Inform	nation:	www.camberpharma.	com						ime:		Soma Raju			
Address:	800 Centennial Av	ve, Suite 1				Address 2:			umber:		732-529-042			
City:	Piscataway				State:		<b>08854</b>	Gi	oup E-mail:		somaraju@	heterousa.cor	<u>n</u>	
Key Contact:	Customer Service	9			Email:	customerservice@can	hberpharma.com						N	1
Phone Number:	1-866-827-3647	Xanthine oxidase inh	ibitor		Fax:	732-562-8788			tions for product in any				No	-
Product Therapeutic Classification	n:	Aanthine oxidase inn	IDILOI		]			Sp	ecial returns requirement	is for this product?			No	
	ADDITI	ONAL PRODUCT INF				PRODUCT DESC	RIPTION INFORMATION	d. Store product	(unit of sale) upright?				No	1
The product is?			Is the Product	Direct-Ship C	nlv				otect product (unit of sa	ale) from light?			No	л 1
a legend device?		No	Is the Product	Neither	any		100 ct	e. Shelf life:	olect product (unit of sa	ile) ironi light?			24	Months
if yes, enter class #			Orphan Drug Status			Size:			itial shelf life at launch (	if different):			27	Months
a product kit?		No				Strength:	300 mg			-				
if yes, list NDCs of			FDA Approval Status			ou engui.				ORDER INFORM	ATION			
component parts		N.				Dosage Form:	Tablet		it of Colo		What is the	NDC selling		
reverse numbered? co-licensed?		No No	Allergens Present					0	nit of Sale x Bottle		1 Bottle of 1		unit?	
latex-free?		Yes	Allergens Fresent				Round		Box/Carton			.g. 1 Box of 1	) Vials)	
preservative-free?		Yes				Product Shape:			Ampule		(	· · · · · · · · · · · · · · · · · · ·		
correctional institution block?		No				Product Color:	White to off-white		Glass		Minimum o	rder quantity	?	Yes
opioid?		No				i roudor obiorr			Tube					
Cannabinoid?	unit dans for	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		K Vee herr	many of whi		4
If Unit Dose, is item bar coded to u hospital scanning?	unit dose for		Is this product covered u	nder the			the other side		Vial Liquid Multi Vial Powder Sgl			many of whi Each	ch package	type?
If Unit Dose, indicate NDC here:			Trade Agreements Act (		No				Vial Powder Multi		24	Inner/Carton	/Pack	
					I				Other: Write In			Case		
			FOR GENERIC DRUG PR	ODUCTS										
											_			
					Au		uthorized Generic, other	PHARMACY ORDER / BILL UNIT						
I. Orange Book Rating:	AB					Sec	tion fields are not applicable	Rec. sell unit to	customer?	-	Rx billing u	nit to pharma	acy:	
II. Generic Equivalent to What Bra	and?:	Zyloprim							( - 1)			Each		
		DRUG SUPPLY	CHAIN SECURITY ACT (	DSCSA) INFOR	MATION			(Write-in, e.g. 1 \ HCPCS J-Code:	(ial)			Gram Milliliter		
		5.000001121								1				
Does supplier meet DSCSA definit	ition of manufactur	rer?	Yes		GLN:	0331722498975			ITEN	I AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			No											
If yes, select exemption:					GCP:				Weight Lbs.	Dimensi	ons (US msr	nts.)	Volume	Saleable #
Other exemption - Write in:									weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No	_		riginal product purchase	ed	Item/Each:	0.18	1.87	1.87	3.22	11.26	1
Is product sold by manufacturer's Has FDA granted waiver/exception			Yes No	_	direct from n		ackaged product	Box/Carton/Bune	llo/					
If yes, attach documentation from			110		FIOVICE SOUL	rce manufacturer for rep	ackaged product	Inner Pack:	lie/					
								Case:	4.9	14.5	7.0	4.5	202.20	24
		GTIN	AND HIBCC PRODUCT II	NFORMATION					4.9	11.5	7.6	4.5	393.30	24
		<b>_</b>						Pallet:						
Saleable Unit of Measure	RFID tag(Y/N)		HIBCC		GTI	IN-14	Unit of Use GTIN-14							
x Item/Each	N	Quantity			003	331722253017								
Box/Carton/Bundle/Inner Pack									COST INFORMATION			WHOLESAL	ER US <u>E ONL</u>	Y:
X Case	N	24			203	331722253011								
A Outo								Regular Cost			Vendor #:			
Pallet														
								Invoice Cost (WA	AC) (\$)	\$21.00	Whsl. Code			
										\$21.00	Whsl. Code Fineline Co			
								Invoice Cost (WA	3/18/2024	\$21.00				
										\$21.00				
			Attach copy of SAFETY DA	TA SHEET (SD	S) or non haza	ard letter, PACKAGE INSI	ERT, LABEL AND PHOTO OF F	As of date:	3/18/2024	\$21.00				
			Attach copy of SAFETY DA	TA SHEET (SD	S) or non haza		ERT, LABEL AND PHOTO OF F gnated Drop Ship Only.	As of date:	3/18/2024	\$21.00				

## **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:



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

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?	