

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

Version 2021						Introduction Typ	e: Post Launch Change		x Final Version			Date:	6/23	3/2024
			PRODUCT INFORMA	TION					SPECIAL HAP	IDLING AND STOR	RAGE REQUI	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc. Application: ANDA							a. Temperature – Indicate the USP temperature range for this product.							
Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): 206574							Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)							
Medical Device Class, if applicable:														
DUNS:	11-856-3719							_	Other Temperature Range	Requirement				
Proprietary Name (If Applicable) a	and Established Na	me: Duta	steride Capsules 0.5 mg						(write in)					
Selling Unit NDC:	31722-131-90		Unit of Use NDC		31722-131-90		31722131902		Notes					
UDI			CVX Code:			MVX Code:								
Description: Dutasteride Capsules 0.5 mg Is this product to be shipped to customers on ice? No									1					
_									Is this product to be shippe				No	1
Active Ingredient(s):		Dutasteride												
								b. Contact fo	or temperature excursion qu	estions:				
URL for Additional Product Inform		www.camberpharr	ma.com					41	Name:		Soma Raju			
Address:	800 Centennial A	ve, Suite 1			State:	Address 2:	7:	-	Number:		732-529-042			
City: Key Contact:	Piscataway Customer Service				Email:	customerservice@c	Zip: 08854	Group E-mail: somara				heterousa.co	<u>m</u>	
Phone Number:	1-866-827-3647	•			Fax:	732-562-8788	amberphamia.com	c Special re	gulations for product in any	states?			No	٦
Product Therapeutic Classification		5 alpha-reductase	inhibitor		l ux.	702 002 0700		c. opeciai re	Special returns requirement				No	-
Troduct Therapeutic Glassification	//··	o dipria reductase	, in illibitor						Special returns requiremen	ts for this product:			140	_
	ADDITI	ONAL PRODUCT I	NEORMATION			PRODUCT DE	SCRIPTION INFORMATION	d Store pro	duct (unit of sale) upright?				No	٦
The weeduct is 2				Direct-Ship (Only	I NOSCO I DE		T a. otore pro		ala) faama liinka				1
The product is? a legend device?		No	Is the Product Is the Product	Unit of Use	Jilly		90 ct	e. Shelf life:	Protect product (unit of s	ale) from light?			No 24	Mantha
if yes, enter class #		INO	Orphan Drug Status	Offit of Ose		Size:	90 61	e. Shell life:	Initial shelf life at launch	if different).			24	Months Months
a product kit?		No	Orphan Drug Status				0.5 mg		illidai Sileli ille at laulicii	ii dillerent).				Months
if yes, list NDCs of		140	FDA Approval Status			Strength:	0.0 mg			ORDER INFORM	MATION			
component parts						B	Liquid filled, soft gelatin							
reverse numbered?		No				Dosage Form:	capsule		Unit of Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present				-		x Bottle		1 Bottle of 9	0 Capsules		
latex-free?		Yes	Anima	I, Alcohol		Product Shape:	Oblong		Box/Carton		(Write-in, e.	.g. 1 Box of 1	0 Vials)	
preservative-free?		Yes	Amina	i, Alconoi		1 Todact Onapc.			Ampule					
correctional institution block?		No				Product Color:	Yellow		Glass		Minimum o	rder quantity	y?	Yes
opioid?		No							Tube					
Cannabinoid?		No	Country of Origin	USA		Product Imprin	t: Printed with 'AT131' with		Vial Liquid Sgl					
If Unit Dose, is item bar coded to unhospital scanning?	unit dose for		Is this product covered			-	black ink		Vial Liquid Multi Vial Powder Sql			many of wh	ich package	type?
If Unit Dose, indicate NDC here:			Trade Agreements Act (Yes				Vial Powder Sgi Vial Powder Multi		24	Inner/Cartor	a/Pook	
il Offit Dose, indicate NDC fiere.			Trade Agreements Act (IAA):	162				Other: Write In			Case	I/Fack	
			FOR GENERIC DRUG PF	CODUCTS		<u> </u>			Guion Willom			Guoo		
			TOR GENERIO DROGTT	.000010										
					Aut	horized Generic *I	f Authorized Generic, other		PI	HARMACY ORDER	R / BILL UNIT			
I Oranga Book Batings	AB						ection fields are not applicable	Rec sell uni	t to customer?			nit to pharm	2011	
I. Orange Book Rating: II. Generic Equivalent to What Bra		Avodart						itee. sen um	t to dustomer .		KX billing u	Each	acy.	
ii. Generic Equivalent to What Bra	iliu:.	rtvodait						(Write-in, e.g	ı. 1 Vial)			Gram		
		DRUG SUPF	PLY CHAIN SECURITY ACT	(DSCSA) INFO	RMATION			, , , , ,	,,			Milliliter		
Does supplier meet DSCSA defini	ition of manufactur	rer?	Yes		GLN:	0331722498975			ITE	II AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			No											
If yes, select exemption:					GCP:					Dimensi	ions (US msr	nts.)	Volume	Saleable #
Other exemption - Write in:								_	Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		If yes, was ori	ginal product purcha	ased	Item/Each:	0.18	1.83	1.83	3.4	11.39	1
Is product sold by manufacturer's			Yes		direct from mi					1.00	1.00	0.4	11.00	
Has FDA granted waiver/exception		oduct?	No		Provide source	e manufacturer for re	epackaged product	Box/Carton/	Bundle/					
If yes, attach documentation fro	m FDA.							Inner Pack:						
		0.	TIN AND HIBCC PRODUCT I	NEODMATION				Case:	4.55	12.25	8.4	4	411.60	24
		G	TIN AND RIBCC PRODUCT	NFORMATION				Pallet:			-		-	
Saleable Unit of Measure	c	Saleable Quantity	HIBCC		GTIN	1.14	Unit of Use GTIN-14	Pallet:						
X Item/Each	3	1	TIBOO			31722131902	00331722131902	П						
Box/Carton/Bundle/Inner Pack					0000		0000112101002		COST INFORMATION	_		WHOLESAL	ER USE ONL	LY:
X Case		24			1033	31722131909								
Pallet								Regular Cos	t		Vendor #:			
								Invoice Cost	(WAC) (\$)	\$32.99	Whsl. Code	#:		
								11			Fineline Co	de:		
								As of date:	11/4/2016					
								П			1			
ļ.								11			1			
			Attach copy of SAFETY D	ATA SHEET (SI	DS) or non hazar		ISERT, LABEL AND PHOTO OF	PRODUCT PACK						
	formation on page	2				See new n 3 for De	esignated Drop Ship Only.		Signature:					



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For Designated Drop Ship Only Products, Please Use Page 3

MATERIAL HAZARD CLASSIFICATION and TRANSPORTATION							
Is this product (check all that apply):	SDS Hazard Classification						
a. Cytotoxic? b. CA Prop. 65 Carcinogen or Reproductive Toxicant?	SDS Hazard Classification						
Is the product a CA Prop 65 carcinogen?	x Organic Corrosive						
Is the product a CA Prop 65 reproductive toxicant?	Inorganic Oxidizer						
Does the product label bear a CA Prop 65 warning?	Steroid/Androgen Contact Hazard						
c. Contact Hazard?	Does the product have an Aerosol class? If yes, No						
d. Does this product require special clean-up instructions?	identify NFPA Storage Level:						
(If yes, attach SDS with special instructions.)	NFPA Storage Level:						
e. Does the product contain DEHP?							
Is this product regulated for shipment by DOT?	Is the product a NIOSH hazardous drug?						
(if yes, answer a-e below and provide SDS)	If yes, indicate which: Group 3 items (primarily adverse reproductive effects)						
a. UN/Identification Number							
b. Proper Shipping Name							
c. DOT Hazard Class	Hazardous Waste Identification						
d. Packing Group e. Inhalation Hazard?	EPA Hazardous Waste Code: Waste Characteristics						
	EFATI Idzaiduus Waste Code:						
Is this product regulated for shipment by IATA? (if yes, answer a-e below and provide SDS)	REMS or REGISTRY RESTRICTIONS						
a. UN/Identification Number	REIRO DI REGISTAT RESTRICTIONS						
b. Proper Shipping Name	Is there a REMS on this product?						
c. DOT Hazard Class	If Yes, is it managed with a pharmacy registry?						
d. Packing Group	Website URL:						
e. Inhalation Hazard?							
Is the product restricted for air shipment? If so, indicate restriction:	Med Guide Required No						
Passenger	Limited Distribution Requirement						
Cargo	Comments / Details: (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? No	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 #: NORDER*						
Limited Quantity Consumer Commodity, ORM-D	Site Enrollment Number assigned by Supplier: NPI #:						
Small Quantity (49 CFR 173.4)	by Supplier.						
Special Permit; DOT-SP	Comments						
Special Provision (listed in Column 7 of 49 CFR 172.101);							
SP#	Registry: No						
	Registry Program Contact Name: Phone:						
ADD'L STORAGE INFORMATION	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						
Schedule No. Is it a scheduled listed chemical product?: No	Is product returnable for credit: Yes						
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:	Special regulations or returns requirements for this						
Restricted to hospital, clinics, and physician offices only:	product in certain states?						
Restricted from US territories? (explain in comments)	If so, which states? Other requirements? Comments?						
Comments:							
MISCELLANE(DUS NOTES and/or Image of Product Barcode:						
Dutasteride is absorbed through the skin. Dutasteride capsules should not be handled by women who are							
absorption of dutasteride and the subsequent potential risk to a developing male fetus.	program of the season beginning because of the potential for						



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FOR DESIGNATED DROP SHIP PRODUCT ONLY - if not a designated drop ship, do not complete.

Order Method for Designated Drop S	nip Product	Standard Order Receipt and Processing
Purchase orders may be accepted by: a. EDI		Purchase order daily receipt cut off time by supplier Cut off time:
b. Autofax c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Phone:	per:	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 Designa	ed Drop Ship Fees:	Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each order: Drop Ship service fee billed with each order:		Overnight receipt available: PO Receipt cut off time:
Drop Ship miscellaneous fees billed: Comments:		Days of week overnight is available: Monday Tuesday Wednesday Thursday Friday
		Priority Overnight receipt available:
Class of Trade Restriction		PO Receipt Cut off time:
No restriction: Select YES if sold to retail pharmacy, hospitals, clinic Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	s and physician offices	Saturday Overnight receipt available: PO Receipt Cut off time: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:
Other Data Information Required to F	rocess 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?
Miscellaneous Notes:		
		ADDITIONAL INFORMATION
		Is product order for scheduled patient procedure? Is product order for restocking purposes?