

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

Version 2024						Introduction T	Type: New Item		x Final Version			Date:	10/10	0/2024	
			PRODUCT INFORMAT	ION					SPECIAL HAN	IDLING AND STOP	RAGE REQUI	REMENTS*			
Company Name: Camber Pharmaceuticals, Inc.				Application: ANDA		a. Temperatu	a. Temperature – Indicate the USP temperature range for this product.								
Application Number for NDA/AN	DA/BLA; PMA/510(k): 217166				NDA 505(b) Type:	NOT APPLICABLE	Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)								
Medical Device Class, if applicat	1														
DUNS:	11-856-3719	1.1						_	Other Temperature Range	Requirement					
Proprietary Name (If Applicable) a Selling Unit NDC:	31722-305-10	ame: K	Ketorolac Tromethamine Injection, Unit of Use NDC:	USP 15 mg/ml	_ (Single-Dose)	UPC:	331722305105	_	(write in) Notes						
UDI	31722-303-10		CVX Code:			MVX Code:	331722305105	-	NULES						
Description:	Ketorolac Tromett	hamine Injection	n, USP 15 mg/mL (Single-Dose Via						Is this product to be shippe	d to customers on	ice2		No	1	
Description.	Retorolac Homet	namme mjection	i, oor to mg/me (ongle-bose via	13)					Is this product to be shippe				No	-	
Active Ingredient(s): Ketorolac tromethamine, USP															
									b. Contact for temperature excursion questions:						
URL for Additional Product Inform Address:		www.camberpl	harma.com			Address 2:	1		Name:		Soma Raju				
City:	800 Centennial Av Piscataway	ve, Suite 1			State:	NJ	Zip: 08854		Number: Group E-mail:		732-529-042		m		
Key Contact:	Customer Service	•			Email:		@camberpharma.com	-	Group E-mail:						
Phone Number:	1-866-827-3647				Fax:				c. Special regulations for product in any states?				No		
Product Therapeutic Classification	n:	Nonsteroidal a	anti-inflammatory drug (NSAID)						Special returns requirement	ts for this product?			No		
														_	
	ADDITI	ONAL PRODUC	CT INFORMATION			PRODUCT	DESCRIPTION INFORMATION	d. Store prod	uct (unit of sale) upright?				No		
The product is?			Is the Product	Direct-Ship (Dnly			.	Protect product (unit of s	ale) from light?			No		
a legend device?		No	Is the Product	Unit Dose		Size:	10 x 1 mL single-dose,	e. Shelf life:					24	Months	
if yes, enter class #		No	Orphan Drug Status				flip top vials		Initial shelf life at launch	(If different):				Months	
a product kit? if yes, list NDCs of		No	FDA Approval Status			Strength:	15 mg/mL per single- dose vial			ORDER INFORM					
component parts			T DA Approval Glatas				Sterile clear solution								
reverse numbered?		No				Dosage Form	n:		Unit of Sale		What is the	NDC selling	unit?		
co-licensed?		No	Allergens Present					_	Bottle			x 1 mL Single			
latex-free?		Yes				Product Sha	pe: N/A		x Box/Carton		(Write-in, e	.g. 1 Box of 1	0 Vials)		
preservative-free? correctional institution block?		Yes					Slightly yellow	-	Ampule x Glass		Minimum o	rder quantity	0	Yes	
opioid?		No No				Product Cold	or:		Tube		Willing the second	ruer quantity		163	
Cannabinoid?		No	Country of Origin	India		Product Imp	N/A	-	x Vial Liquid Sgl						
If Unit Dose, is item bar coded to u	init dose for					Froduct imp			Vial Liquid Multi		If Yes, how	many of whi	ich package	type?	
hospital scanning?		Yes	Is this product covered up						Vial Powder Sgl		1	Each			
If Unit Dose, indicate NDC here:		31722-305-01	Trade Agreements Act (T	AA) (No				Vial Powder Multi Other: Write In	I		Inner/Cartor Case	1/Pack		
			FOR GENERIC DRUG PRO	DUCTS		1			Other. write in						
					Au	thorized Generic	*If Authorized Generic, other		Pł	IARMACY ORDER	R / BILL UNIT				
I. Orange Book Rating:	AP						section fields are not applicable	Rec. sell unit	Rec. sell unit to customer? Rx b				x billing unit to pharmacy:		
II. Generic Equivalent to What Bra	nd?:	Toradol							Each						
		DBUG SI	JPPLY CHAIN SECURITY ACT (I					(Write-in, e.g.				Gram			
		DRUG SC	JPPLT CHAIN SECORIT FACT (L	JSCSA) INFO	(WATION			HCPCS J-Co	J1885			Milliliter			
Does supplier meet DSCSA definit	tion of manufactur	rer?	Yes		GLN:	0331722498975				AND PACKING I	NFORMATIO	N			
Is product exempt from DSCSA?			No												
If yes, select exemption:				_	GCP:					Dimens	ions (US msr	nts.)	Volume	Saleable #	
Other exemption - Write in:									Weight Lbs.	Depth	Width	Height	(Cube)	Pieces	
Is product repackaged?			No			iginal product		Item/Each:	0.18	3.39	1.48	2	10.03	1	
Is product sold by manufacturer's			Yes	_	-	rect from mfr?						_			
Has FDA granted waiver/exception If yes, attach documentation from		roduct?	No		Provide sour	ce manufacturer fo	or repackaged product	Box/Carton/E Inner Pack:	Sundle/						
in yes, attach documentation nor	II FDA.							Case:							
			GTIN AND HIBCC PRODUCT IN	FORMATION					6.55	10.5	8.5	5.5	490.88	30	
								Pallet:							
Saleable Unit of Measure	RFID tag(Y/N)		HIBCC		GTI	N-14	Unit of Use GTIN-14								
X Item/Each	N	Quantity 1			003	31722305105									
Box/Carton/Bundle/Inner Pack	IN IN				003	0.1.22000100			COST INFORMATION			WHOLESAL	ER US <u>E ONI</u>	LY:	
x Case	N	30			203	31722305109									
Pallet								Regular Cost			Vendor #:				
							-	Invoice Cost	(WAC) (\$)	\$18.75	Whsl. Code				
				_			-	As of date:	10/10/2024		Fineline Co	ae:			
							-	As of uale:	10/10/2024						
							-								
			Attach copy of SAFETY DAT	TA SHEET (SD	S) or non haza	rd letter, PACKAGE	INSERT, LABEL AND PHOTO O	F PRODUCT PACK	AGING and BARCODE.						
*Please provide any additional inf	ormation on page	2.				See new p. 3 for	Designated Drop Ship Only.		Signature:						

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

Version 2024 For Designation	ted Drop Ship Only Products, Please Use Page 3					
MATERIAL HA	ZARD CLASSIFICATION and TRANSPORTATION					
Is this product (check all that apply): a. Cytotoxic? 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? Does the product label bear a CA Prop 65 warning? No	x Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard					
c. Contact Hazard? d. Does this product require special clean-up instructions? (If yes, attach SDS with special instructions.) e. Does the product contain DEHP? Is this product regulated for shipment by DOT? (if yes, answer a-e below and provide SDS) a. UN/dentification Number	Does the product have an Aerosol class? If yes, identify No NFPA Storage Level: NFPA Storage Level: Is the product a NIOSH hazardous drug? No If yes, indicate which: If yes, indicate which:					
b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	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?	REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? No If Yes, is it managed with a pharmacy registry? No Website URL: Image: Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2"Colspan="					
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: DEA #: Provider Name: DEA #: Site Enrollment Number assigned NCPDP#: by Supplier: NPI #:					
Special Provision (listed in Column 7 of 49 CFR 172.101); SP# ADD'L STORAGE INFORMATION	Registry: No Registry Program Contact Name: Phone: Comments					
Is the Product Controlled Substance Code Controlled Substance (List of II) ARCOS Reportable? No If yes, indicate which: Schedule No. CLASS OF TRADE RESTRICTION:	RETURN INSTRUCTIONS Contact tel. # if product received damaged: 1-866-827-3647 Is product returnable for credit: Yes					
CLASS OF TRADE RESTRICTION: No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Yes Restricted to retail pharmacy only: No Restricted to hospital, clinics, and physician offices only: No Restricted from US territories? (explain in comments) No Comments:	URL/Link to returns policy: contact - customerservice@camberpharma.com Special regulations or returns requirements for this product in certain states? No If so, which states? Other requirements? Comments?					
MISCELLANE	OUS NOTES and/or Image of Product Barcode:					



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Version 2024 FOR DESIGNATED DROP SHIP PRODUCT ONLY - in	f 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 Site Address: Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Name:	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:
Phone:	
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 second
	Priority Overnight receipt available:
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: 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?