

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

Version 2021					Introduction Type	: New Item		Final Version			Date:	4/10/2	2024
		PRODUCT INFORMA	TION					SPECIAL HAN	DLING AND STOP	RAGE REQUI	REMENTS*		
Company Name:	SOLA Pharmaceuticals				Application	ANDA	a. Temperature – Indi	ate the USP temp	erature range for t	his product.			
Application Number for NDA/AN	DA/BLA (drug); PMA/510(k)(r	ned device):	A21	18204				ature Range	Controlled Room		and 25 C (68	° – 77° F)	
Medical Device Class, if applicat													
DUNS:	080121345							emperature Range I	Requirement				
Proprietary Name (If Applicable) a	nd Established Name: 70512-790-05	Ketorolac Tromethamine Ophthaln Unit of Use NDC		6 5mL	UPC: 37	0512790059		rite in)					
Selling Unit NDC: UDI	70512-790-05	CVX Code:			MVX Code:	J512/90059	Notes						
Description:	Katavalaa Teemathamina Cal						la thia a	naduatta ha ahinna	d ta avatamana an i	2		No	
Description: Ketorolac Tromethamine Solution/Drops; Ophthalmic 0.5% Is this product to be shipped to customers on ice? No Is this product to be shipped to customers on dry ice? No													
Active ingredient(s): Ketorolac Tromethamine													
b. Contact for temperature excursion questions:													
URL for Additional Product Inform Address:	ation: 655 Highlandia Dr.			1	Address 2:		Name:			866-747-736	5		
City:	Baton Rouge			State:		p: 70810	Numbe			info@sola			
Key Contact:	Baton Hougo			Email:	info@solameds.us					into@solu	incus.us		
Phone Number:	866-747-7365			Fax:	800-754-9550		c. Special regulations	for product in any	states?			No	
Product Therapeutic Classification	n: Nonsteroi	dal Anti-inflammatory Drug (NSAID)]			Special	returns requirement	ts for this product?			No	
				4			_						
	ADDITIONAL PRO	DUCT INFORMATION			PRODUCT DES	CRIPTION INFORMATION	d. Store product (unit					Yes	
The product is?		Is the Product	Direct-Ship O	only				product (unit of sa	ale) from light?			Yes	
a legend device? if yes, enter class #	No	Is the Product Orphan Drug Status	Unit of Use		Size:	5mL	e. Shelf life:	helf life at launch (if different)			24	Months Months
a product kit?	No	Orphan Drug Status				0.5%		nen me at launch (n amerenty:				wonths
if yes, list NDCs of		FDA Approval Status			Strength:	0.070			ORDER INFORM	IATION			
component parts					Dosage Form:	Solution/Drops							
reverse numbered?	No				Dosuge Form.		Unit of				NDC selling	unit?	
co-licensed? latex-free?	No	Allergens Present					X	Bottle Box/Carton		1 Bottle	n 1 Dev ef 1	0 \/iele)	
preservative-free?	Yes	-			Product Shape:			Ampule		(write-in, e.	g. 1 Box of 1	u viais)	
correctional institution block?	No				Burn de la de la sec			Glass		Minimum or	rder quantity	?	Yes
opioid?	No				Product Color:			Tube				L	
Cannabinoid?	No	Country of Origin	India		Product Imprint:			Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	nit dose for	In this woodust sevened a	und an the					Vial Liquid Multi				ch package t	ype?
hospital scanning? If Unit Dose, indicate NDC here:		Is this product covered Trade Agreements Act (No				Vial Powder Sql Vial Power Multi		48	Each Inner/Carton	Pack	
in onit bose, indicate NDO here.		riddo / igroomonio / ior (110				Other: Write In			Case	and dork	
		FOR GENERIC DRUG PR	ODUCTS					_			4		
				AL		Authorized Generic, other ction fields are not applicable			IARMACY ORDER				
I. Orange Book Rating: II. Generic Equivalent to What Bran	AT nd?: ACULAR®				500		Rec. sell unit to custo 1 Bot			Rx billing u x	nit to pharma Each	acy:	
II. Generic Equivalent to what Bra	ACOLAIN	y					(Write-in, e.g. 1 Vial)	ue		-	Gram		
	DRU	G SUPPLY CHAIN SECURITY ACT	(DSCSA) INFOR	MATION							Milliliter		
		No.			00705 (000000)			1751			1		
Does supplier meet DSCSA definit Is product exempt from DSCSA?	tion of manufacturer?	Yes		GLN:	0370512000004			ITEN	I AND PACKING I	NFURMATIO	N		
If yes, select exemption:				GCP:	0370512				Dimensi	ions (US msn	nts.)	Volume	Saleable #
Other exemption - Write in:				JUF.	0010012			Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?		No			riginal product purchas	ed	Item/Each:	0.056	1.4567	1.378	2.7953	5.6110968	1
Is product sold by manufacturer's		No		direct from n				0.000	1.4007	1.570	2.1900	5.0110300	
Has FDA granted waiver/exception If yes, attach documentation from		No		Provide sour	ce manufacturer for re	packaged product	Box/Carton/Bundle/ Inner Pack:					0	
in yes, attach documentation non	IT DA.						Case:	0.4000	9.0551	5.7087	6.1024	315.45044	48
		GTIN AND HIBCC PRODUCT	NFORMATION					3.1288	9.0551	5.7087	6.1024	315.45044	48
Salashia Linit of Massura				0.7			Pallet:	593.184	48	40	47.44	91084.8	8640
Saleable Unit of Measure	Saleable Qu	antity HIBCC			IN-14 370512790059	Unit of Use GTIN-14 00370512790059							
Box/Carton/Bundle/Inner Pack				000			<u> </u>	ST INFORMATION			WHOLESAL	ER USE ONL'	Y:
X Case	48			503	370512790054								
Pallet							Regular Cost			Vendor #:			
	-			_			Invoice Cost (WAC) (\$)	\$28.00	Whsl. Code			
	-	_		-			As of date:	6/10/2024		Fineline Co	ue:		
				-			As of date.						
		Attach copy of SAFETY D	ATA SHEET (SD	S) or non haza	ard letter, PACKAGE INS	ERT, LABEL AND PHOTO OF	PRODUCT PACKAGING a	d BARCODE.					
*Please provide any additional infe	ormation on page 2.				See new p. 3 for Des	ignated Drop Ship Only.	Signatu	ire:					

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

Version 2021	For Design	ated Drop Ship Only Products, Please Use Page 3	
	MATERIAL H	AZARD CLASSIFICATION and TRANSPORTATION	
Is this product (check all that apply): a. Cytotoxic? b. CA Prop. 65 Carcinogen or Reproductive Tox Is the product a CA Prop 65 carcinogen? Is the product a CA Prop 65 reproductive to Does the product label bear a CA Prop 65 c. Contact Hazard? d. Does this product require special clean-up ins (If yes, attach SDS with special inst e. Does the product contain DEHP? Is this product regulated for shipment by DOT? (if yes, answer a-e below and provide SDS)	vicant? No warning? No structions? No	x Organic Inorganic Inorganic Steroid/Androgen Does the product have an Aerosol class? If yes, identify NFPA Storage Level: NFPA Storage Level: NFPA Storage Level: Is the product a NIOSH hazardous drug? If yes, indicate which:	DS Hazard Classification Corrosive Oxidizer Contact Hazard No No No
a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	No	Hazardous Waste Code:	ardous Waste Identification 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 c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	No	Is there a REMS on this product? If Yes, is it managed with a pharmacy registry? Website URL:	No No
Is the product restricted for air shipment? If so, ind Passenger Cargo Passenger & Cargo Is this a reportable quantity? No RQ Threshold: Is this a marine pollutant? No Is this product shipped utilizing an authorized DOT No (if yes, identify method below) Limited Quantity Consumer Commodity, ORM-D Small Quantity (49 CFR 173.4) Special Permit, DOT-SP		Med Guide Required Limited Distribution Requirement Comments / Details: (For example, iPledge program?) REMS: REMS Program Manager Name: Supplier Manages REMS registry exclusively: Wholesale distributor support: Provider Name: Site Enrollment Number assigned by Supplier: Comments	No Phone:
Special Provision (listed in Column 7 of 45 SP#) CFR 172.101); GE INFORMATION	Registry: Registry Program Contact Name: Comments	No Phone:
Controlled by State(s)? No Liste	rolled Substance Code d Chemical (List I or II) No ss, indicate which:	F Contact tel. # if product received damaged:	RETURN INSTRUCTIONS
Schedule No. Is it	a scheduled listed chemical product?:	Is product returnable for credit: URL/Link to returns policy:	Yes
Restricted to retail pharmacy only: Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices Restricted from US territories? (explain in commen Comments:	No No	Special regulations or returns requirements for this product in certain states? If so, which states? Other requirements? Comments?	No
	MISCELLAN	EOUS NOTES and/or Image of Product Barcode:	



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Version 2021	FOR DESIGNATED DROP SHIP PRODUCT ONLY - in	f not a designated drop ship, do not complete.
Order Method	l 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 #:	Fax Number: Fax Number: Fax Number: Phone No.: Site Address:	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 Ch	arges or Other Designated Drop Ship Fees:	Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each ord Drop Ship service fee billed with each ord Drop Ship miscellaneous fees billed: Comments:		Overnight receipt available: Image: Comparison of the second
CI	ass of Trade Restriction:	PO Receipt Cut off time:
	oharmacy, hospitals, clinics and physician offices	Saturday Overnight receipt available: PO Receipt Cut off time: Order receipt method: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:
Other Data Ir	nformation 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?