

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

Version 2021						Introduction Ty	/pe: New Iter	m		Final Version			Date:	4/10/	2024	
			PRODUCT INFORMA	TION						SPECIAL HAN	DLING AND STOR	RAGE REQUI	REMENTS*			
Company Name: SOLA Pharmaceuticals Application: ANDA								4	a. Temperature – Indicate the USP temperature range for this product.							
Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device):  A218204  Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)																
Medical Device Class, if applica																
DUNS:	080121345									Other Temperature Range F	Requirement					
Proprietary Name (If Applicable) a		lame: Ketor	rolac Tromethamine Ophthalm	nic Solution 0.5	% 10mL					(write in)						
Selling Unit NDC:	70512-790-10		Unit of Use NDC:				370512790103		1	Notes						
UDI			CVX Code:			MVX Code:										
Description:	Ketorolac Trome	thamine Solution/Dro	ops; Ophthalmic 0.5%							s this product to be shipped				No		
									1	s this product to be shipped	d to customers on o	Iry ice?		No		
Active Ingredient(s): Ketorolac Tromethamine									h Camtant faut							
URL for Additional Product Information:										emperature excursion que Name:	estions:					
Address:	655 Highlandia [	)r			1	Address 2:				Number:		866-747-736	35			
City:	Baton Rouge					LA	<b>Zip</b> : 70810			Group E-mail:		info@sola				
Key Contact:				info@solameds												
Phone Number:	866-747-7365				Fax:	800-754-9550			c. Special regu	lations for product in any	states?			No		
Product Therapeutic Classification	n:	Nonsteroidal Anti-	Nonsteroidal Anti-inflammatory Drug (NSAID)							Special returns requirement	s for this product?			No		
					1											
	ADDIT	IONAL PRODUCT II	NFORMATION			PRODUCT D	ESCRIPTION INFORMA	ATION	d. Store product (unit of sale) upright?							
The product is?			Is the Product	Direct-Ship (	Only				F	Protect product (unit of sa	ile) from light?			Yes		
a legend device?		No	Is the Product	Unit of Use		Size:	10mL		e. Shelf life:					24	Months	
if yes, enter class #			Orphan Drug Status			0.20.			I	nitial shelf life at launch (	if different):				Months	
a product kit?		No				Strength:	0.5%				ORDER INFORM	ATION				
if yes, list NDCs of			FDA Approval Status				Solution/Drops				ORDER INFORM	IATION				
component parts reverse numbered?		No				Dosage Form	: Solution/Diops		l ,	Jnit of Sale		What is the	NDC selling	unit?		
co-licensed?		No	Allergens Present							x Bottle		1 Bottle				
latex-free?		Yes	J			Door door to Ohio oo				Box/Carton			g. 1 Box of 1	0 Vials)		
preservative-free?		Yes				Product Shap	e:			Ampule			•	•		
correctional institution block?		No				Product Color	<b>.</b> .			Glass		Minimum or	rder quantity	/?	Yes	
opioid?		No				Froduct Color				Tube						
Cannabinoid?		No	Country of Origin	India		Product Impri	nt:		_	Vial Liquid Sgl					_	
If Unit Dose, is item bar coded to	unit dose for		In this case does a consequent						-	Vial Liquid Multi Vial Powder Sql			many of whi	ich package t	type?	
hospital scanning? If Unit Dose, indicate NDC here:			Is this product covered u  Trade Agreements Act (		No				_	Vial Powder Sqi Vial Power Multi		48	Lacn Inner/Cartor	n/Pack		
ii onit bose, indicate NBC here.			Trade Agreements Act (	1701):	140					Other: Write In			Case	I/I dok		
			FOR GENERIC DRUG PR	ODUCTS					_				1.			
									1							
					Au		*If Authorized Generic, o		PHARMACY ORDER / BILL UNIT							
I. Orange Book Rating:	AT						section fields are not app	plicable	Rec. sell unit to	customer?		Rx billing u	nit to pharm	acy:		
II. Generic Equivalent to What Bra	and?:	ACULAR®								1 Bottle		х	Each	•		
									(Write-in, e.g. 1	Vial)			Gram			
		DRUG SUPP	PLY CHAIN SECURITY ACT (	DSCSA) INFO	RMATION								Milliliter			
Does supplier meet DSCSA defin	ition of manufactu	12023	Yes		GLN:	0370512000004				ITEM	I AND PACKING II	NEODMATIO	N			
Is product exempt from DSCSA?	ition of manufacti	liei r	No No	-	GLN.	0370312000004				11 = 11	I AND FACRING II	VI OKWATIOI	N.			
If ves. select exemption:					GCP:	0370512					Dimonei	ons (US msn	nte )	Volume	Saleable #	
Other exemption - Write in:					GUF.	03/03/2				Weight Lbs.	Depth	Width	Height	(Cube)	Pieces	
Is product repackaged?			No		If ves, was o	riginal product purcl	hased		Item/Each:							
Is product sold by manufacturer's	s exclusive distrib	outor?	No		direct from n					0.0613	1.4567	1.378	2.7953	5.6110968	1	
Has FDA granted waiver/exception	n/exemption for p		No		Provide sour	rce manufacturer for	repackaged product		Box/Carton/Bui	ndle/				0		
If yes, attach documentation fro	m FDA.								Inner Pack:					Ů		
		0.7	TIN AND HIBCC PRODUCT II	UEODMATION					Case:	3.3819	9.0551	5.7087	6.1024	315.45044	48	
		GI	TIN AND RIBCC PRODUCT II	NFORMATION					Pallet:							
Saleable Unit of Measure		Saleable Quantity	HIBCC		GT	IN-14	Unit of Use GTII	N_14	railet.	638.742	48	40	47.44	91084.8	8640	
X Item/Each		1	TIIBOO			370512790103	0037051279010									
Box/Carton/Bundle/Inner Pack	Box/Carton/Bundle/Inner Pack						COST INFORMATION			WHOLESALER USE ONLY:						
X Case		48			503	370512790108										
Pallet	_								Regular Cost			Vendor #:				
									Invoice Cost (W	/AC) (\$)	\$56.00	Whsl. Code				
									As of data:	6/10/2024		Fineline Co	de:			
									As of date:	0/10/2024						
•			Attach copy of SAFETY DA	ATA SHEET (SI	OS) or non haza	ard letter, PACKAGE	INSERT, LABEL AND PH	HOTO OF P	RODUCT PACKAG	SING and BARCODE.		•				
		e 2.	.,	(	,		Designated Drop Ship C			Signature:						



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

#### Version 2021 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 No a. Cytotoxic? b. CA Prop. 65 Carcinogen or Reproductive Toxicant? Is the product a CA Prop 65 carcinogen? No x Organic Corrosive Is the product a CA Prop 65 reproductive toxicant? Inorganic Oxidizer No Does the product label bear a CA Prop 65 warning? Steroid/Androgen No Contact Hazard c. Contact Hazard? Does the product have an Aerosol class? If yes, No No identify NFPA Storage Level: d. Does this product require special clean-up instructions? No (If yes, attach SDS with special instructions.) NFPA Storage Level: e. Does the product contain DEHP? No Is this product regulated for shipment by DOT? No Is the product a NIOSH hazardous drug? No (if yes, answer a-e below and provide SDS) If yes, indicate which: a. UN/Identification Number b. Proper Shipping Name Hazardous Waste Identification c. DOT Hazard Class d. Packing Group EPA Hazardous Waste Code: Waste Characteristics e. Inhalation Hazard? No Is this product regulated for shipment by IATA? No (if yes, answer a-e below and provide SDS) **REMS or REGISTRY RESTRICTIONS** a. UN/Identification Number b. Proper Shipping Name Is there a REMS on this product? If Yes, is it managed with a pharmacy registry? c. DOT Hazard Class No d. Packing Group Website URL: e. Inhalation Hazard? No Is the product restricted for air shipment? If so, indicate restriction: Med Guide Required No No Passenger Limited Distribution Requirement No Comments / Details: (For example, iPledge program?) Cargo Passenger & Cargo Is this a reportable quantity? REMS: No Phone: RQ Threshold: REMS Program Manager Name: Is this a marine pollutant? Supplier Manages REMS registry exclusively: Wholesale distributor support: Is this product shipped utilizing an authorized DOT exception or Special Permit? Provider Name: (if yes, identify method below) 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: Phone: Registry Program Contact Name: ADD'L STORAGE INFORMATION Comments Is the Product RETURN INSTRUCTIONS Controlled Substance? No Controlled Substance Code Controlled by State(s)? No Listed Chemical (List I or II) No 866-747-7365 ARCOS Reportable? No If yes, indicate which: Contact tel. # if product received damaged: Schedule No. Is it a scheduled listed chemical product?: Yes Is product returnable for credit: CLASS OF TRADE RESTRICTION: URL/Link to returns policy: No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Yes Restricted to retail pharmacy only: No Special regulations or returns requirements for this product in certain states? Restricted to hospital, clinics, and physician offices only: No No Restricted from US territories? (explain in comments) If so, which states? Other requirements? Comments? No Comments:

MISCELLANEOUS NOTES and/or Image of Product Barcode



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

#### Version 2021

### FOR DESIGNATED DROP SHIP PRODUCT ONLY - if 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  Minimum Order Quantity:  Supplier's Customer Service Number:  Contracted 3PL company / contact #:  Name: Phone:	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 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:  PO Receipt cut off time:  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, 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:  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:  Miscellaneous Notes:	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?							
	ADDITIONAL INFORMATION							
	Is product order for scheduled patient procedure? Is product order for restocking purposes?							