

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

Version 2020				Introduction Type:	Open Stock	7	Final Version			Date:	7/10/	/2024	
	PR	ODUCT INFORMATION				_	SPECIAL HAN	DLING AND STOP	RAGE REQUI	REMENTS*			
Company Name: Xiromed LLC ANDA							a. Temperature – Indicate the USP temperature range for this product.						
Application Number for NDA/AN	DA/BLA (drug); PMA/510(k)(med device):	20970	01				Temperature Range	Controlled Room	– between 20	and 25 C (68	° – 77° F)		
DUNS:	080228637					_	Other Temperature Range	Requirement					
Proprietary Name (If Applicable) a				1100 07070	2400450	41	(write in)						
Selling Unit NDC: UDI	70700-106-15	Unit of Use NDC: CVX Code:		UPC: 370700 MVX Code:	0106150		Notes						
Description:	Clobetasol Propionate Ointment USP, 0.05% 1						Is this product to be shipped	I to customers on i	ce?		No		
	,,,,,,,,,,	-3					Is this product to be shipped				No	-	
Active Ingredient(s):	Active Ingredient(s): Clobetasol Propionate												
URL for Additional Product Inform	b. Contact for temperature excursion questions: Name: Xiromed Quality												
Address:	180 Park Ave			Address 2: Suite	101		Number:		844-947-663	,			
City:	Florham Park		State:	NJ Zip:			Group E-mail:		US-Quality	/-Xiromed@	exiromed.co	om	
Key Contact: Phone Number:	Xiromed Regulatory 844-947-6633		Email: Fax:	usregulatory@xiron 862-286-0932	<u>ned.com</u>		gulations for product in any				Ne		
Product Therapeutic Classification			T dA.	002-200-0332		c. Special reg	Special returns requirement				No No	-	
							opoolal lotarito loquitoritori	o for the product.				-	
	ADDITIONAL PRODUCT INFORMAT	rion		PRODUCT DESCR	RIPTION INFORMATION	d. Store proc	luct (unit of sale) upright?						
The product is?		Product Direct-Ship Only	у				Protect product (unit of sa	ale) from light?					
a legend device?		Product Neither		Size:	15gm	e. Shelf life:		·····			24	Months	
if yes, enter class # a product kit?	No No	an Drug Status			0.05%		Initial shelf life at launch (if different):				Months	
if yes, list NDCs of		Approval Status		Strength:				ORDER INFORM	MATION				
component parts				Dosage Form:	Topical				M/h = 4 1 = 4 h =				
reverse numbered? co-licensed?	No Aller	gens Present					Unit of Sale Bottle		1 Box contai	NDC selling	unit ?		
latex-free?	Ne	Not made with natural rubber latex.		Product Shape:			x Box/Carton			g. 1 Box of 1	0 Vials)		
preservative-free?	Yes	Not made with natural rubber latex.		Product Shape:			Ampule						
correctional institution block? opioid?	Yes			Product Color:			Glass Tube		Minimum o	rder quantity	?	Yes	
Cannabinoid?	No Coun	try of Origin Spain					Vial Liquid Sgl						
If Unit Dose, is item bar coded to u				Product Imprint:			Vial Liquid Multi				ich package f	type?	
scanning?		s product covered under the					Vial Powder Sql		48	Each			
If Unit Dose, indicate NDC here:	Irade	e Agreements Act (TAA)? Y	es				Vial Power Multi Other: Write In			Inner/Cartor Case	Pack		
	FOR G	ENERIC DRUG PRODUCTS		<u>.</u>									
									-				
		L	Auth		norized Generic, other section are not applicable			ARMACY ORDER					
I. Orange Book Rating: II. Generic Equivalent to What Bra	AB Temovate			lieius e		Rec. sell unit	to customer? 1 Tube	1	Rx billing u	nit to pharm Each	acy:		
II. Generic Equivalent to What Bra	inu :.					(Write-in, e.g		1		Gram			
	DRUG SUPPLY CHAIN	N SECURITY ACT (DSCSA) INFORM	IATION				,			Milliliter			
Does supplier meet DSCSA defini	ition of manufacturer?	Yes GLN:		037070000007			ITEN	AND PACKING I	NFORMATIO	N			
Is product exempt from DSCSA?	No			0010100000001									
If yes, select exemption:						-	Weight Lbs.	Dimensi	ions (US msn	nts.)	Volume	# Pieces:	
Other exemption - Write in:							weight Lbs.	Depth	Width	Height	(Cube)	# Fleces.	
Is product repackaged? Is product sold by manufacturer's	No No		s, was origi t from mfr?	nal product purchased		Item/Each:	0.033	1.457	4.528	1.083	7.1448716	1	
Has FDA granted waiver/exception				cumentation from FDA.		Box/Carton/E	Bundle/				0		
						Inner Pack:					U		
	GTIN AND H	IIBCC PRODUCT INFORMATION				Case:	3.5	12.6	7.5	5.1	481.95	48	
Saleable Unit of Measure	Quantity HIBC	C	GTIN-	14	Unit of Use GTIN-14	Pallet:					0		
x Item/Each	1			700106150							0		
Box/Carton/Bundle/Inner Pack			40070	700106157			COST INFORMATION				ER USE ONL	M .	
X Case Pallet	48		10370	100106157			COST INFORMATION			WHOLESAL	ER USE ONL	.1:	
			-			Regular Cost	t		Vendor #:				
						Invoice Cost		\$12.00	Whsl. Code				
	4					As of date:	8/2/2021		Fineline Co	de:			
						As of date.	0/2/2021		1				
	Attach	copy of SAFETY DATA SHEET (SDS)) or non haza	ard letter, PACKAGE INSE	RT, LABEL AND PHOTO OF	PRODUCT PACK	AGING and BARCODE.		·				
					nated Drop Ship Only.		Signature:						

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

Version 2020 For Designated Drop Ship Only Products, Please Use Page 3						
MATERIAL HAZ	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? Is the product a CA Prop 65 reproductive toxicant? Does the product label bear a CA Prop 65 warning? No	SDS Hazard Classification Organic Corrosive Inorganic Oxidizer X Steroid/Androgen Contact Hazard					
c. Contact Hazard? No d. Does this product require special clean-up instructions? No (If yes, attach SDS with special instructions.) e. Does the product contain DEHP? No Is this product regulated for shipment by DOT? No (if yes, answer a-e below and provide SDS)	Aerosol Class; Identify NFPA Storage Level: Is the product a NIOSH hazardous drug? If yes, indicate which:					
a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard? No	Hazardous Waste Identification EPA Hazardous Waste Code: Waste Characteristics					
	REMS or REGISTRY RESTRICTIONS					
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. Deaking Course	Is there a REMS on this product? No If Yes, is it managed with a pharmacy registry? Website URL:					
d. Packing Group e. Inhalation Hazard? No	Med Guide Required No					
Is the product restricted for air shipment? If so, indicate restriction: Passenger Cargo Passenger & Cargo	Limited Distribution Requirement No Comments / Details: (For example, iPledge program?)					
Is this a reportable quantity? No RQ Threshold: Is this a marine pollutant? <u>No</u> Is this product shipped utilizing an authorized DOT exception or Special Permit? <u>No</u> (if yes, identify method below) Limited Quantity Consumer Commodity, ORM-D Small Quantity (49 CFR 173.4) Special Permit; DOT-SP Special Perovision (listed in Column 7 of 49 CFR 172.101);	REMS: Phone: Supplier Manages REMS registry exclusively: Phone: Wholesale distributor support: DEA #: Provider Name: DEA #: Site Enrollment Number assigned PCPDP#: by Supplier: NPI #:					
SP#	Registry:					
ADD'L STORAGE INFORMATION Is the Product	Registry Program Contact Name: Phone: Phone: Comments					
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: If yes, indicate which:	Contact tel. # if product received damaged: Is product returnable for credit: URL/Link to returns policy:					
Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only:	Special regulations or returns requirements for this product in certain states?					
Restricted from US territories? (explain in comments) Comments:	If so, which states? Other requirements? Comments?					
MISCELLANE	OUS NOTES and/or Image of Product Barcode:					



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Version 2020	FOR DESIGNATED DROP SHIP PRODUCT ONLY - if	not a designated drop ship, do not complete.					
Order Metho	od 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: Phone No.: Site Address: 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 C	Charges or Other Designated Drop Ship Fees:	Overnight and Priority Overnight PO Processing					
Expedited freight fees billed with each of Drop Ship service fee billed with each of 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					
(Class of Trade Restriction:	PO Receipt Cut off time:					
No restriction: Select YES if sold to retain Restricted to retail pharmacy only: Restricted to hospital, clinics, and physic Restricted from US territories? (explain in Comments:		Saturday Overnight receipt available: PO Receipt Cut off time: PO Receipt Cut off time: Phone: Order receipt method: Phone: Fax: EDI: Overnight Fees apply: Image: Constraint of the state of th					
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?					