

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

Version 2020						Introduction Type:	Post Launch Change		Final Version			Date:	7/10	0/2024
			PRODUCT INFORMA	TION					SPECIAL HA	NDLING AND STO	RAGE REQUI	REMENTS*		
Company Name: Xiromed LLC Application: ANDA							a. Temperature – Indicate the USP temperature range for this product.							
Application Number for NDA/AN		/510(k)(med devi	ice):	21261	2		!	·	Temperature Range	Controlled Room		and 25 C (68	° – 77° F)	
DUNS:	080228637								Other Temperature Range	Requirement				
Proprietary Name (If Applicable) a	and Established Name	e: GLYC0	OPYRROLATE						(write in)					
Selling Unit NDC:	70700-165-25		Unit of Use NDC:				165256		Notes					
UDI			CVX Code:			MVX Code:								
Description:	Glycopyrrolate Inject	tion, USP 0.2 mg/r	mL Single Dose Vial						Is this product to be shippe	d to customers on i	ce?		No	_
									Is this product to be shippe	d to customers on o	Iry ice?		No	_
Active Ingredient(s): GLYCOPYRROLATE								41						
URL for Additional Product Inform	mation:							b. Contact to	r temperature excursion qι Name:	iestions:	Xiromed Qu	ality		
Address:	180 Park Ave					Address 2: Suite 1	01		Number:		844-947-663			
City:	Florham Park				State:	NJ Zip:	07932		Group E-mail:				xiromed.co	om
Key Contact:	Xiromed Regulatory				Email:	usregulatory@xirom	ed.com	<u></u>						
Phone Number:	844-947-6633				Fax:	862-286-0932		c. Special reg	gulations for product in any				No	_
Product Therapeutic Classification	on:								Special returns requiremen	ts for this product?			No	_
	ADDITION	AL DRODUCT IN	FORMATION			PROPUST PESSE	UDTION INCODMATION						.,	
	ADDITION	AL PRODUCT IN				PRODUCT DESCR	RIPTION INFORMATION	d. Store prod	uct (unit of sale) upright?				Yes	_
The product is?			Is the Product	Direct-Ship Only	'				Protect product (unit of sa	ale) from light?			Yes	-
a legend device? if yes, enter class #	N	lo	Is the Product	Neither		Size:	25 x 1 mL Single Dose Vials	e. Shelf life:	luitial abalf life at lavuab	if different).			24	Months Months
a product kit?		lo	Orphan Drug Status				0.2MG/ML		Initial shelf life at launch	ıτ αιπerent):				Months
if yes, list NDCs of	1,	10	FDA Approval Status			Strength:	O.ZIVIO/IVIE			ORDER INFORI	MATION			
component parts						Dosage Form:	INJECTABLE							
reverse numbered?		lo				Dosage Form.			Unit of Sale		What is the		unit?	
co-licensed?		lo	Allergens Present						Bottle		1 Box of 25			
latex-free?		'es				Product Shape:			x Box/Carton Ampule		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free? correctional institution block?		lo 'es							Glass		Minimum o	der quantity	2	Yes
opioid?	_	lo				Product Color:			Tube		William Ci	der quartity	•	103
Cannabinoid?		lo	Country of Origin	India		Product Imprint:			Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	unit dose for					Froduct imprint.			Vial Liquid Multi		If Yes, how	many of whi	ch package t	type?
hospital scanning?			Is this product covered u						Vial Powder Sql		12	Each		
If Unit Dose, indicate NDC here:			Trade Agreements Act (ΓΑΑ)? <u>N</u>	0				Vial Power Multi			Inner/Cartor	n/Pack	
									Other: Write In			Case		
			FOR GENERIC DRUG PF	ODUCIS				_						
					Autho	rized Generic *If Auth	orized Generic, other section	PHARMACY ORDER / BILL UNIT						
I. Orange Book Rating:	AP			_			re not applicable	Rec. sell unit	to customer?	Rx billing unit to pharmacy:				
II. Generic Equivalent to What Bra	and?:	Robinul®						1 Vial x Each						
								(Write-in, e.g. 1 Vial) Gram						
		DRUG SUPPI	LY CHAIN SECURITY ACT	(DSCSA) INFORM.	ATION							Milliliter		
Does supplier meet DSCSA defini	ition of manufacturar	2	Yes	GLN:		0370700000007			ITE	M AND PACKING I	NEORMATIO	N		
Is product exempt from DSCSA?	naon or manufacturer	• —	No	_ GLN.		007070000007								
If yes, select exemption:				_						Dimens	ions (US msn	nts.)	Volume	
Other exemption - Write in:									Weight Lbs.	Depth	Width	Height	(Cube)	# Pieces:
Is product repackaged?	_		No			al product purchased		Item/Each:	0.47	3.54	3.54	1.77	22.180932	1
Is product sold by manufacturer's			No No		from mfr?			Box/Carton/E						
Has FDA granted waiver/exceptio	on/exemption for prod	uct?	INO	if yes,	, attach doc	umentation from FDA.		Inner Pack:	sunale/				0	
		GT	IN AND HIBCC PRODUCT	NFORMATION				Case:	0.045	7.400	7 400	F 000	323.5874	12
									6.615	7.402	7.402	5.906	323.5874	12
Saleable Unit of Measure		Quantity	HIBCC		GTIN-1		Unit of Use GTIN-14	Pallet:					0	
X Item/Each		1			003707	700165256								
Box/Carton/Bundle/Inner Pack X Case	-	12			20370	700165250			COST INFORMATION			WHO! ESAL	ER USE ONL	γ.
Pallet	-	12			200701	33.33200			SOOT IN CHIRATION			MIOLEGAL	EN OOL ONE	
	7							Regular Cost	:		Vendor #:			
								Invoice Cost		\$67.00	Whsl. Code			
	↓ □							1			Fineline Co	de:		
								As of date:			-			
Attach copy of SAFETY DATA SHEET (SDS) or non hazard letter, PACKAGE INSERT, LABEL AND PHOTO OF PRODUCT PACKAGING and BARCODE.														
1			Allach copy of SAFETY L	ATA SHEET (SDS)	or non naza	ard letter, PACKAGE INSE		RODUCT PACK						
*Please provide any additional inf	formation on nage ?					See new p. 3 for Design	ated Dron Ship Only		Signature:					



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For Designated Drop Ship Only Products, Please Use Page 3

MATE	RIAL HAZARD 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?	No SDS Hazard Classification No Organic Corrosive No Inorganic Oxidizer No Steroid/Androgen Contact Hazard
Does the product label bear a CA Prop 65 warning? 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?	No Steroid/Androgen Contact Hazard No Aerosol Class; Identify NFPA Storage Level: No Is the product a NIOSH hazardous drug? No If yes, indicate which:
Is this product regulated for shipment by DOT? (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?	Hazardous Waste Identification EPA Hazardous Waste Code: 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	No REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? If Yes, is it managed with a pharmacy registry? Website URL:
e. Inhalation Hazard? Is the product restricted for air shipment? If so, indicate restriction: 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)	REMS: REMS Program Manager Name: Supplier Manages REMS registry exclusively: Wholesale distributor support: Provider Name: Site Enrollment Number assigned by Supplier: PCPDP#: NPI #:
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101); SP# ADD'L STORAGE INFORMATION	Registry: Registry Program Contact Name: Comments Phone:
Is the Product Controlled Substance? Controlled Substance Code Controlled by State(s)? ARCOS Reportable? No If yes, indicate which:	No Contact tel. # if product received damaged:
Schedule No. Is it a scheduled listed chemical product?: CLASS OF TRADE RESTRICTION: No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Restricted to retail pharmacy only:	No Is product returnable for credit: URL/Link to returns policy: Yes Consider stude for a set was possible for this
Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	Special regulations or returns requirements for this product in certain states? If so, which states? Other requirements? Comments?
MISC	ELLANEOUS NOTES and/or Image of Product Barcode:



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

Order Method for Designation	ated Drop Ship Product	Standard Order Receipt and Processing					
Purchase orders may be accepted by:		Purchase order daily receipt cut off time by supplier					
a. EDI		Cut off time:					
b. Autofax	Fax Number:						
c. Fax	Fax Number:	Shipping lead time of PO: Hours	Days				
d. Phone only	Phone No.:						
e. Supplier Web Site only	Site Address:	Ships same day for next day receipt:					
Minimum Order Quantity:		Ships for second day receipt:					
Supplier's Customer Service Number:		Ships regular ground for 3-10 days receipt:					
Contracted 3PL company / contact #: Name: Phone:							
Expedited Freight Charges or Ott	her Designated Drop Ship Fees:	Overnight and Priority Overnight PO Proc	cessing				
Expedited freight fees billed with each order:		Overnight receipt available:					
Drop Ship service fee billed with each order:		PO Receipt cut off time:					
Drop Ship miscellaneous fees billed:		Days of week overnight is available:	Monday				
Comments:			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, ho	ospitals, clinics and physician offices	Saturday Overnight receipt available:					
Restricted to retail pharmacy only:		PO Receipt Cut off time:					
Restricted to hospital, clinics, and physician offices onl	y:	Order receipt method: Phone: Phone #:					
Restricted from US territories? (explain in comments)		Fax: Fax #:					
Comments:		EDI:					
		Overnight Fees apply:					
		Other fees apply:					
Other Data Information F	Required to Process PO:	Return Instructions					
Patient Procedure Date:		Contact # if product is received damaged:					
Physician Name:		Is product returnable for credit:					
Physician/Clinic Phone #		URL/Link to returns policy:					
Physician State License #							
Physician/Clinic DEA #:		Special regulations or returns requirements for this product in certain	n states?				
Physician/Clinic Specialty:		If so, which states? Other requirements? Comments?					
Miscellaned	ous Notes:						
		ADDITIONAL INFORMATION					
		Is product order for scheduled patient procedure?					
		Is product order for restocking purposes?					
		1 .					