

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

Version 2020				Introduction Type:	Post Launch Change		Final Version			Date:	7/10/)/2024
		PRODUCT INFORMATION					SPECIAL HAN	IDLING AND STO	RAGE REQUI	REMENTS*		
Company Name:	Xiromed LLC	d desdeed.	212995	Application:	ANDA		- Indicate the USP temp			and 25 C (69	0° 77° F\	
	NDA/BLA (drug); PMA/510(k)(med	d device):	Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)									
DUNS:	080228637)-fT-bl-t- 000				. °	Other Temperature Range	Requirement				
Proprietary Name (If Applicable) a Selling Unit NDC:	70700-271-30	Deferasirox Tablets, 360mg Unit of Use NDC:		UPC: 37070	0271308	N N	(write in) lotes					
UDI	10100-211-30	CVX Code:		MVX Code:	027 1300		ioles					
Description:	Deferasirox Tablets, 360mg - 30			<u> </u>			s this product to be shipped				No	
Active Ingredient(s):	Deferasirox						s this product to be shipped		dry ice?		No	-
URL for Additional Product Infor	mation:						emperature excursion qu lame:	iestions:	Xiromed Qu	ality		
Address:	180 Park Ave			Address 2: Suite	101		lumber:		844-947-663			
City:	Florham Park		State:	NJ Zip:			Froup E-mail:				@xiromed.co	om
Key Contact:	Xiromed Regulatory		Email:	usregulatory@xiror			•					
Phone Number:	844-947-6633		Fax:	862-286-0932		c. Special regula	ations for product in any	states?			No	_
Product Therapeutic Classification	Iron Chelating	g Agent				S	pecial returns requirement	ts for this product?	•		No	_
	ADDITIONAL PRODUC	CT INFORMATION		PRODUCT DESC	RIPTION INFORMATION	d. Store product	t (unit of sale) upright?				Yes	_
The product is?		Is the Product Dis	rect-Ship Only			P	rotect product (unit of s	ale) from light?			Yes	_
a legend device?	No		either	Size:	30 Count Bottle	e. Shelf life:					24	Months
if yes, enter class #		Orphan Drug Status				In	nitial shelf life at launch ((if different):				Months
a product kit?	No	FDA Assessed Otator		Strength:	360mg			OBDER INFOR	MATION			
if yes, list NDCs of component parts		FDA Approval Status			Tablets			ORDER INFOR	MATION			
reverse numbered?	No			Dosage Form:	Tablets	ll u	Init of Sale		What is the	NDC selling	unit?	
co-licensed?	No	Allergens Present				ll É	x Bottle		1 Bottle			
latex-free?	No	Not made with natural	rubbor latov	Product Shape:	Film-coated, oval biconvex		Box/Carton		(Write-in, e.	.g. 1 Box of 1	0 Vials)	
preservative-free?	Yes	Not made with natural	rubber latex.	Flouuci Sliape.			Ampule					
correctional institution block?				Product Color:	Blue	<u> </u>	Glass		Minimum o	rder quantit	y?	Yes
opioid?	No						Tube					
Cannabinoid?	No No	Country of Origin Inc	dia	Product Imprint:	Debossed "360" on one side		Vial Liquid Sgl		If Van haw		lah naskans	4
If Unit Dose, is item bar coded to a scanning?	unit dose for nospital	Is this product covered under	the		side	 	Vial Liquid Multi Vial Powder Sql			Each	ich package t	type?
If Unit Dose, indicate NDC here:		Trade Agreements Act (TAA)?					Vial Power Multi		2.7	Inner/Cartor	n/Pack	
							Other: Write In			Case		
		FOR GENERIC DRUG PRODU	CTS			_				_		
									_			
			Autho		horized Generic, other section		PH	IARMACY ORDER	R / BILL UNIT			
I. Orange Book Rating:	AB			fields	are not applicable	Rec. sell unit to customer?			Rx billing unit to pharmacy:			
II. Generic Equivalent to What Br	and?: Jadenu						1 Bottle		х	Each		
	22112.01	URBLY ALLEN ARALIBITY A AT (DAG	OAN INTODIAL TION			(Write-in, e.g. 1	Vial)			Gram		
	DRUG S	UPPLY CHAIN SECURITY ACT (DSC	SA) INFORMATION							Milliliter		
Does supplier meet DSCSA defin	sition of manufacturor?	Yes	GLN:	0370700000007			ITEN	AND PACKING	INFORMATIO	N		
Is product exempt from DSCSA?		No	OLN.	007070000007						•		
If yes, select exemption:				l .		•		Dimens	sions (US msn	nts)	Volume	
Other exemption - Write in:							Weight Lbs.	Depth	Width	Height	(Cube)	# Pieces:
Is product repackaged?		No	If Yes, was origin	nal product purchased		Item/Each:	0.12	1.705	3.26	1.705	9.4769015	1
Is product sold by manufacturer'	s exclusive distributor?	No	direct from mfr?				0.12	1.705	3.26	1.705	9.4769015	1
Has FDA granted waiver/exception	on/exemption for product?	No	If yes, attach doo	cumentation from FDA.		Box/Carton/Bun	ndle/				0	
						Inner Pack:						
		GTIN AND HIBCC PRODUCT INFOR	RMATION			Case:	4.32	7.625	3.375	5.75	147.97266	24
Saleable Unit of Measure	Quantity	HIBCC	GTIN-	14	Unit of Use GTIN-14	Pallet:				-	-	
Item/Each	quality	нвес		700271308	Offic of Ose GTIN-14	railet.					0	
Box/Carton/Bundle/Inner Pack	-		000701					1				
x Case	24		103707	700271305			COST INFORMATION			WHOLESAL	ER USE ONL	Y:
Pallet												
						Regular Cost			Vendor #:			
						Invoice Cost (W	/AC) (\$)	\$60.00	Whsl. Code			
	_					As of date:			Fineline Co	ae:		
						i i As oi date:						
		Attach copy of SAFETY DATA S	SHEET (SDS) or non baza	ard letter PACKAGE INSI	ERT LABEL AND PHOTO OF		SING and BARCODE					
*Please provide any additional in	formation on page 2.	Attach copy of SAFETY DATA S	SHEET (SDS) or non haza		ERT, LABEL AND PHOTO OF	PRODUCT PACKAG	GING and BARCODE.					



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For Designated Drop Ship Only Products, Please Use Page 3

MATERIAL HA	ZARD CLASSIFICATION and TRANSPORTATION						
Is this product (check all that apply): a. Cytotoxic? No	SDS Hazard Classification						
b. CA Prop. 65 Carcinogen or Reproductive Toxicant?	SDS Hazard Classification						
Is the product a CA Prop 65 carcinogen?	Organic Corrosive						
Is the product a CA Prop 65 reproductive toxicant?	Inorganic Oxidizer						
Does the product label bear a CA Prop 65 warning?	Steroid/Androgen Contact Hazard						
c. Contact Hazard?	Aerosol Class; Identify NFPA Storage Level:						
d. Does this product require special clean-up instructions?							
(If yes, attach SDS with special instructions.)	Is the product a NIOSH hazardous drug? No						
e. Does the product contain DEHP?	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	Hazardous Waste Identification						
b. Proper Shipping Name							
c. DOT Hazard Class	EPA Hazardous Waste Code: Waste Characteristics						
d. Packing Group							
e. Inhalation Hazard?							
Is this product regulated for shipment by IATA?	REMS or REGISTRY RESTRICTIONS						
(if yes, answer a-e below and provide SDS)	Is there a REMS on this product? No						
a. UN/Identification Number	If Yes, is it managed with a pharmacy registry?						
b. Proper Shipping Name	Website URL:						
c. DOT Hazard Class							
d. Packing Group							
e. Inhalation Hazard?	Med Guide Required No						
Is the product restricted for air shipment? If so, indicate restriction:	Limited Distribution Requirement No						
Passenger	Comments / Details: (For example, iPledge program?)						
Cargo							
Passenger & Cargo							
Is this a reportable quantity? No	REMS:						
RQ Threshold:	REMS Program Manager Name: Phone:						
Is this a marine pollutant? No	Supplier Manages REMS registry exclusively:						
Is this product shipped utilizing an authorized DOT exception or Special Permit?	Wholesale distributor support:						
No (if yes, identify method below)	Provider Name: DEA #:						
Limited Quantity	Site Enrollment Number assigned PCPDP#:						
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:						
	Registry Program Contact Name: Phone:						
ADD'L STORAGE INFORMATION	Comments						
Is the Product							
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:	Contact tel. # if product received damaged:						
Schedule No. Is it a scheduled listed chemical product?: No	ls product returnable for credit:						
CLASS OF TRADE RESTRICTION:	URL/Link to returns policy:						
Restricted to retail pharmacy only:	Special regulations or returns requirements for this						
Restricted to hospital, clinics, and physician offices only:	product in certain states?						
Restricted from US territories? (explain in comments)	If so, which states? Other requirements? Comments?						
Comments:	1						
MISCELLANE	OUS 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 Designated Drop Ship Pr	oduct	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 Number:		Shipping lead time of PO: Hours Days
d. Phone only Phone No.:		
e. Supplier Web Site only Site Address: Minimum Order Quantity:		Ships same day for next day receipt: Ships for second day receipt:
Supplier's Customer Service Number:		Ships regular ground for 3-10 days receipt:
Contracted 3PL company / contact #: Name:		Chips regular ground for 5-10 days receipt.
Phone:		
Expedited Freight Charges or Other Designated Dr	op Ship Fees:	Overnight and Priority Overnight PO Processing
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, hospitals, clinics and	nhysician offices	Saturday Overnight receipt available:
Restricted to retail pharmacy only:	priysician onices	PO Receipt Cut off time:
Restricted to hospital, clinics, and physician offices only:		Phone #
Restricted from US territories? (explain in comments)		Order receipt method: Fax: Fax#:
Comments:		EDI:
		Overnight Fees apply:
		Other fees apply:
Other Data Information Required to Proces	s 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 #		Special regulations or returns requirements for this product in certain states?
Physician/Clinic DEA #: Physician/Clinic Specialty:		If so, which states? Other requirements? Comments?
		il so, which states? Other requirements? Comments?
Miscellaneous Notes:		
		ADDITIONAL INFORMATION
		Is product order for scheduled patient procedure?
		Is product order for restocking purposes?