

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

Version 2021				Introduction Type:	New Item		Final Version			Date:	4/19/2024	
		PRODUCT INFORMATION	N				SPECIAL HAN	DLING AND STOR	AGE REQUIF	REMENTS*		
Company Name:	XIROMED LLC			Application:	ANDA	a. Temperature – Indi	cate the USP tempe	erature range for the	his product.			
Application Number for NDA/AN		ed device):	215351			Temper	ature Range	Controlled Room -	- between 20	and 25 C (68	° – 77° F)	
Medical Device Class, if applicat												
DUNS:	080228637	TESTOSTERONE CYPIONATE INJEC					emperature Range F	Requirement	NO			
Proprietary Name (If Applicable) a Selling Unit NDC:	70700-290-22	Unit of Use NDC:	TION, USP	UPC: 370	700290224	(W Notes	rrite in)					
UDI	10100 200 22	CVX Code:		MVX Code:	100230224	140(65						
Description:	TESTOSTERONE CYPIONAT					ls this n	roduct to be shipped	to customers on in	?		No	
Description							roduct to be shipped				No	
Active Ingredient(s):	TESTOSTE	ERONE CYPIONATE, USP										
						b. Contact for tempera	ature excursion que	estions:				
URL for Additional Product Inform Address:	180 PARK AVE			Address 2: #10		Name: Numbe			VIPUL GANE 973-953-786			
City:	FLORHAM PARK		State:		: 07932	Group			VIPUL.GANE		ED.COM:	
Key Contact:	DAVID HERNANDEZ		Email:	DAVID.HERNANDEZ								
Phone Number:	844-947-6633		Fax:	862-286-0932		c. Special regulations	for product in any	states?			No	
Product Therapeutic Classification	n:					Special	returns requirement	s for this product?			No	
		DUCT INFORMATION		PRODUCT DESC	RIPTION INFORMATION		of colo) us stated				Ver	
The second second second	ADDITIONAL PROL		ireat Chin Ont	PRODUCT DESC	RIFTION INFORMATION	d. Store product (unit		1.) for a 1			Yes	
The product is? a legend device?	No		lirect-Ship Only leither		10 mL multi dose	Protect e. Shelf life:	product (unit of sa	ile) from light?			Yes 24	Months
if yes, enter class #	INO	Orphan Drug Status		Size:	TO THE HIGH dose		helf life at launch (i	if different):			24	Months
a product kit?	No			Strength:	2000 mg/10 mL (200 mg/m		inon nio ut iuunon (i	in annon onty.				montho
if yes, list NDCs of		FDA Approval Status		Strength.				ORDER INFORM	IATION			
component parts				Dosage Form:	INJECTABLE		- ·		What is the			
reverse numbered? co-licensed?	No	Allergens Present		-		Unit of	Sale		Single Vial in		unit?	
latex-free?	Yes					X	Box/Carton		(Write-in, e.g) Vials)	
preservative-free?	No	oppers are not manufacture	ed with natural rubbe	Product Shape:			Ampule			5	,	
correctional institution block?				Product Color:			Glass		Minimum or	der quantity	?	Yes
opioid?	No		PAIN				Tube					
Cannabinoid? If Unit Dose, is item bar coded to u	No No	Country of Origin S	PAIN	Product Imprint:			Vial Liquid Sgl Vial Liquid Multi		If Yes, how i	many of whi	ch nackano t	vno?
hospital scanning?		Is this product covered under	r the				Vial Powder Sql			Each	cii package i	ype:
If Unit Dose, indicate NDC here:		Trade Agreements Act (TAA))? Yes				Vial Power Multi			Inner/Carton	/Pack	
							Other: Write In			Case		
		FOR GENERIC DRUG PRODU	JCTS									
			Au	uthorized Generic *If A	uthorized Generic, other		РН	ARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AO				on fields are not applicable	Rec. sell unit to custo	mer?		Rx billing ur	nit to pharma	acv:	
I. Generic Equivalent to What Brand?: DEPO-TESTOSTERONE						Single Vial in Carton X Each						
	(Write-in, e.g. 1 Vial)				Gram							
		S SUPPLY CHAIN SECURITY ACT (DSC	SA) INFORMATION			-				Milliliter		
Does supplier meet DSCSA definit	tion of manufacturer?	Yes	GLN:	37070000007			ITEM	I AND PACKING IN	FORMATION			
Is product exempt from DSCSA?		No										
If yes, select exemption:			GCP:				Weight Lbs.		ons (US msm			Saleable #
Other exemption - Write in: Is product repackaged?		No		riginal product purchase	d	Item/Each:	-	Depth	Width	Height	(Cube)	Pieces
Is product sold by manufacturer's	exclusive distributor?	No	direct from m			item/Each.	0.092	7.559055118	7.6771654	10.433071	605.45318	1
Has FDA granted waiver/exception		No	Provide sour	ce manufacturer for repart	ackaged product	Box/Carton/Bundle/					0	
If yes, attach documentation from	m FDA.					Inner Pack:						
		GTIN AND HIBCC PRODUCT INFO	RMATION			Case:	3.68	52.75590551	38.818898	47.086614	96429.906	40
						Pallet:					0	
Saleable Unit of Measure	Saleable Qua	ntity HIBCC		IN-14	Unit of Use GTIN-14						U	
x Item/Each	1		003	370700290224								
Box/Carton/Bundle/Inner Pack	40	_	103	370700290221		CO	ST INFORMATION		\	WHOLESALI	ER USE ONL	1.
Pallet	40		100	10100230221		Regular Cost			Vendor #:			
						Invoice Cost (WAC) (\$	i)	\$81.11	Whsl. Code			
									Fineline Coo	le:		
	-	_				As of date:						
		Attach copy of SAFETY DATA	SHEET (SDS) or non haza	ard letter, PACKAGE INSE	RT, LABEL AND PHOTO OF F	PRODUCT PACKAGING a	nd BARCODE.		I			
*Please provide any additional infe	ormation on page 2.	, ,	()		gnated Drop Ship Only.	Signatu						
L												

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

Version 2021 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? No 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 a Bel bear a CA Prop 65 warning? C. Contact Hazard? O. Does this product require special clean-up instructions? (If yes, attach SDS with special instructions.)	SDS Hazard Classification Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard Does the product have an Aerosol class? If yes, identify NFPA Storage Level: NFPA Storage Level: NFPA Storage Level: NFPA Storage Level:							
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) a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group	Is the product a NIOSH hazardous drug? No If yes, indicate which: Hazardous Waste Identification							
e. Inhalation Hazard?	EPA Hazardous Waste Code: Waste Characteristics							
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. Packing Group e. Inhalation Hazard? No Is the product restricted for air shipment? If so, indicate restriction: Passenger Cargo Passenger & Cargo Is this a reportable quantity? No RQ Threshold: 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) Special Permit; DOT-SP Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101); SP#	REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? No If Yes, is it managed with a pharmacy registry? No Website URL: No Med Guide Required No Limited Distribution Requirement No Comments / Details: (For example, iPledge program?) No REMS: Phone: REMS Program Manager Name: Phone: Supplier Manages REMS registry exclusively: DEA #: Provider Name: DEA #: Site Enrollment Number assigned NCPDP#: by Supplier: NPI #:							
	Registry Program Contact Name: Phone:							
ADD'L STORAGE INFORMATION	Comments							
Is the Product Controlled Substance? Yes Controlled Substance Code CIII Controlled by State(s)? Yes Listed Chemical (List I or II) No ARCOS Reportable? No If yes, indicate which: Is Schedule No. 3 Is it a scheduled listed chemical product?: No CLASS OF TRADE RESTRICTION: No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Restricted to retail pharmacy only: Is Initial Select YES if sold to retail pharmacy only:	RETURN INSTRUCTIONS Contact tel. # if product received damaged: Is product returnable for credit: URL/Link to returns policy: Special regulations or returns requirements for this							
Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	If so, which states? Other requirements? Comments?							
	OUS NOTES and/or Image of Product Barcode:							



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Version 2021 FOR DESIGNATED DROP SHIP PRODUCT ONLY - i	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 Fax Number: c. Fax Fax Number: d. Phone only Phone No.: e. Supplier Web Site only Site Address: 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: Image: Comparison of the second
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: Order receipt method: Phone: Fax: Fax #: EDI: Other fees apply:
Other Data Information Required to Process PO:	Return Instructions
Patient Procedure Date:	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?