

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

Version 2021 Introduction Type:								Fina	l Version			Date:	4/13/2	2023			
PRODUCT INFORMATION								SPECIAL HANDLING AND STORAGE REQUIREMENTS*									
Company Name: XIROMED LLC Application: ANDA							a. Temperature – Indicate the USP temperature range for this product.										
Application Number for NDA/ANDABLA (drug); PMA/510(k)(med device): 215351 Application Number for NDA/ANDABLA (drug); PMA/510(k)(med device): 215351 Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)																	
Medical Device Class, if applicable:																	
DUNS:	080228637								Other Temperature Range Requirement				No				
Proprietary Name (If Applicable) a		ne: TESTO	OSTERONE CYPIONATE IN	JECTION, USP					_	(write in))						
Selling Unit NDC: UDI	70700-288-22		Unit of Use NDC:			UPC:	3707002	288221	-	Notes							
	CVX Code: MVX Code:																
Description: TESTOSTERONE CYPIONATE INJECTION, USP 1000 mg/10 mL Is this product to be shipped to customers on ice?																	
Active Ingredient(s): Is this product to be shipped to customers on dry ice? No TESTOSTERONE CYPIONATE																	
Active ingredient(s): ESTOS LENONE CYPIONALE																	
URL for Additional Product Inforn	t Information:								Name:				VIPUL GANDHI				
Address:	180 PARK AVE				Address 2: #101			Number:				973-953-7867					
City:	FLORHAM PARK	RHAM PARK			State:	NJ	Zip:		Group E-mail:				VIPUL.GANDHI@XIROMED.COM;				
Key Contact:	DAVID HERNANDE				Email:		DAVID.HERNANDEZ@XIROMED.COM										
Phone Number:	844-947-6633				Fax:	862-286-0932	362-286-0932			c. Special regulations for product in any states?				No			
Product Therapeutic Classificatio	n:									Special return	s requiremen	ts for this product?			No		
	ADDITION	NAL PRODUCT INF	FORMATION			PRODUCT	DESCRIP.	TION INFORMATION			-)				V		
	ADDITION	NAL PRODUCT IN				PRODUCT DESCRIPTION INFORMATION			d. Store prod	luct (unit of sal			Yes				
The product is?			Is the Product	Direct-Ship C Neither	only		44	0 mL multi dose vial	e. Shelf life:	Protect produ	uct (unit of s	ale) from light?			Yes 24	Months	
a legend device? if yes, enter class #	-	No	Orphan Drug Status	Neither		Size:	10	u mL muiti dose viai	e. Shelf life:	Initial shelf lif	fo at launch /	if difforant):			24	Months	
a product kit?		No	Orphan Drug Status				10	00 mg/mL		illitiai Sileli III	ie at iauricii (ii dillerent).	24		24	Months	
if yes, list NDCs of	FDA Approval Status					Strength:	'	oo mgm.				ORDER INFORM	ATION				
component parts						Dosage For	rm:	NJECTABLE									
reverse numbered?	-	No				Dosage i oi				Unit of Sale			What is the		unit?		
co-licensed?	No Allergens Present							Bottle				1 Carton of 1 Vial					
latex-free?	Yes Vial stoppers are not manufactured natural rubber latex			with Product Shape:				Box/Carton (Write-in, e.g. 1 Box of 10 Vials) Ampule				0 Vials)					
preservative-free? correctional institution block?		No No	naturai ru	bber latex			_			Glas			Minimum o	rder quantity	,2 T	Yes	
opioid?		No				Product Co	olor:			Tube			wiiiiiiiiiiiiiiiiiiiiiiiiiiiiiiiiiiiii	idei quantiti	,. L	163	
Cannabinoid?	No. Country of Origin SPAIN							x Vial Liquid Sgl									
If Unit Dose, is item bar coded to u	If Unit Dose, is item bar coded to unit dose for							Vial Liquid Multi If Yes, how many of which package type?					type?				
hospital scanning?								Vial Powder Sql			40 Each						
If Unit Dose, indicate NDC here: Trade Agreements Act (TAA)? Ye				Yes				Vial Power Multi			Inner/Carton/Pack						
Other: Write In Case																	
			FOR GENERIC DRUG PRO	DDUCTS													
Authorized Generic *If Authorized Generic, other										PHARMACY ORDER / BILL UNIT							
acetian fields are not applicable																	
I. Orange Book Rating: II. Generic Equivalent to What Brand?: DEPO-TESTOSTERONE								1 Vial				Rx billing unit to pharmacy: x Each					
II. Generic Equivalent to What Brand?: DEPO-TESTOSTERONE									(Write-in, e.g. 1 Vial)				Gram				
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION														Milliliter			
Does supplier meet DSCSA defini	ition of manufacture	er?	Yes		GLN:	3707000000074	1				ITEM	AND PACKING IN	IFORMATION	١			
Is product exempt from DSCSA?			No														
If yes, select exemption:					GCP:					w	eight Lbs.		ons (US msn			Saleable #	
Other exemption - Write in:									1	**	J.g.ii. ED3.	Depth	Width	Height	(Cube)	Pieces	
Is product repackaged?			No			riginal product			Item/Each:		0.046	7.559055118	7.6771654	10.433071	605.45318	1	
Is product sold by manufacturer's Has FDA granted waiver/exception			No No			irect from mfr? rce manufacturer i	for ropock	raged product	Box/Carton/E	Pundle/							
If yes, attach documentation from		ductr	140		FIOVIUE SOUI	ce manufacturer	101 Tepack	ageu product	Inner Pack:	ouritie/					0		
,									Case:		2.94	52.75590551	20.040000	47.086614	06420.006	40	
		GTIN	AND HIBCC PRODUCT IN	IFORMATION							2.94	52.75590551	30.010090	47.000014	90429.900	40	
									Pallet:						0		
Saleable Unit of Measure	Sal	leable Quantity	HIBCC			IN-14	_	Unit of Use GTIN-14									
x Item/Each	1 00370700288221							COST INFORMATION WHOLESALER USE ONLY:									
Box/Carton/Bundle/Inner Pack X Case	40 10370700288				70700288228			COST INFORMATION				WHOLESALER USE UNLY:					
Pallet	-	40			103	77 30200220			Regular Cost	ŧ			Vendor #:				
	T I								Invoice Cost			\$41.40	Whsl. Code	#:			
	<u> </u>												Fineline Co				
	Į [As of date:								
 							- 1110		II		2005		l				
*Please provide any additional inf	formation on page 2		Attach copy of SAFETY DAT	IA SHEET (SDS) or non nazai			LABEL AND PHOTO OF Ited Drop Ship Only.	PRODUCT PACK	AGING and BAI Signature:	KCODE.						



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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 a. Cvtotoxic? No b. CA Prop. 65 Carcinogen or Reproductive Toxicant? Organic Is the product a CA Prop 65 carcinogen? Nο 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 Contact Hazard No Does the product have an Aerosol class? If yes, c. Contact Hazard? No identify NFPA Storage Level: d. Does this product require special clean-up instructions? Yes (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? Is the product a NIOSH hazardous drug? No 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? c. DOT Hazard Class If Yes, is it managed with a pharmacy registry? d. Packing Group Website URL: e. Inhalation Hazard? No Is the product restricted for air shipment? If so, indicate restriction: No Med Guide Required No Passenger Limited Distribution Requirement No Comments / Details: (For example, iPledge program?) Cargo Passenger & Cargo Is this a reportable quantity? REMS: RQ Threshold: REMS Program Manager Name: Phone: 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? (if ves. identify method below) Provider Name: DEA #: Limited Quantity Site Enrollment Number assigned NCPDP#: Consumer Commodity, ORM-D NPI#: by Supplier: 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 Code CIII RETURN INSTRUCTIONS Controlled Substance? Yes Controlled by State(s)? Yes 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 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 Restricted to retail pharmacy only: Special regulations or returns requirements for this product in certain states? Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) If so, which states? Other requirements? Comments? Comments: MISCELLANEOUS 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 Product	Standard Order Receipt and Processing						
Purchase orders may be accepted by: a. EDI b. Autofax Fax Number:	Purchase order daily receipt cut off time by supplier Cut off time:						
c. Fax d. Phone only e. Supplier Web Site only Site Address:	Shipping lead time of PO: Hours Days Ships same day for next day receipt:						
Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Phone:	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:	Overnight receipt available:						
Drop Ship service fee billed with each order:	PO Receipt cut off time:						
Drop Ship miscellaneous fees billed: Comments:	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: Order receipt method: 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:	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?						