

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

Version 2021						Introduction 1	Гуре:	New Item		Final Version			Date:	05.08.	.2023
PRODUCT INFORMATION							SPECIAL HAND	LING AND STOR	AGE REQUI	REMENTS*					
Company Name: Eugia US LLC (f/k/a AuroMedics Pharma LLC) Application: ANDA						ANDA	a. Temperature – Indicate the USP temperature range for this product.								
Application Number for NDA/AN		1A/510(k)(med de	evice):	21	5169				Tempera	ature Range	Cold – between 2	and 8 C (36°	– 46° F)		
Medical Device Class, if applicat															
DUNS:	968961354		and a state of the							emperature Range F	Requirement			GHT, STORE	
Proprietary Name (If Applicable) a	55150-0394-02	me: Fulv	vestrant Injection Unit of Use NDC:		55150 0204 0	2 UPC:	355150	204020		rite in)		ORIGINAL C	ARTON UN	TIL TIME OF	USE.
Selling Unit NDC: UDI	55150-0394-02		CVX Code:		55150-0394-02	MVX Code:	305100	594029	Notes						
_	Eulus strent lais stie	- 050			and the sector in it.									Na	
Description:	Fulvestrant injection	n 250 mg/5 mL (5	50 mg/mL) Both single-dose pre	efilied syringes	must be admin	istered to receive tr	ne 500 m	g dose.		oduct to be shipped oduct to be shipped				No No	
Active Ingredient(s):		Fulvestrant Inject	tion						15 1115 P					NO	
b. Contact for temperature excurs							ture excursion que	estions:							
URL for Additional Product Inform		Eugiaus.com				Address 2:			Name:			Kevin Cagne			
Address: City:	279 Princeton-High East Windsor	itstown Road			State:	NJ Zip: 08520			Number Group E	-		732.839.940 kcagnetti@		om	
Key Contact:					Email:		Ζιρ.	00320	Group L	-111a11.		KCagnettie		<u>.0111</u>	
Phone Number:	888-238-7880				Fax:	732-355-9449			c. Special regulations	for product in any	states?			No	
Product Therapeutic Classificatio	on:	Estrogen recepto	or antagonist		1					returns requirements				No	
-	I.				4						·				
	ADDITIO		INFORMATION			PRODUCT	DESCRIP	TION INFORMATION	d. Store product (unit	of sale) upright?					
The product is?			Is the Product	Direct-Ship C	Dnly				Protect	product (unit of sa	le) from light?				
a legend device?		No	Is the Product	Neither		Size:	2	2 PFS's	e. Shelf life:					24	Months
if yes, enter class #			Orphan Drug Status						Initial sh	nelf life at launch (i	f different):				Months
a product kit? if yes, list NDCs of		No	FDA Approval Status			Strength:		250 mg/5 mL 50 mg/mL)			ORDER INFORM				
component parts			PDA Approval Status				-				ORDER INFORM	IATION			
reverse numbered?		No				Dosage Forr	m: li	njection (Liquid)	Unit of S	Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present				_			Bottle		1 Box of 2 P	FSs		
latex-free?		Yes				Product Sha	npe: F	Prefilled Syringes	X	Box/Carton		(Write-in, e.	g. 1 Box of 2	PFS's)	
preservative-free?		Yes					.p.e.	ionica officigeo		Ampule					Ň
correctional institution block?		No				Product Col	or:			Glass Tube		Minimum o	der quantit	/?	Yes
opioid? Cannabinoid?		No No	Country of Origin	India			-			Vial Liquid Sgl					
If Unit Dose, is item bar coded to u			Obdinary of Origin	India		Product Imp	orint:			Vial Liquid Multi		If Yes. how	manv of wh	ich package f	type?
hospital scanning?			Is this product covered u	nder the						Vial Powder Sql			Each		.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
If Unit Dose, indicate NDC here:			Trade Agreements Act (T	AA)?	No					Vial Power Multi			Inner/Cartor	/Pack	
										Other: Write In		1	Case		
			FOR GENERIC DRUG PRO	DUCTS											
					Διι	thorized Generic	*If Autho	orized Generic. other		PHA		/ BILL UNIT			
I. Orange Book Rating:	AO					anonzeu Generie		fields are not applicable	Rec. sell unit to custor				nit to phorm	201/:	
II. Generic Equivalent to What Bra		Faslodex Injectio	n, 250 mg/5 mL (50 mg/mL)						Rec. sell unit to customer? Rx billing unit to pharmacy: 2 PFS's x						
						(Write-in, e.g. 2 PFS's)				Gram					
		DRUG SUPP	PLY CHAIN SECURITY ACT ([OSCSA) INFOR	RMATION							x	Milliliter		
Does supplier meet DSCSA defini	ition of monufacture	~~?	Yes	-	GLN:					ITEM	AND PACKING IN		1		
Is product exempt from DSCSA definition		err	No	-	GLN:						AND FACKING IN		l l		
If yes, select exemption:	Ĺ				GCP:						Dimensi	ons (US msn	ite)	Volume	Saleable #
Other exemption - Write in:					GCF.					Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		If yes, was or	iginal product			Item/Each:	0.00	•			. ,	4
Is product sold by manufacturer's			No		•	rect from mfr?				0.22	9.13	1.73	5	78.9745	
Has FDA granted waiver/exceptio		oduct?	No		Provide source	ce manufacturer f	or repacl	kaged product	Box/Carton/Bundle/	1.36	5.51	3.78	9.64	200.77999	5
If yes, attach documentation fro	om FDA.								Inner Pack:						
		GI	TIN AND HIBCC PRODUCT IN	FORMATION					Case:	6.83	18.89	12	10.23	2318.9364	20
									Pallet:	101.001	10	40			
Saleable Unit of Measure	Sa	leable Quantity	HIBCC		GTI	N-14		Unit of Use GTIN-14		191.361	48	40	50.5	96960	560
X Item/Each		1				55150394029									
Box/Carton/Bundle/Inner Pack		5				55150394020			COS	T INFORMATION		١	VHOLESAL	ER USE ONL	.Y:
Case		20				55150394024 55150394028			Bogular Coot			Vandar #			
Pallet		560			7035	00100394028	-		Regular Cost Invoice Cost (WAC) (\$	`	\$300.00	Vendor #: Whsl. Code	#•		
										,	ψ500.00	Fineline Co			
							1		As of date:	9/19/2023					
		_	Attach copy of SAFETY DAT	A SHEET (SD	S) or non hazar									0	
*Please provide any additional inf	formation on page 2	2.	Attach copy of SAFETY DAT	A SHEET (SD	S) or non hazar			, LABEL AND PHOTO OF F ated Drop Ship Only.	PRODUCT PACKAGING ar Signatu				Narender	Chamala	

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

Version 2021 For Designation	ted Drop Ship Only Products, Please Use Page 3					
MATERIAL H	AZARD 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? 	SDS Hazard Classification Organic Corrosive Inorganic Oxidizer 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.) No 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) No a. UN/Identification Number UN/Identification Number b. Proper Shipping Name UN/Identification Number c. DOT Hazard Class UN/Identification Number	Does the product have an Aerosol class? If yes, No identify NFPA Storage Level: NFPA Storage Level: Is the product a NIOSH hazardous drug? If yes, indicate which: Hazardous Waste Identification					
d. Packing Group	EPA Hazardous Waste Code: Waste Characteristics					
e. Inhalation Hazard? No Is this product regulated for shipment by IATA? No	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 e. Inhalation Hazard? No Is the product restricted for air shipment? If so, indicate restriction: Passenger Cargo Passenger & Cargo Is this a reportable quantity? RQ Threshold: Is this a marine pollutant? Is this product shipped utilizing an authorized DOT exception or Special Permit? (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# ADD'L STORAGE INFORMATION Is the Product	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: No REMS Program Manager Name: No Supplier Manages REMS registry exclusively: No Wholesale distributor support: Provider Name: Site Enrollment Number assigned DEA #: by Supplier: NPI #: Comments No Registry No Registry Program Contact Name: Phone: Comments Phone:					
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: If yes, indicate which: Schedule No. 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 Yes	Contact tel. # if product received damaged: Is product returnable for credit: URL/Link to returns policy: https://eugiaus.com/policies/return-policy/					
Restricted to retail pharmacy only: No Restricted to hospital, clinics, and physician offices only: No Restricted from US territories? (explain in comments) No Comments:	Special regulations or returns requirements for this product in certain states? If so, which states? Other requirements? Comments?					
MISCELLAN	EOUS NOTES and/or Image of Product Barcode:					



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Version 2021	FOR DESIGNATED DROP SHIP PRODUCT ONL	Y - 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 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:	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 Cha	rges 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 time: PO Receipt cut off time: Image: Comparison of time: Days of week overnight is available: Image: Monday Tuesday Image: Comparison of time: Image: Comparison of time: Image: Compariso
	ss of Trade Restriction:	Priority Overnight receipt available:
	narmacy, hospitals, clinics and physician offices	PO Receipt Cut off time: PO Receipt Cut off time: PO Receipt Cut off time: PO Receipt Cut off time: Order receipt method: Phone: Fax: Phone #: EDI: Fax #: Other fees apply: Image: Content for the state of the
Other Data Inf	ormation 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?
N	/liscellaneous Notes:	
		ADDITIONAL INFORMATION Is product order for scheduled patient procedure? Is product order for restocking purposes?