

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

Version 2024						Introduction 1	Гуре: Р	ost Launch Change		Final Version			Date:	13.1	2.24
			PRODUCT INFORMATI	ON						SPECIAL HAN	DLING AND STOR	AGE REQUI	REMENTS*		
Company Name:	Eugia US LLC					Applicat	tion:	ANDA	a. Temperature – Indica	ate the USP temp	erature range for 1	this product.			
Application Number for NDA/AN	IDA/BLA; PMA/510	(k):	214579						Tempera	ature Range	Controlled Room	– between 20	and 25 C (6	3° – 77° F)	
Medical Device Class, if applicat															
DUNS:	968961354									mperature Range	Requirement			to 77°F) [see	•
Proprietary Name (If Applicable) a Selling Unit NDC:	55150-373-25	ime:	Ephedrine Sulfate Injection USP, 50 Unit of Use NDC:		55150-373-01	UPC:	355150373	253	(wr Notes	ite in)		USP Contro	lled Room Te	imperaturej.	
UDI	00100 010 20		CVX Code:		00100 010 01	MVX Code:	000100010	200	Notes						
Description: Ephedrine Sulfate Injection USP, 50mg/mL [Single-Dose Vial] Is this product to b								oduct to be shipped	d to customers on i	ce?		No			
Description.	Ephedrine Suitate	injection 001 ,	Song/IIE [Single-Dose viai]								d to customers on a			No	
Active Ingredient(s):		Ephedrine Su	Ifate USP												
									b. Contact for temperat	ture excursion qu	estions:				
URL for Additional Product Inform Address:	279 Princeton-High	https://eug	laus.com/			Address 2:			Name:	_		Eugia US C 888-238-788		rice	
City:	East Windsor	nisiown Rodu			State:	NJ	Zip: 08	520	Number Group E					igial IS com	
Key Contact:		Email:					p.		Group E-mail: CustomerService@EugiaUS.com						
Phone Number:	888-238-7880				Fax:	732-355-9449			c. Special regulations f	or product in any	states?			No	
Product Therapeutic Classification	n:	Sympathomir	netic amine (alpha- and beta-adren	ergic agonist)					Special r	eturns requiremen	ts for this product?			No	
	ADDITIC	ONAL PRODU	CT INFORMATION			PRODUCT	DESCRIPTIC	IN INFORMATION	d. Store product (unit o	of sale) upright?				No	
The product is?			Is the Product	Direct-Ship O	nly					product (unit of s	ale) from light?				
a legend device? if yes, enter class #		No	Is the Product	Neither		Size:	25 x Vials	1 ml Single-Dose	e. Shelf life:	alf life at launah ((if different).			24	Months Months
a product kit?		No	Orphan Drug Status					, ig per mL	initial Sh	helf life at launch ((ir different):				wonths
if yes, list NDCs of			FDA Approval Status			Strength:		.9			ORDER INFORM	IATION			
component parts						Dosage Form	m: Solu	tion							
reverse numbered?		No							Unit of S	1			NDC selling	unit?	
co-licensed? latex-free?		No Yes	Allergens Present				Vial	Pack	x	Bottle Box/Carton		1 Box of 25	vials .g. 1 Box of 1	0 \/iale)	
preservative-free?		Yes				Product Sha	ape:	dok		Ampule		(11111111111111111111111111111111111111	.g. 1 D0x 01 1	0 viais)	
correctional institution block?		No				Product Col	or:			Glass		Minimum o	rder quantit	y?	Yes
opioid?		No				FIGULE	or.			Tube					
Cannabinoid?		No	Country of Origin	India		Product Imp	orint:			Vial Liquid Sgl Vial Liquid Multi					(
If Unit Dose, is item bar coded to u hospital scanning?	init dose for		Is this product covered un	der the						Vial Powder Sql		If Yes, now	Each	ich package	type?
If Unit Dose, indicate NDC here:			Trade Agreements Act (T		No					Vial Power Multi			Inner/Cartor	/Pack	
										Other: Write In		1	Case		
			FOR GENERIC DRUG PRO	DUCTS											
				_ [Aut	horized Generic		ed Generic, other Is are not applicable			ARMACY ORDER				
	AP	41(0)/47					section neit	is are not applicable	Rec. sell unit to custon				nit to pharm	acy:	
II. Generic Equivalent to What Bra	and?:	AKOVAZ							(Write-in, e.g. 1 Vial)	S		x	Each Gram		
		DRUG SI	UPPLY CHAIN SECURITY ACT (D	SCSA) INFORI	MATION				HCPCS J-Code:				Milliliter		
													1		
Does supplier meet DSCSA defini	tion of manufactur	rer?	Yes	_	GLN:					ITEM	AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			No												
If yes, select exemption:					GCP:					Weight Lbs.		ons (US msn		Volume (Cube)	Saleable # Pieces
Other exemption - Write in: Is product repackaged?			No		lf ves, was or	iginal product			Item/Each:		Depth	Width	Height	· /	
Is product sold by manufacturer's	s exclusive distribu	itor?	No			rect from mfr?				0.5437	3.54	3.54	1.97	24.687	25
Has FDA granted waiver/exception	n/exemption for pr	oduct?			Provide source	e manufacturer f	or repackag	ed product	Box/Carton/Bundle/	6.961	11.26	7.64	4.92	423.25	300
If yes, attach documentation from	m FDA.								Inner Pack:	0.501	11.20	7.04	4.52	420.20	000
			GTIN AND HIBCC PRODUCT INF						Case:	15.245	15.944	12.4	6.299	1245.35	600
				ORMATION					Pallet:						
Saleable Unit of Measure	RFID tag(Y/N)	Saleable	HIBCC		GTI	N-14	Un	it of Use GTIN-14		1130.707	48	40	55.39	106348.8	43200
		Quantity													
X Item/Each		25				5150373253									N.
X Box/Carton/Bundle/Inner Pack X Case		300				5150373254 5150373258	-		COS	T INFORMATION		,	WHOLESAL	ER USE ONL	.Y:
x Pallet		600 43200				5150373252			Regular Cost			Vendor #:			
									Invoice Cost (WAC) (\$)		\$300.00	Whsl. Code	#:		
												Fineline Co	de:		
							-		As of date:	12/17/2024		-			
┝┹─────			Attach copy of SAFETY DATA	A SHEET (SDS) or non hazar	Letter PACKAGE	INSERT 1 A	BEL AND PHOTO OF P	LI PRODUCT PACKAGING and	d BARCODE		1			
*Please provide any additional inf	ormation on page	2.	Auton copy of OAI LTT DATA	. 511221 (000	, or non nazdit			Drop Ship Only.	Signatur			-	D.Venkata Si	irender Redd	v
	on paye					200 p. 0 101			Signatu						,

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

Version 2024 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? No b. CA Prop. 65 Carcinogen or Reproductive Toxicant? Is the product a CA Prop 65 carcinogen? No Is the product a CA Prop 65 reproductive toxicant? Does the product label bear a CA Prop 65 warning? No c. Contact Hazard? (If ves, 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: No NFPA Storage Level: Image: Contact Hazard					
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	Is the product a NIOSH hazardous drug? No If yes, indicate which:					
d. Packing Group						
e. Inhalation Hazard? No	EPA Hazardous Waste Code: Waste Characteristics					
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 c. DOT Hazard Class d. Packing Group e. Inhalation Hazard? No	Is there a REMS on this product? No If Yes, is it managed with a pharmacy registry? Website URL:					
Is the product restricted for air shipment? If so, indicate restriction: No 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) Special Permit; DOT-SP Special Perovision (listed in Column 7 of 49 CFR 172.101);	REMS: No REMS Program Manager Name: Phone: Supplier Manages REMS registry exclusively: DEA Wholesale distributor support: DEA #: Provider Name: DEA #: Site Enrollment Number assigned NCPDP#: by Supplier: NPI #:					
SP#	Registry:					
ADD'L STORAGE INFORMATION	Registry Program Contact Name: Phone: Comments					
Is the Product Controlled Substance? No Controlled Substance Code	RETURN INSTRUCTIONS					
Controlled by State(s)? No Listed Chemical (List I or II) Yes ARCOS Reportable? No If yes, indicate which: Ephedrine Schedule No. Is it a scheduled listed chemical product?: Yes CLASS OF TRADE RESTRICTION:	Contact tel. # if product received damaged: Is product returnable for credit: URL/Link to returns policy:					
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices YES	http://auromedics.com/policies/return-policy/					
Restricted to retail pharmacy only: No Restricted to nospital, clinics, and physician offices only: No Restricted from US territories? (explain in comments) No	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 2024 FOR DESIGNATED DRO	DP SHIP PRODUCT ONLY - if not a designated drop ship, do not complete.
Order Method for Designated Drop Ship Prod	luct 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 #: 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: PO Receipt cut off time: Days of week overnight is available: Monday Tuesday Wednesday Thursday Friday
	Priority Overnight receipt available:
Class of Trade Restriction: No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and phy Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	ysician offices Saturday Overnight receipt available: Order receipt method: PO Receipt Cut off time: Order receipt method: Phone: EDI: Fax: Overnight Fees apply: Image: Content of the state of
Other Data Information Required to Process F	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?