

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

						Introduction Type		X	Final Version			Date:	3/17/	2020
			PRODUCT INFORMAT	TION					SPECIAL HAN	DLING AND STOR	AGE REQUIF	EMENTS*		
Company Name: Jubila	ant Cadista Ph	armaceuticals Inc.				Application:	ANDA	a. Temperature – Indica	ate the USP tempe	rature range for th	is product.			
Application Number for NDA/ANDA/BL						NDA 505(b) Type:	NOT APPLICABLE	-	ture Range	Controlled Room -		and 25 C (68	° – 77° F)	
Medical Device Class, if applicable:								1	3 -	-		· ·		
	94141							Other Te	mperature Range R	Requirement				
Proprietary Name (If Applicable) and Est		e: Escitalo	pram Tablets						ite in)					
	6-281-90		Unit of Use NDC:			UPC: 3-5	9746-281-90-8	Notes						
UDI			CVX Code:			MVX Code:								
Description: Escitalopram 20mg 90ct Tablets								Is this pro	oduct to be shipped	to customers on ic	e?		No	
									to customers on di			No		
Active Ingredient(s): Escitalopram						11								
						b. Contact for temperat	ure excursion que							
URL for Additional Product Information:			/products/full-product	-list				Name:			Customer Se			
	Township Line	ne Road			Address 2: Suite 325			Number:			(800) 313-46			
City: Yardio		•				State: PA Zip: 19067 Email: customer.service@cadista.com			-mail:		customer.s	ervice@ca	aista.com	
					customer.service@ N/A	vcaulsta.com	c. Special regulations f	or product in any	etates?			No		
Product Therapeutic Classification:		Anti-depressant			ı ax.	IVA		-					No	
Frouder Therapeutic Classification:	1	un-ucpressant						Special r	eturns requirements	a ioi una product?			INU	
	ADDITIO	NAL PRODUCT INFO	ORMATION			PRODUCT DES	CRIPTION INFORMATION	d. Store product (unit o	of sale) upright?				No	
The westerning	ADDITIO	WILL RODGOT IN		Disease Of the C	ali.	T RODGOT DES	JAN THER IN ORMATION	11		I-1 (I' 1				
The product is?	Ε.	11-	Is the Product	Direct-Ship Or Neither	nıy		00 count		product (unit of sa	ie) from light?			No 24	Months
a legend device? if yes, enter class #		No	Is the Product Orphan Drug Status	Meiniel		Size:	90 count	e. Shelf life:	elf life at launch (i	f different\.			24	Months Months
a product kit?	T.	No	Orphian Drug Status				20mg	Initial Sh	en me at launch (l'	i unierent):				WORTHS
if yes, list NDCs of		NU	FDA Approval Status			Strength:	Zomy			ORDER INFORM	ATION			
component parts			. S				TABLET							
reverse numbered?		No				Dosage Form:		Unit of S	ale		What is the	NDC selling	unit?	
co-licensed?	-	No	Allergens Present						Bottle		1 Bottle of 90	tablets		
latex-free?	-	Yes				Product Shape:	Oval, Biconvex, scorred		Box/Carton		(Write-in, e.	g. 1 Box of 1) Vials)	
preservative-free?		No				i roddot onape.			Ampule					
correctional institution block?		Yes				Product Color:	White		Glass		Minimum or	der quantity	?	Yes
opioid?	-	No		110			ID 41 4 10 I		Tube					
Cannabinoid?		No	Country of Origin	US		Product Imprint:	"B4" / "3"		Vial Liquid Sgl		W.V 1			
If Unit Dose, is item bar coded to unit dos	se for		Is this product covered ur	nder the		_			Vial Liquid Multi Vial Powder Sql			•	ch package t	ype?
hospital scanning? If Unit Dose, indicate NDC here:	-		Trade Agreements Act (T		Yes				Vial Powder Sql Vial Power Multi			Each Inner/Carton	/Pack	
ii onit bose, indicate NDC nere:				, .	100				Other: Write In			nner/Carton Case	n auk	
L						<u> </u>								
			FOR GENERIC DRUG PRO	ODUCTS										
			FOR GENERIC DRUG PRO	ODUCTS										
			FOR GENERIC DRUG PRO	ODUCTS	Aut	horized Generic *If	Authorized Generic, other		PH.	ARMACY ORDER	BILL UNIT			
L Orange Rook Rating:			FOR GENERIC DRUG PRO	ODUCTS	Aut		Authorized Generic, other tion fields are not applicable	Rec. sell unit to custom		ARMACY ORDER		it to pharm	acv.	
I. Orange Book Rating: II. Generic Equivalent to What Brand?:	li		FOR GENERIC DRUG PRO	ODUCTS	Aut			Rec. sell unit to custom	ner?	ARMACY ORDER	/ BILL UNIT Rx billing ur X	iit to pharma	асу:	
I. Orange Book Rating: II. Generic Equivalent to What Brand?:	<u>[i</u>	_exapro®	FOR GENERIC DRUG PRO	ODUCTS	Aut				ner?	ARMACY ORDER	Rx billing ur		acy:	
	Į.	_exapro®	FOR GENERIC DRUG PRO					1 Bottle of 90	ner?	ARMACY ORDER	Rx billing ur	Each	асу:	
II. Generic Equivalent to What Brand?:		_exapro® DRUG SUPPLY	CHAIN SECURITY ACT (I	DSCSA) INFOR	MATION	sec		1 Bottle of 90 (Write-in, e.g. 1 Vial)	ner? I tablets]	Rx billing ur X	Each Gram Milliliter	acy:	
II. Generic Equivalent to What Brand?: Does supplier meet DSCSA definition of		_exapro® DRUG SUPPLY	' CHAIN SECURITY ACT (I	DSCSA) INFOR				1 Bottle of 90 (Write-in, e.g. 1 Vial)	ner? I tablets	ARMACY ORDER	Rx billing ur X	Each Gram Milliliter	асу:	
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II. Generic Equivalent to What Brand?: Does supplier meet DSCSA definition of Is product exempt from DSCSA?		_exapro® DRUG SUPPLY	Y CHAIN SECURITY ACT (I Yes No	DSCSA) INFOR	MATION GLN:	0359746000004		1 Bottle of 90 (Write-in, e.g. 1 Vial)	ner? I tablets	AND PACKING IN	Rx billing ur X IFORMATION	Each Gram Milliliter		Saleable # Pieces
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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. Cytotoxic? No b. CA Prop. 65 Carcinogen or Reproductive Toxicant? Is the product a CA Prop 65 carcinogen? Nο Organic Corrosive Is the product a CA Prop 65 reproductive toxicant? Oxidizer No Inorganic Does the product label bear a CA Prop 65 warning? Steroid/Androgen Contact Hazard No c. Contact Hazard? No Does the product have an Aerosol class? If yes, identify No NFPA Storage Level: d. Does this product require special clean-up instructions? No (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? No Is the product a NIOSH hazardous drug? 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 Is there a REMS on this product? b. Proper Shipping Name 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 Cargo Comments / Details: (For example, iPledge program?) Passenger & Cargo Is this a reportable quantity? REMS: RQ Threshold: REMS Program Manager Name: Phone: Is this a marine pollutant? Nο Supplier Manages REMS registry exclusively: Is this product shipped utilizing an authorized DOT exception or Special Permit? Wholesale distributor support: (if yes, identify method below) Provider Name: DEA #: Site Enrollment Number assigned NCPDP#: Limited Quantity 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: Phone: Registry Program Contact Name: ADD'L STORAGE INFORMATION Comments Is the Product... RETURN INSTRUCTIONS Controlled Substance? No Controlled Substance Code 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 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 Yes 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 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:	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: 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: Miscellaneous Notes:	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?					
	ADDITIONAL INFORMATION					
	Is product order for scheduled patient procedure? Is product order for restocking purposes?					