

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

Version 2021						Introduction Ty	/pe: Post Launch Chan	le	x	Final Version			Date:	7/12	/2021
			PRODUCT INFO	RMATION			1			SPECIAL HAN	DLING AND STOR	AGE REQUI	REMENTS*	1	
Company Name: Jubilant Cadista Pharmaceuticals Inc.					Application: ANDA			a. Temperature – Indicate the USP temperature range for this product.							
Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): 077563							•	ature Range	Controlled Room		and 25 C (68	8° – 77° F)			
Medical Device Class, if applicat	Medical Device Class, if applicable:														
DUNS:	022490515								Other T	emperature Range	Requirement				
Proprietary Name (If Applicable) a		ime:	Cyclobenzaprine HCI Tablets							rite in)					
Selling Unit NDC:	59746-177-10		Unit of Use	IDC:		UPC: MVX Code:	3-59746-177-10-2		Notes						
UDI			CVX Code			WIVA Code:									1
Description: Cyclobenzaprine Hydrochloride 10mg 1000ct Tablet										roduct to be shippe				No	
Active Ingredient(s): Cyclobenzaprine Hydrochloride									a to customers on a	ary ice?		No			
b. Contact for temperature excursion questions:															
URL for Additional Product Inform		www.cadis	ta.com/products/full-pr	duct-list					Name:			Customer S (800) 313-4			
Address:	207 Kiley Drive				Address 2:				Number:						
City: Key Contact:	Salisbury Jackie Emershaw					MD Jackie.Emersha	Zip: 21801		Group E-mail: <u>customer.service@cadista.co</u>					idista.com	
Phone Number:	(410) 912-3722					(215) - 443 - 9646		c. Special regulations for product in any states?			states?			No	1
Product Therapeutic Classification		Analgesic M	uscle Relaxant			(210) 110 0010		0.00	-	returns requiremen				No	
									opoola	rotanio roquironion	to for the product.				
	ADDITI	ONAL PRODU	ICT INFORMATION			PRODUCT D	ESCRIPTION INFORMATIO	d. Sto	ore product (unit	of sale) upright?				No]
The product is?			Is the Product	Direct-Ship	Only				Protect	product (unit of s	ale) from light?			No	
a legend device?		No	Is the Product	Neither		Size:	1000 count	e. Sh	elf life:					24	Months
if yes, enter class #			Orphan Drug Stat	IS		5126.			Initial s	helf life at launch (if different):				Months
a product kit?		No				Strength:	10mg				ORDER INFORM	ATION			
if yes, list NDCs of component parts			FDA Approval Sta	us			TABLETS				ORDER INFORM	IATION			
reverse numbered?		No				Dosage Form	:		Unit of	Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Presen						X	Bottle		1 bottle of 1	-		
latex-free?		Yes				Product Shap	Round			Box/Carton		(Write-in, e	g. 1 Box of 1	0 Vials)	
preservative-free?		No				i i ouuor onup				Ampule				_	
correctional institution block? opioid?		Yes				Product Colo	r: Yellow			Glass Tube		Minimum o	rder quantity	?	Yes
Cannabinoid?		No No	Country of Origin	US			TL 177			Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	init dose for	140	,g			Product Impri	int:			Vial Liquid Multi		If Yes, how	many of whi	ch package	type?
hospital scanning?			Is this product cov							Vial Powder Sql		12	Each		
If Unit Dose, indicate NDC here:			Trade Agreements	Act (TAA)?	Yes					Vial Power Multi			Inner/Carton	/Pack	
										Other: Write In			Case		
			FOR GENERIC DRU	PRODUCTS											
					A	uthorized Generic	*If Authorized Generic, other		PHARMACY ORDER / BILL UNIT						
I. Orange Book Rating:	AB				I		section fields are not applica	le Rec.	sell unit to custo	mer?		Rx billing u	nit to pharm	acv.	
II. Generic Equivalent to What Bra		Flexeril							1 bottle of 1			X	Each		
								(Writ	te-in, e.g. 1 Vial)		_		Gram		
		DRUG S	SUPPLY CHAIN SECURITY	CT (DSCSA) INFO	RMATION								Milliliter		
Does supplier meet DSCSA definit	tion of manufactur	er?	Yes		GLN:	0359746000004				ITEN	I AND PACKING IN	NFORMATIO	N		
Is product exempt from DSCSA?			No		02.11										
If yes, select exemption:					GCP:	0359746					Dimensi	ons (US msr	nts.)	Volume	Saleable #
Other exemption - Write in:										Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No			riginal product		Item/	'Each:	0.4	2.4	2.4	4.75	27.36	1
Is product sold by manufacturer's			No		-	irect from mfr?									
Has FDA granted waiver/exception If yes, attach documentation from		oduct?	No		Provide soul	rce manufacturer for	r repackaged product		Carton/Bundle/ Pack:					0.00	
in yes, adden documentation not								Case		5.40	40.05	7.5	5.40	004.07	10
			GTIN AND HIBCC PRODU	CT INFORMATION	l					5.16	10.25	7.5	5.13	394.37	12
Onland to the total and the second								Palle	et:					0.00	
Saleable Unit of Measure	S	aleable Quant	ity HIBCC			IN-14 359746177102	Unit of Use GTIN-14								
X Item/Each Box/Carton/Bundle/Inner Pack		1			003	00740177102			cos	ST INFORMATION			WHOLESALI	ER US <u>E ONL</u>	.Y:
X Case		12			403	359746177100									
Pallet	_								ılar Cost			Vendor #:			
								Invoi	ce Cost (WAC) (\$	i)	\$67.46	Whsl. Code			
	-				_				data			Fineline Co	de:		
	-				_			As of	uate:						
			Attach copy of SAFE	Y DATA SHEET (S	DS) or non haza	ard letter, PACKAGE	INSERT, LABEL AND PHOT	OF PRODUC	T PACKAGING a	nd BARCODE.					
*Please provide any additional info	ormation on page	2.		(-			Designated Drop Ship Only.		Signatu						

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

Version 2021 For Designated Drop Ship Only Products, Please Use Page 3							
MATERIAL HAZARD 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? No Does the product label bear a CA Prop 65 warning? No c. Contact Hazard? No d. Does this product require special clean-up instructions? (If yes, attach SDS with special instructions.) e. Does the product contain DEHP? No (If yes, answer a-e below and provide SDS) a. UN/Identification Number	SDS Hazard Classification Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard Does the product have an Aerosol class? If yes, identify No NFPA Storage Level: NFPA Storage Level: Is the product a NIOSH hazardous drug? No If yes, indicate which: No						
b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	Hazardous Waste Identification 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 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 Pervision (listed in Column 7 of 49 CFR 172.101); SP#	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: Phone: Wholesale distributor support: DEA #: Provider Name: DEA #: Site Enrollment Number assigned NCPDP#: by Supplier: NPI #:						
ADD'L STORAGE INFORMATION Is the Product Controlled Substance? No Controlled Substance Code	Comments 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 Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	Contact tel. # if product 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?						
MISCELLA	NEOUS NOTES and/or Image of Product Barcode:						



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Version 2021 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:	Overnight receipt available: PO Receipt cut off time: Days of week overnight is available: Monday Tuesday Wednesday Thursday Friday
Class of Trade Restriction:	Priority Overnight receipt available: 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)	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:	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?