

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

						Introduction Type	: New Item		X Final Version			Date:	1/20	/2023
			PRODUCT INFORMAT	TION					SPECIAL HANI	DLING AND STOR	AGE REQUI	REMENTS*		
Company Name: Jubilant Cadista Pharmaceuticals Inc.				Application	: ANDA	a Temperature	a. Temperature – Indicate the USP temperature range for this product.							
Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): 211320						Application	. 704071		Temperature Range Controlled Room – between 20 and 25 C (68° – 7					
Medical Device Class, if applical		MA/3 TO(K)(IIIed device	=).	21	1020			- I	emperature reange	Controlled (Contr	Detween 20	ana 20 0 (00	, ,, ,,	
DUNS:	022490515							<u>-</u>	Other Temperature Range F	2 oguiromont				
Proprietary Name (If Applicable) a		Chlorth	alidone Tablets						(write in)	requirement				
Selling Unit NDC:	59746-760-36	ine. Chiota	Unit of Use NDC:			UPC: 3-	59746-760-36-6	- I	lotes					
UDI	33740-700-30		CVX Code:			MVX Code:	39740-700-30-0	- "	votes					
			OVA Code.			MIVX Gode.		-						
Description:	Chlorthalidone 25	mg 1000ct Tablets							s this product to be shipped				No	
		lou a ni						_ Is	s this product to be shipped	to customers on d	Iry ice?		No	
Active Ingredient(s):		Chlorthalidone						11						
		P. v	1 1 1 16 11 1					-	emperature excursion que	estions:	0			
URL for Additional Product Inform Address:		www.cadista.con	n/products/full-product	<u>t-list</u>		Address 2:			lame:		Customer Se			
	207 Kiley Drive				State:		24.004	-	lumber:		(800) 313-46		areas areas	
City: Key Contact:	Salisbury Customer Service				Email:		ip: 21801	-	Group E-mail:		customer.	service@ca	idista.com	
Phone Number:	(800) 313-4623	•			Fax:	<u>customer.service</u> (215) - 443 - 9646	wcauista.com	o Special regul	ations for product in any	ototoo?			No	1
		Oral I I mantanaire			I ax.	(213) - 443 - 9040								
Product Therapeutic Classificatio	n:	Oral Hypertensive						5	Special returns requirement	s for this product?			No	
	ADDITI	ON AL DRODUCT INC	ORMATION			PRODUCT DEC	ACRIPTION INFORMATION	.						1
	ADDITI	ONAL PRODUCT INF	ORMATION			PRODUCT DES	CRIPTION INFORMATION	d. Store produc	t (unit of sale) upright?				No	
The product is?			Is the Product	Direct-Ship C	Only			P	Protect product (unit of sa	le) from light?			No	
a legend device?		No	Is the Product	Neither		Size:	1000 count	e. Shelf life:					36	Months
if yes, enter class #			Orphan Drug Status			0.20.		Ir	nitial shelf life at launch (i	f different):				Months
a product kit?		No				Strength:	25mg							
if yes, list NDCs of			FDA Approval Status							ORDER INFORM	IATION			
component parts						Dosage Form:	Tablet							
reverse numbered?		No							Init of Sale		What is the		unit?	
co-licensed?		Yes	Allergens Present						X Bottle		1 Bottle of 1			
latex-free?		Yes				Product Shape:	Flat Round Tablet		Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free?		Yes				•	11.1.26.0		Ampule				_	
correctional institution block?		Yes				Product Color:	Light Yellow	_	Glass		Minimum o	der quantity	?	Yes
opioid?		No	Occupation of October	India			14001	_	Tube					
Cannabinoid?		No	Country of Origin	muia		Product Imprint	: "103"	_	Vial Liquid Sgl		W. W I			
If Unit Dose, is item bar coded to unhospital scanning?	init dose for		Is this product covered u						Vial Liquid Multi		If Yes, how	many or wni Each	сп раскаде	type?
I I														
					NI.			_	Vial Powder Sql		12		(D I	
If Unit Dose, indicate NDC here:		,	Trade Agreements Act (1		No				Vial Power Multi		12	Inner/Carton	/Pack	
If Unit Dose, indicate NDC here:		'	Trade Agreements Act (1	TAA)?	No						12		/Pack	
If Unit Dose, indicate NDC here:		,		TAA)?	No				Vial Power Multi		12	Inner/Carton	/Pack	
If Unit Dose, indicate NDC here:			Trade Agreements Act (1	TAA)?		***************************************	Authorizad Quantin althou		Vial Power Multi Other: Write In	ADMACY ODDER		Inner/Carton	/Pack	
If Unit Dose, indicate NDC here:			Trade Agreements Act (1	TAA)?			Authorized Generic, other		Vial Power Multi Other: Write In	ARMACY ORDER	/ BILL UNIT	Inner/Carton Case		
I. Orange Book Rating:	AB		Trade Agreements Act (1	TAA)?			Authorized Generic, other ction fields are not applicable	Rec. sell unit to	Vial Power Multi Other: Write In PH. customer?	ARMACY ORDER	/ BILL UNIT	Inner/Carton Case		
		Hygroton®	Trade Agreements Act (1	TAA)?				1 Bottle	Vial Power Multi Other: Write In PH. customer? e of 1000 tablets	ARMACY ORDER	/ BILL UNIT	Inner/Carton Case		
I. Orange Book Rating:		Hygroton®	Trade Agreements Act (1	ODUCTS	Au				Vial Power Multi Other: Write In PH. customer? e of 1000 tablets	ARMACY ORDER	/ BILL UNIT	Inner/Carton Case nit to pharm Each Gram		
I. Orange Book Rating:		Hygroton®	Trade Agreements Act (1	ODUCTS	Au			1 Bottle	Vial Power Multi Other: Write In PH. customer? e of 1000 tablets	ARMACY ORDER	/ BILL UNIT	Inner/Carton Case		
I. Orange Book Rating: II. Generic Equivalent to What Bra	nd?:	Hygroton® DRUG SUPPLY	Trade Agreements Act (1) FOR GENERIC DRUG PRO	ODUCTS	Au	se		1 Bottle	Vial Power Multi Other: Write In PH. customer? e of 1000 tablets Vial)		/ BILL UNIT Rx billing u	Inner/Carton Case nit to pharm Each Gram Milliliter		
I. Orange Book Rating: II. Generic Equivalent to What Bra Does supplier meet DSCSA defini	nd?:	Hygroton® DRUG SUPPLY	Trade Agreements Act (1) FOR GENERIC DRUG PRO CHAIN SECURITY ACT (1) Yes	ODUCTS	Au			1 Bottle	Vial Power Multi Other: Write In PH. customer? e of 1000 tablets Vial)	ARMACY ORDER	/ BILL UNIT Rx billing u	Inner/Carton Case nit to pharm Each Gram Milliliter		
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I. Orange Book Rating: II. Generic Equivalent to What Bra Does supplier meet DSCSA defini Is product exempt from DSCSA? If yes, select exemption: Other exemption - Write in: Is product repackaged? Is product sold by manufacturer's Has FDA granted waiver/exceptio If yes, attach documentation from Saleable Unit of Measure X	nd?: tion of manufactu exclusive distribi	Hygroton® DRUG SUPPLY rer? inter? roduct? GTIN taleable Quantity 12	Trade Agreements Act (1) FOR GENERIC DRUG PRO CHAIN SECURITY ACT (1) Yes No No No No No HIBCC PRODUCT IN	DSCSA) INFORMATION	RMATION GLN: GCP: If yes, was or purchased di Provide source GTIII 0038	8904184010027 0359746 iginal product rect from mfr? be manufacturer for re 14-14 19746760366 59746760364	ction fields are not applicable	Item/Each: Box/Carton/Bun Inner Pack: Case: Pallet: Regular Cost Invoice Cost (W As of date:	Vial Power Multi Other: Write In PH. customer? e of 1000 tablets Vial) Weight Lbs. 0.26 adde/ 3.91 COST INFORMATION AC) (\$)	Dimension Depth 2.2 9.25	/ BILL UNIT Rx billing u X IFORMATION ons (US msn Width 2.2 7.09	Inner/Carton Case hit to pharm Each Gram Milliliter Height 4.53 5.71	Volume (Cube) 21.93 0.00 374.48	Pieces 1 12



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 SDS Hazard Classification 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? No Inorganic Oxidizer 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 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 REMS: Is this a reportable quantity? 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: No Special regulations or returns requirements for this product in certain states? No Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) No If so, which states? Other requirements? Comments? Comments: MISCELLANEOUS NOTES and/or Image of Product Barcode



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

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 #:	Fax Number: Fax Number: Phone No.: Site Address: Name: Phone: Phone: rges or Other Designated Drop Ship Fees:	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: Overnight and Priority Overnight PO Processing					
Expedited freight fees billed with each orde		Overnight receipt available:					
Drop Ship service fee billed with each order Drop Ship miscellaneous fees billed: Comments:	T	PO Receipt cut off time: Days of week overnight is available: Monday Tuesday Wednesday Thursday Friday Priority Overnight receipt available:					
Cla	ss of Trade Restriction:	PO Receipt Cut off time:					
No restriction: Select YES if sold to retail ph Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician Restricted from US territories? (explain in contemporary)	narmacy, hospitals, clinics and physician offices a offices only: comments)	Saturday Overnight receipt available: PO Receipt Cut off time: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:					
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:	//iscellaneous 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?					