

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

Version 2021						Introduction 1	Туре:	New Item		Х	Final Version			Date:	6/17	/2024
PRODUCT INFORMATION							SPECIAL HANDLING AND STORAG			AGE REQUI	GE REQUIREMENTS*					
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): 211320 Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)																
Medical Device Class, if applica	ıble:															
DUNS:	022490515										mperature Range R	Requirement				
Proprietary Name (If Applicable) a		lame: Chlorth	halidone Tablets								te in)					
Selling Unit NDC:	59746-761-01		Unit of Use NDC:			UPC:	3-5974	46-761-01-1		Notes						
UDI CVX Code: MVX Code:										1						
									No							
Active Ingredient(s): Sthis product to be shipped to customers on dry ice? No									1							
b. Contact for temperature excursion questions:																
URL for Additional Product Inform	mation:	www.cadista.com	m/products/full-product	-list						Name:			Customer S			
Address:	790 Township Lir	ne Road				Address 2:	Suite 3			Number:			(800) 313-46			
City:	Yardley			State:	PA		19067	Group E-mail:			customer.service@cadista.com					
Key Contact: Phone Number:				Email: Fax:		customer.service@cadista.com N/A		c. Special regulations for product in any states?			No			1		
		Oral Hypertensive			ı ax.	IVA			c. Special re	-						
Product Therapeutic Classification: Oral Hypertensive Special returns requirements for this product? No																
ADDITIONAL PRODUCT INFORMATION PRODUCT DESCRIPTION INFORMATION d. Stor									d. Store proc	luct (unit c	of sale) upright?				No	1
The product is?			Is the Product	Direct-Ship O	nlv					•	product (unit of sa	la) from light?			No	1
a legend device?		No	Is the Product	Neither	,			100 count	e. Shelf life:	riotect	orounce (unit or sa	ie) iroin light:			36	Months
if yes, enter class #		1.10	Orphan Drug Status			Size:		100 oount	0.0.0	Initial sh	elf life at launch (i	f different):				Months
a product kit?		No				Strength:		50mg			•					
if yes, list NDCs of			FDA Approval Status			ou engui.						ORDER INFORM	IATION			
component parts		1				Dosage Form	m:	Tablet					140	NDOIII		
reverse numbered? co-licensed?		No	Allergens Bresent							Unit of S	Bottle		1 Bottle of 1	NDC selling	unit?	
latex-free?		Yes Yes	Allergens Present					Flat Round Tablet			Box/Carton			.g. 1 Box of 1	0 Vials)	
preservative-free?		Yes				Product Sha	ape:	riat Round Tablet			Ampule		(vviito iii, o.	g. I Dox of I	o viais)	
correctional institution block?		Yes				Product Col		Light Green			Glass		Minimum o	rder quantity	/?	Yes
opioid?		No				Product Col	or:				Tube					
Cannabinoid?		No	Country of Origin	India		Product Imp	orint:	'I' / '3'			Vial Liquid Sgl					
If Unit Dose, is item bar coded to	unit dose for		In this was dead account to	a da a tha							Vial Liquid Multi			many of whi	ich package	type?
hospital scanning? If Unit Dose, indicate NDC here:			Is this product covered u Trade Agreements Act (1		No						Vial Powder Sql Vial Power Multi		24	Each Inner/Carton	n/Pack	
il Offit Dose, indicate NDC here.			Trade Agreements Act (1	AA):	INU						Other: Write In			Case	// ack	
			FOR GENERIC DRUG PR	ODUCTS										1		
					Αι	thorized Generic		thorized Generic, other			PH	ARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating: AB					section fields are not applicable			Rec. sell unit to customer?				Rx billing unit to pharmacy:				
II. Generic Equivalent to What Bra	and?:	Hygroton®								Bottle of 100) tablets		Х	Each		
(Write-in, e.g						. 1 Vial)				Gram						
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION Milliliter																
Does supplier meet DSCSA defin	ition of manufactu	ırer?	Yes	7	GLN:	8904184010027					ITEM	AND PACKING II	NEORMATIO	N		
Is product exempt from DSCSA?			No		ULIT.	3304104010027						TAIL THORING II				
If yes, select exemption:					GCP:	0359746						Dimensi	ons (US msr	nts)	Volume	Saleable #
Other exemption - Write in:					GOI .	0339740					Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		If yes, was o	riginal product pur	chased		Item/Each:		0.085	1.54	1.54	3.35	7.94	1
Is product sold by manufacturer's			No		direct from n	nfr?					0.085	1.54	1.54	3.35	7.94	1
Has FDA granted waiver/exception		product?	No		Provide sour	ce manufacturer fo	or repac	kaged product	Box/Carton/E	Bundle/					0.00	
If yes, attach documentation fro	om FDA.								Inner Pack:							
		GTI	N AND HIBCC PRODUCT II	JEORMATION					Case:		2.98	9.45	6.5	4.33	265.97	24
		011	IN AND HIBCC PRODUCT II	II OKWATION					Pallet:							
Saleable Unit of Measure	:	Saleable Quantity	HIBCC		GTI	N-14		Unit of Use GTIN-14	l anot.						0.00	
X Item/Each		1			003	59746761011										
Box/Carton/Bundle/Inner Pack										cos	TINFORMATION			WHOLESAL	ER USE ONL	.Y:
X Case		24			403	59746761019	_		11							
Pallet							_		Regular Cost Invoice Cost			¢22.00	Vendor #: Whsl. Code	. 4.		
									invoice cost	(4) (0711)		\$22.00	Fineline Co			
									As of date:				1			
													1			
							-						<u> </u>			
1			Attach copy of SAFETY DA	TA SHEET (SD	S) or non haza			T, LABEL AND PHOTO OF F	PRODUCT PACK	AGING and	BARCODE.					
*Please provide any additional int	formation on page	2				See new n 3 for	r Design	nated Drop Ship Only		Signatur	е.					



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

Version 2021

For Designated Drop Ship Only Products, Please Use Page 3

MA	TERIAL HAZ	ARD 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?	SDS Hazard Classification Organic Corrosive								
Is the product a CA Prop 65 reproductive toxicant? Does the product label bear a CA Prop 65 warning?	Inorganic Steroid/Androgen	Oxidizer Contact Hazard							
c. Contact Hazard? d. Does this product require special clean-up instructions? (If yes, attach SDS with special instructions.) e. Does the product contain DEHP?	Does the product have an Aerosol class? If yes, identify NFPA Storage Level: NFPA Storage Level:	No							
Is this product regulated for shipment by DOT? (if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name	No	Is the product a NIOSH hazardous drug? If yes, indicate which:	No						
c. DOT Hazard Class				Hazardous Waste Identification					
d. Packing Group e. Inhalation Hazard?	d. Packing Group e. Inhalation Hazard? No			Waste Characteristics					
Is this product regulated for shipment by IATA?	No	EPA Hazardous Waste Code:		Tracto Charactoriorio					
(if yes, answer a-e below and provide SDS)	140	REMS o	r REGISTRY RESTRICTIONS						
a. UN/Identification Number									
b. Proper Shipping Name		Is there a REMS on this product?	No						
c. DOT Hazard Class d. Packing Group		If Yes, is it managed with a pharmacy registry? Website URL:							
e. Inhalation Hazard?	No	Woodle Cite.							
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		5540							
Is this a reportable quantity? No RQ Threshold:		REMS Program Manager Name:		Phone:					
Is this a marine pollutant? No		Supplier Manages REMS registry exclusively:		1 10.10.					
Is this product shipped utilizing an authorized DOT exception or Special Permit?		Wholesale distributor support:							
No (if yes, identify method below)	Provider Name:		DEA #: NCPDP#:						
Limited Quantity Consumer Commodity, ORM-D		Site Enrollment Number assigned by Supplier:		NCPDP#: NPI #:					
Small Quantity (49 CFR 173.4)		зу саррион							
Special Permit; DOT-SP		Comments							
Special Provision (listed in Column 7 of 49 CFR 172.101);									
SP#		Registry:		Phone:					
ADD'L STORAGE INFORMATION		Registry Program Contact Name: Comments		FIIOTIE.					
Is the Product									
Controlled Substance? No Controlled Substance Code		R	ETURN INSTRUCTIONS						
Controlled by State(s)? No Listed Chemical (List I or II)	No								
ARCOS Reportable? Schedule No. If yes, indicate which: Is it a scheduled listed chemical product?:	No	Contact tel. # if product received damaged:							
CLASS OF TRADE RESTRICTION:	140	Is product returnable for credit:							
	Yes	URL/Link to returns policy:							
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices	No								
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)	No No	If so, which states? Other requirements? Comments?							
Comments:	140	ii 30, wiilon states: Other requirements: Other ents:							
Commonto.									
MIS	SCELLANEC	DUS NOTES and/or Image of Product Barcode:							
- Wild		The red dilator image of Floudet 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 S	nip Product	Standard Order Receipt and Processing
Purchase orders may be accepted by: a. EDI		Purchase order daily receipt cut off time by supplier Cut off time:
b. Autofax c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Phone:	per:	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 Designa	ed Drop Ship Fees:	Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each order: Drop Ship service fee billed with each order:		Overnight receipt available: PO Receipt cut off time:
Drop Ship miscellaneous fees billed: Comments:		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, clinic Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	s and physician offices	Saturday Overnight receipt available: PO Receipt Cut off time: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:
Other Data Information Required to F	rocess 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?
Miscellaneous Notes:		
		ADDITIONAL INFORMATION
		Is product order for scheduled patient procedure? Is product order for restocking purposes?