

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

Version 2021						Introduction Ty	/pe: Post Launch Ch	ange		x Final Version			Date:	7/12	/2021
			PRODUCT INFORM	ATION						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): 201845								perature Range	Controlled Room		and 25 C (68	8° – 77° F)			
Medical Device Class, if applicable:															
DUNS:	022490515								Oth	er Temperature Range	Requirement				
Proprietary Name (If Applicable) a		ame:	Losartan - HCTZ Tablets							(write in)					
Selling Unit NDC:	59746-337-90		Unit of Use NDO	:		UPC: MVX Code:	3-59746-337-90-2		Note	es					
UDI			CVX Code:			WVX Code:									1
								is product to be shippe				No No			
Active Ingredient(s): Losartan Potassium; Hydrochlorothiazide								Is this product to be shipped to customers on dry ice? No							
b. Contact for temperature excursion questions:															
URL for Additional Product Inform		www.cadis	ta.com/products/full-produ	<u>ct-list</u>					Nan	ne:		Customer S			
Address:	207 Kiley Drive				0	Address 2:				nber:		(800) 313-46			
City: Key Contact:	Salisbury Jackie Emershaw	,			State:	MD Jackie.Emersha	Zip: 21801		Group E-mail: <u>customer.service@cadista.c</u>				idista.com		
Phone Number:	(410) 912-3722				(215) - 443 - 9646	w@jubi.com		c. Special regulations for product in any states?					No	1	
Product Therapeutic Classificatio	. ,	Antihyperten	sive			(210) 110 0010				cial returns requiremen				No	
									000		lo for the product.				
	ADDITI	ONAL PRODU	ICT INFORMATION			PRODUCT D	ESCRIPTION INFORMATI	ION	d. Store product (u	init of sale) upright?				No	1
The product is?			Is the Product	Direct-Ship	Only				Pro	tect product (unit of sa	ale) from light?			No	1
a legend device?		No	Is the Product	Neither		0	90 count		e. Shelf life:		ile) il elli ligitti			24	Months
if yes, enter class #		1	Orphan Drug Status			Size:			Initi	al shelf life at launch (if different):				Months
a product kit?		No				Strength:	50mg-12.5mg								
if yes, list NDCs of			FDA Approval Status			g	7404570				ORDER INFORM	IATION			
component parts reverse numbered?		No				Dosage Form	: TABLETS		Unit	of Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present							X Bottle		1 bottle of 9	-	unit:	
latex-free?		Yes	Anergens Fresent			Des des color	Oval			Box/Carton			g. 1 Box of 1	0 Vials)	
preservative-free?		No				Product Shap	e:			Ampule					
correctional institution block?		Yes				Product Colo	Yellow			Glass		Minimum o	rder quantity	?	Yes
opioid?		No	o	110						Tube					
Cannabinoid?	wit dooo for	No	Country of Origin	US		Product Impri	nt: C / 337			Vial Liquid Sgl Vial Liquid Multi		If Voc. how	mony of whi	oh naokaga	turno?
If Unit Dose, is item bar coded to u hospital scanning?	Init dose for		Is this product covered	under the						Vial Powder Sql			Each	ch package	type?
If Unit Dose, indicate NDC here:			Trade Agreements Act		Yes					Vial Power Multi			Inner/Cartor	/Pack	
			<u>`</u>		·					Other: Write In			Case		
			FOR GENERIC DRUG P	RODUCTS		-									
					Au		*If Authorized Generic, oth section fields are not appli				ARMACY ORDER				
I. Orange Book Rating:	AB						section neios are not appli	Cable	Rec. sell unit to cu		1		nit to pharm	acy:	
II. Generic Equivalent to What Bra	nd?:	Hyzaar								of 90 tablets		X	Each Gram		
		DRUG S	UPPLY CHAIN SECURITY ACT	(DSCSA) INFO	RMATION				(Write-in, e.g. 1 Via	u)			Milliliter		
				(1						
Does supplier meet DSCSA defini	tion of manufactu	rer?	Yes		GLN:	0359746000004				ITEN	I AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			No												
If yes, select exemption:					GCP:	0359746				Weight Lbs.		ons (US msn	-	Volume	Saleable #
Other exemption - Write in:										Weight Ebs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?	avaluaine diatriku		No	_		riginal product irect from mfr?			Item/Each:	0.12	1.9	1.9	4	14.44	1
Is product sold by manufacturer's Has FDA granted waiver/exception			No	_	-		repackaged product		Box/Carton/Bundle	2/					
If yes, attach documentation from		ouuor.			i i ovide sour		repairingen product		Inner Pack:					0.00	
									Case:	3.92	11.5	7.75	5	445.63	24
			GTIN AND HIBCC PRODUCT	INFORMATION						0.52	11.0	1.10		440.00	27
Saleable Unit of Measure					07				Pallet:					0.00	
X Item/Each	8	Saleable Quant	ity HIBCC		_	N-14 59746337902	Unit of Use GTIN-	14							
Box/Carton/Bundle/Inner Pack										COST INFORMATION			WHOL <u>ESAL</u>	ER USE ONL	.Y:
X Case		24			403	59746337900									
Pallet	_								Regular Cost			Vendor #:			
					_				Invoice Cost (WAC	;) (\$)	\$18.80	Whsl. Code			
	-				_				A (.]			Fineline Co	de:		
	-				-				As of date:						
			Attach copy of SAFETY D	ATA SHEET (SE	S) or non haza	rd letter, PACKAGE	NSERT, LABEL AND PHC	TO OF P	RODUCT PACKAGIN	G and BARCODE.		1			
*Please provide any additional inf	ormation on page	2.		(Designated Drop Ship On			nature:					

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: NFPA Storage Level: No 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?