

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

Version 2020				Introduction Type:	New Item		x Final Version			Date:	7/24/	/2023
		PRODUCT INFORMATION					SPECIAL HAN	DLING AND STO	RAGE REQUI	REMENTS*		
Company Name: Viona Pharmaceuticals Inc. ANDA							a. Temperature – Indicate the USP temperature range for this product.					
	IDA/BLA (drug); PMA/510(k)(med devi	ice): 20	5443				mperature Range	Controlled Room		and 25 C (68	° – 77° F)	
DUNS:	081468959					01	her Temperature Range F	Requirement				
Proprietary Name (If Applicable) a	and Established Name: Febuxe	ostat Tablets, 40 mg					(write in)					
Selling Unit NDC:	72578-136-06	Unit of Use NDC:		UPC: 372578	136062	No	otes					
UDI	N/A	CVX Code:		MVX Code: N/A								
Description: White to off-white, beveled-edge, oval-shaped tablets debossed with "401" on one side and plain on the other side. Is this product to be shipped to customers on ice? No is this product to be shipped to customers on dry ice? No										No No		
Active Ingredient(s):	b. Contact for temperature excursion questions:											
URL for Additional Product Inform	mation: www.vionausa.	com					ime:		Customer S	ervice		
Address:	20 Commerce Drive			Address 2: Suite 34	40	Number: 888-304-5022						
City:	Cranford		State:		07016	Group E-mail:			customerservice@vionausa.com			
Key Contact:	Chris Urbanski		Email:	Curbanski@vionausa	.com							
Phone Number:	908-956-0600		Fax:	908-514-4005			tions for product in any				No	
Product Therapeutic Classification	n: Xanthine Oxidase I	Inhibitors				Sp	ecial returns requirement	s for this product?			No	
		CORMATION										
	ADDITIONAL PRODUCT IN			PRODUCT DESCR	IPTION INFORMATION		(unit of sale) upright?				Yes	
The product is?		Is the Product Direct-Ship	Only				otect product (unit of sa	le) from light?			Yes	
a legend device?	No	Is the Product Neither		Size:	30 ct	e. Shelf life:					24	Months
if yes, enter class # a product kit?	No	Orphan Drug Status				In	itial shelf life at launch (i	f different):				Months
if yes, list NDCs of	NO	FDA Approval Status		Strength:	40 mg			ORDER INFORI	MATION			
component parts reverse numbered?	N.			Dosage Form:	Tablets		it of Colo		What is the	NDC selling	unit?	
co-licensed?	No No	Allergens Present					nit of Sale x Bottle		1 Bottle of 3		unit?	
latex-free?	Yes	Allergens i Tesent					Box/Carton			.g. 1 Box of 1) Vials)	
preservative-free?	Yes			Product Shape:	Oval		Ampule		(11110 111, 0	.g. 1 20/ 01 11	, that of	
correctional institution block?	No			Desident Onland			Glass		Minimum o	rder quantity	?	Yes
opioid?	No			Product Color:	White to off-white		Tube					
Cannabinoid?	No	Country of Origin India		Product Imprint:	"401"		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	init dose for hospital			rioddet imprint.	401		Vial Liquid Multi			many of whi	ch package	type?
scanning?		Is this product covered under the					Vial Powder Sql		24	Each		
If Unit Dose, indicate NDC here:		Trade Agreements Act (TAA)?	No				Vial Power Multi Other: Write In			Inner/Carton	/Pack	
		FOR GENERIC DRUG PRODUCTS					Other: write in		1	Case		
		FOR GENERIC DRUG PRODUCTS										
			Auth	prized Generic *If Author	prized Generic, other section		PH	ARMACY ORDER	R / BILL UNIT			
L Orenge Beek Beting:	AD		Addin		e not applicable	Rec. sell unit to				nit to pharm		
I. Orange Book Rating: II. Generic Equivalent to What Bra	AB and?: Uloric						1 Bottle	T		Each	acy:	
n. Generie Equivalent to Milat Bit						(Write-in, e.g. 1 \		l	^	Gram		
	DRUG SUPPL	Y CHAIN SECURITY ACT (DSCSA) INFO	RMATION			(Milliliter		
Does supplier meet DSCSA defin	ition of manufacturer?		LN:	0372578000004			ITEM	AND PACKING I	INFORMATIO	N		
Is product exempt from DSCSA?		No										
If yes, select exemption:					_		Weight Lbs.		ions (US msr	,	Volume	# Pieces:
Other exemption - Write in:		Ni-						Depth	Width	Height	(Cube)	
Is product repackaged?				nal product purchased		Item/Each:	0.07	1.61	1.61	2.98		1
Is product sold by manufacturer's Has FDA granted waiver/exceptio			rect from mfr?	cumentation from FDA.		Box/Carton/Bun						
Has I DA granted walver/exCeptio			yes, anden 00			Inner Pack:						
	GTI	N AND HIBCC PRODUCT INFORMATION				Case:						
							2.79	9.72	6.5	3.81		24
Saleable Unit of Measure	Quantity	HIBCC	GTIN-		Unit of Use GTIN-14	Pallet:	570.02	47	39	49		4,752
X Item/Each	1		00372	578136062			510.02	47		-10		4,102
Box/Carton/Bundle/Inner Pack												~
x Case	24			578136060			COST INFORMATION			WHOLESALI	ER USE ONL	.r:
x Pallet	4,752		50372	578136067		Bogular Cost			Vendor #:			
	┥ ┝━━━┥		-			Regular Cost Invoice Cost (W)	AC) (\$)		Whsl. Code	# •		
	┥ ┝━━━┥						, (Ψ)		Fineline Co			
	1 1					As of date:						
									-			
		Attach copy of SAFETY DATA SHEET (SI	DS) or non haza	ard letter, PACKAGE INSER	RT, LABEL AND PHOTO OF I	PRODUCT PACKAG	ING and BARCODE.					
i i i i i i i i i i i i i i i i i i i												
*Please provide any additional inf	formation on page 2.			See new p. 3 for Design	ated Drop Ship Only.	Si	gnature:					

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

Version 2020 For Designated Drop Ship Only Products, Please Use Page 3							
MATERIAL HAZ	ZARD 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? Does the product label bear a CA Prop 65 warning? C. Contact Hazard? No	SDS Hazard Classification Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard Aerosol Class; Identify NFPA Storage Level:						
d. Does this product require special clean-up instructions? No (If yes, attach SDS with special instructions.) No e. Does the product contain DEHP? No Is this product regulated for shipment by DOT? No (if yes, answer a-e below and provide SDS) No	Is the product a NIOSH hazardous drug?						
a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard? No	Hazardous Waste Identification EPA Hazardous Waste Code: Waste Characteristics						
Is this product regulated for shipment by IATA? No	REMS or REGISTRY RESTRICTIONS						
(if yes, answer a-e below and provide SDS) a. UN/dentification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group	Is there a REMS on this product? No If Yes, is it managed with a pharmacy registry? Website URL:						
e. Inhalation Hazard?	Med Guide Required No						
Is the product restricted for air shipment? If so, indicate restriction:	Limited Distribution Requirement No						
Passenger Cargo Passenger & Cargo	Comments / Details: (For example, iPledge program?)						
Is this a reportable quantity? No RQ Threshold:	REMS: Phone: REMS Program Manager Name: Phone: Supplier Manages REMS registry exclusively: Phone: Wholesale distributor support: Provider Name: Provider Name: DEA #: Site Enrollment Number assigned PCPDP#: by Supplier: NPI #:						
SP#	Registry:						
	Registry Program Contact Name: Phone:						
ADD'L STORAGE INFORMATION	Comments						
Is the Product							
Controlled Substance? No Controlled Substance Code	RETURN INSTRUCTIONS						
Controlled by State(s)? No Listed Chemical (List I or II) ARCOS Reportable? No If yes, indicate which:	Contact tel. # if product received damaged: 888-304-5022						
Schedule No. Is it a scheduled listed chemical product?:							
CLASS OF TRADE RESTRICTION:	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	www.vionausa.com						
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:							
MISCELLANE	DUS NOTES and/or Image of Product Barcode:						



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Version 2020 FOR DESIGNATED DROP SHIP PRODUCT ONLY	Y - 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 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:
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) 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: 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?