

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

Version 2020				Introduction Type:	Post Launch Change]	x Final Version			Date:	9/6/2	2022		
		PRODUCT INFORMATION					SPECIAL HAN	DLING AND STO	RAGE REQUI	REMENTS*				
Company Name:	Viona Pharmaceuticals Inc. Application: ANDA DA/BLA (drug): PMA/510(k)(med device): 208970 208970					a. Temperature – Indicate the USP temperature range for this product. Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)								
Application Number for NDA/AN		2 2 ned device):]	Temperature Range	Controlled Room	– between 20	and 25 C (68	° – 77° F)						
DUNS:	081468959						Other Temperature Range F	Requirement						
Proprietary Name (If Applicable) a Selling Unit NDC:	72578-057-16	Felbamate Tablets, USP 600 mg Unit of Use NDC:		UPC: 372578	057169		(write in) Notes							
UDI	N/A	CVX Code:		MVX Code: N/A		11	1000							
Description:	Light pink to pink colored, capa	sule shaped biconvex tablets, debossed with "10" a	and "54"separate	ed by breakline on one side	and plain on other.	<u>i</u> l	Is this product to be shipped				No			
Active Ingredient(s):		Is this product to be shipped		dry ice?		No								
URL for Additional Product Inform	b. Contact for temperature excursion questions: Name: Customer Service													
Address:	20 Commerce Drive			Address 2: Suite 340			Number:			888-304-5022				
City:	Cranford Chris Urbanski	Cranford State: NJ Zip: 07016 Chris Urbanski Email: Curbanski@vionausa.com						Group E-mail:				customerservice@vionausa.com		
Key Contact: Phone Number:	908-956-0600		Fax:	Curbanski@vionausa 908-514-4005	1.com	c. Special reg	ulations for product in any	states?			No			
Product Therapeutic Classificatio		ic/Anticonvulsant]	Special returns requirement				No			
	ADDITIONAL PROD			PRODUCT DESCR	IPTION INFORMATION	d. Store produ	uct (unit of sale) upright?				No			
The product is? a legend device?	Ne	Is the Product Direct-Ship	Only			e. Shelf life:	Protect product (unit of sa	le) from light?			No 24	Mantha		
if yes, enter class #	No	Is the Product Neither Orphan Drug Status		Size:	90 ct	e. Sneif life:	Initial shelf life at launch (f different):			24	Months Months		
a product kit? if yes, list NDCs of	No	FDA Approval Status		Strength:	600 mg			ORDER INFORI						
component parts				Dosage Form:	Tablets									
reverse numbered? co-licensed?	No No	Allergens Present					Unit of Sale x Bottle		What is the 1 Bottle of 90		unit?			
latex-free?	Yes	Allergena Freaent		Des des t Oliverse	Ormanda		Box/Carton			g. 1 Box of 10) Vials)			
preservative-free?	Yes			Product Shape:	Capsule		Ampule			-				
correctional institution block?	No			Product Color:	Light pink to pink		Glass		Minimum o	der quantity	?	Yes		
opioid? Cannabinoid?	No No	Country of Origin India					Tube Vial Liquid Sgl							
If Unit Dose, is item bar coded to u				Product Imprint:	"10" ; "54"		Vial Liquid Multi		If Yes, how	many of whi	ch package	type?		
scanning?		Is this product covered under the					Vial Powder Sql		15	Each				
If Unit Dose, indicate NDC here:		Trade Agreements Act (TAA)?	No				Vial Power Multi Other: Write In		1	Inner/Carton Case	/Pack			
		FOR GENERIC DRUG PRODUCTS				<u>1</u>	Other. White in			Case				
			Auth		orized Generic, other section			ARMACY ORDER						
I. Orange Book Rating:					re not applicable	Rec. sell unit		T	Rx billing u		acy:			
II. Generic Equivalent to What Bra	Felbatol					(Write-in, e.g.	1 Bottle	l	x	Each Gram				
	DRUG	SUPPLY CHAIN SECURITY ACT (DSCSA) INFO	RMATION			(Trino III, o.g.	, they			Milliliter				
Does supplier meet DSCSA defini	ition of manufacturer?	Yes	LN:	0372578000004			ITEN	AND PACKING I	NEORMATIO	J				
Is product exempt from DSCSA?		No	2.11.	0012010000004										
If yes, select exemption:				•			Weight Lbs.	Dimens	ions (US msn	nts.)	Volume	# Pieces:		
Other exemption - Write in:		No					Weight Ebs.	Depth	Width	Height	(Cube)	#110003.		
Is product repackaged? Is product sold by manufacturer's	s exclusive distributor?		Yes, was origit irect from mfr?	nal product purchased		Item/Each:	0.26	2.12	2.12	3.97		1		
Has FDA granted waiver/exceptio				cumentation from FDA.		Box/Carton/B	undle/							
						Inner Pack:								
		GTIN AND HIBCC PRODUCT INFORMATION				Case:	7.49	10.71	6.46	5.11		15		
Saleable Unit of Measure	Quantity	HIBCC	GTIN-		Unit of Use GTIN-14	Pallet:	1062.99	47.24409	39.3701	49.21		2,100		
X Item/Each Box/Carton/Bundle/Inner Pack	1		00372	578057169		└────								
X Case	15	1	40372	578057167			COST INFORMATION			WHOLESAL	ER USE ONL	.Y:		
X Pallet	2,100		50372	578057164										
	-	┥ ┝────	-			Regular Cost Invoice Cost (Vendor #: Whsl. Code	#•				
						invoice Cost ((III.C.) (I)		Fineline Co					
						As of date:		•						
μ														
*Please provide any additional inf	formation on name 2	Attach copy of SAFETY DATA SHEET (S	DS) or non haza	ard letter, PACKAGE INSE See new p. 3 for Design		PRODUCT PACK	AGING and BARCODE. Signature:							
I IGAGE PROVIDE any adultional in	ionnation on page 2.			occ new p. 5 for Design	atea brop omp omy.		orgnature.							

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	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? Does the product label bear a CA Prop 65 warning? No	SDS Hazard Classification Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard						
c. Contact Hazard? No d. Does this product require special clean-up instructions? No (If yes, attach SDS with special instructions.) e. Does the product contain DEHP? No Is this product regulated for shipment by DOT? No	Aerosol Class; Identify NFPA Storage Level: Is the product a NIOSH hazardous drug? If yes, indicate which:						
(if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group	Hazardous Waste Identification 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) a. UN/dentification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group	REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? No If Yes, is it managed with a pharmacy registry? Website URL:						
e. Inhalation Hazard? Is the product restricted for air shipment? If so, indicate restriction: Passenger Cargo	Med Guide Required No Limited Distribution Requirement No Comments / Details: (For example, iPledge program?) Image: Comment of the second sec						
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 Permit; DOT-SP Special Perovision (listed in Column 7 of 49 CFR 172.101);	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:						
ADD'L STORAGE INFORMATION	Registry Program Contact Name: Phone: Comments						
Is the Product Controlled Substance? No Controlled Substance Code	RETURN INSTRUCTIONS						
Controlled Substance ? No Controlled Substance Code Controlled by State(s)? No Listed Chemical (List I or II) ARCOS Reportable? No If yes, indicate which: Schedule No. Is it a scheduled listed chemical product?:	Contact tel. # if product received damaged: 888-304-5022 Is product returnable for credit: Yes URL/Link to returns policy: Yes						
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices	www.vionausa.com						
Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	Special regulations or returns requirements for this product in certain states?						
MISCELLANEC	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?