

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

Version 2021					Introduction	Туре:	New Item	x	Final Version			Date:	8/10	2021
		PRODUCT IN	ORMATION						SPECIAL HAN	IDLING AND STOR	AGE REQUI	REMENTS*		
Company Name:					Application: ANDA		a. Temperature – Indicate the USP temperature range for this product.							
Application Number for NDA/AND	DA/BLA (drug); PMA/510(k)(n	ned device):	21	1977					perature Range	Controlled Room		and 25 C (68	° – 77° F)	
Medical Device Class, if applicable:														
	82-667-4775							Othe	r Temperature Range	Requirement				
Proprietary Name (If Applicable) and		Esomeprazole Magnesium		sules 20MG 900					(write in)					
	31722-664-90	Unit of Us			UPC: MVX Code:	33172266	64905	Note	s					
UDI		CVX Co	e:		MVX Code:									
Description:	Oral Solid, Capsule, White, H	/E2							s product to be shippe				No	
Active Ingredient(s): Esomeprazole Magnesium														
Active ingrementations context for temperature excursion questions:														
URL for Additional Product Information	ation:							Nam		631013.				
Address:	800 Centennial Ave.				Address 2:			Num	ber:					
	Piscataway			State:	NJ	Zip: 0)8854	Grou	ıp E-mail:					
	Customer Service			Email:	customerservice@camberpharma.com									
	1-866-827-3647			Fax:	732-562-8788				ns for product in any				No	
Product Therapeutic Classification):							Spec	ial returns requiremen	ts for this product?			No	
	ADDITIONAL PRO	DUCT INFORMATION			PRODUCT	DESCRIPT	ION INFORMATION	d. Store product (u	nit of sale) unright?				No	
The medication			Direct-Ship 0	Only	11000001	DEGORIFI				ala) from Kato				
The product is? a legend device?	No	Is the Product Is the Product		July		90	let.	e. Shelf life:	ect product (unit of s	are) from light?			No 24	Months
a legend device? if yes, enter class #	INU	Orphan Drug S			Size:	90			I shelf life at launch (if different).			24	Months
a product kit?	No	orphan brug of				20	Img	inde		in americity.			24	Months
if yes, list NDCs of		FDA Approval S	tatus		Strength:		-			ORDER INFORM	IATION			
component parts					Dosage For	m: Or	al Solid - Capsule							
reverse numbered?	No				J. J. J.				of Sale		What is the		unit?	
co-licensed?	No Yes	Allergens Prese	nt			0	apsule	X			1 bottle of 90) capsules g. 1 Box of 1	() Viele)	
latex-free? preservative-free?	Yes	-			Product Sha	ape:	apsule		Box/Carton Ampule		(write-in, e.	g. I bux ui i	U VIAIS)	
correctional institution block?	No					. w	hite		Glass		Minimum or	der quantity	?	Yes
opioid?	No				Product Col	lor:			Tube					
Cannabinoid?	No	Country of Origin	India		Product Imp	orint:	E2		Vial Liquid Sgl					
If Unit Dose, is item bar coded to un									Vial Liquid Multi				ch package	type?
hospital scanning? If Unit Dose, indicate NDC here:	No	Is this product of Trade Agreemer		No					Vial Powder Sql Vial Power Multi			Each Inner/Cartor	/De els	
Il Unit Dose, Indicate NDC here:		Trade Agreemen	IS ACI (TAA)?	NO					Other: Write In			Case	Pack	
		FOR GENERIC DI	UG PRODUCTS								1	louoo		
				Au	uthorized Generic	*If Author	ized Generic, other	-	Pł	IARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AB					section fie	elds are not applicable	Rec. sell unit to customer? Rx billing unit to pharmacy:						
II. Generic Equivalent to What Brand?: Nexium Delayed-Release Capsules								Each						
							(Write-in, e.g. 1 Vial)			Gram			
	DRU	G SUPPLY CHAIN SECURIT	Y ACT (DSCSA) INFO	RMATION								Milliliter		
Does supplier meet DSCSA definit	ion of manufacturor?	Yes		GLN:	031722000000				ITEN	AND PACKING I		N		
Is product exempt from DSCSA?		No		OLN.	001722000000									
If yes, select exemption:				GCP:						Dimensi	ons (US msm	nts.)	Volume	Saleable #
Other exemption - Write in:				001.					Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?		No		If yes, was o	riginal product pu	rchased		Item/Each:	0.1		2	3.5	0	1
Is product sold by manufacturer's		No		direct from n							2	3.0	0	1
Has FDA granted waiver/exception				Provide sour	ce manufacturer f	or repacka	ged product	Box/Carton/Bundle	/				0	
If yes, attach documentation from	n FDA.							Inner Pack: Case:						
		GTIN AND HIBCC PRO	DUCT INFORMATION					Case:	3.5	11.5	5	8.25	0.275	24
								Pallet:					0	
Saleable Unit of Measure	Saleable Qua	antity HIBCC		GT	IN-14	ι	Unit of Use GTIN-14						0	
X Item/Each	1			003	31722664905									
Box/Carton/Bundle/Inner Pack						_		c	OST INFORMATION			WHOLESAL	ER USE ONL	Y:
X Case Pallet	24			303	31722664906	-		Regular Cost			Vendor #:			
F allos		-				-		Invoice Cost (WAC)	(\$)	\$30.00	Whsl. Code	#:		
						-				φ30.00	Fineline Co			
				1		-		As of date:						
											1			
μ								L <u>I</u>						
		Attach copy of SAF	ETY DATA SHEET (SI	DS) or non haza			LABEL AND PHOTO OF P							
*Please provide any additional information on page 2. See new p. 3 for Designated Drop Ship Only. Signature:														

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	For Designated Drop Ship Only Products, Please Use Page 3					
MAT	TERIAL HAZARD 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? c. Contact Hazard? d. Does this product require special clean-up instructions?	X Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard Does the product have an Aerosol class? If yes, identify NFPA Storage Level: Image: Contact Hazard					
(If yes, attach SDS with special instructions.) e. Does the product contain DEHP? Is this product regulated for shipment by DOT?	NFPA Storage Level: No Is the product a NIOSH hazardous drug?					
(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?	If yes, indicate which: Hazardous Waste Identification EPA Hazardous Waste Code: Waste Characteristics					
Is this product regulated for shipment by IATA? (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 REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? No If Yes, is it managed with a pharmacy registry? No Website URL: No					
Is the product restricted for air shipment? If so, indicate restriction: Passenger Cargo Passenger & Cargo Is this a reportable quantity?	No Med Guide Required Limited Distribution Requirement Comments / Details: (For example, iPledge program?) REMS:					
RQ Threshold: Is this a marine pollutant? No Is this product shipped utilizing an authorized DOT exception or Special Permit? (if yes, identify method below) Limited Quantity Consumer Commodity, ORM-D Small Quantity (49 CFR 173.4) Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101);	REMS Program Manager Name: Phone: Supplier Manages REMS registry exclusively: Phone: Wholesale distributor support: Provider Name: Provider Name: DEA #: Site Enrollment Number assigned NCPDP#: by Supplier: NPI #:					
SP#ADD'L STORAGE INFORMATION	Registry: No Registry Program Contact Name: Phone: Comments					
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?:	No RETURN INSTRUCTIONS No Contact tel. # if product received damaged: 1-866-827-3647 No Is product returnable for credit: URL/Link to returns policy:					
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:	Yes No No Special regulations or returns requirements for this product in certain states? No If so, which states? Other requirements? Comments?					
	ISCELLANDOLIS NOTES and/or lingge of Broduct Dependen					
MIS	SCELLANEOUS 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: Drop Ship service fee billed with each order: Drop Ship miscellaneous fees billed: Comments:	Overnight receipt available: Image: Comparison of the co
Class of Trade Restriction:	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:	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?