

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

Version 2021					Introduction	Туре:	New Item		x Final Version			Date:	10/5	/2021
		PRODUCT IN	ORMATION						SPECIAL HAN	IDLING AND STOP	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					Temperature Range	Controlled Room		and 25 C (68	° – 77° F)	
Medical Device Class, if applicable:														
	82-667-4775								Other Temperature Range	Requirement				
Proprietary Name (If Applicable) and		Esomeprazole Magnesium		sules 40MG 900					(write in)					
	31722-665-90	Unit of Us			UPC: MVX Code:	3317226	65902	1	Notes					
UDI		CVX Co	e:		MVX Code:									
Description:	Oral Solid, Capsule, White, H	/E3							s this product to be shippe				No	
Active Ingredient(s): Esomeprazole Magnesium														
Active Ingredient(s): Esomeprazole Magnesium b. Contact for temperature excursion questions:														
URL for Additional Product Inform	ation.								Name:	estions.				
Address:	800 Centennial Ave.				Address 2:				Number:					
City:	Piscataway State:			NJ	NJ Zip: 08854			Group E-mail:						
	Customer Service			Email:	customerserv	customerservice@camberpharma.com								
	1-866-827-3647	-866-827-3647 Fax:			732-562-8788			c. Special regulations for product in any states? No						
Product Therapeutic Classification	1:							5	Special returns requiremen	ts for this product?			No	
					DRODUEZ	DECODUC								1
	ADDITIONAL PRO	DUCT INFORMATION	_		PRODUCT	DESCRIPT	TION INFORMATION	-	ct (unit of sale) upright?				No	1
The product is?		Is the Product		Only		_			Protect product (unit of sa	ale) from light?			No	
a legend device?	No	Is the Product.			Size:	90	Oct	e. Shelf life:	nisial abalf life of low 1	if different ()			24	Months
if yes, enter class # a product kit?	No	Orphan Drug S	atus			10	0mg		nitial shelf life at launch (if different):			24	Months
if yes, list NDCs of	INU	FDA Approval S	tatus		Strength:	40	onig			ORDER INFORM	ATION			
component parts					Decese Fee		oral Solid - Capsule							
reverse numbered?	No				Dosage For	m:		L	Unit of Sale		What is the	NDC selling	unit?	
co-licensed?	No	Allergens Prese	nt						X Bottle		1 bottle of 90			
latex-free?	Yes				Product Sha	ape:	apsule	_	Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free?	Yes						<i>II</i> . 1	_	Ampule					N/s s
correctional institution block? opioid?	No	_			Product Col	lor:	/hite	_	Glass Tube		Minimum or	rder quantity	<i>c</i>	Yes
Cannabinoid?	No	Country of Origin	India			. н	I/E3	_	Vial Liquid Sgl					
If Unit Dose, is item bar coded to un					Product Imp	print:			Vial Liquid Multi		If Yes, how	many of whi	ch package	type?
hospital scanning?		Is this product of						Vial Powder Sql 24 Each						
If Unit Dose, indicate NDC here:		Trade Agreemer	ts Act (TAA)?						Vial Power Multi			Inner/Carton/Pack		
							L	Other: Write In			Case			
	FOR GENERIC DRUG PRODUCTS													
Authorized Generic *If Authorized Generic, other								PHARMACY ORDER / BILL UNIT						
L Orenne Beek Beting	AB				anonzed Genene		ields are not applicable	Rec. sell unit to						
I. Orange Book Rating: AB Section rields are not applicable II. Generic Equivalent to What Brand?: Nexium Delayed-Release Capsules								Rec. sell unit to customer? Rx billing unit to pharmacy:						
in Selecte Lyandick to that Datati.							(Write-in, e.g. 1	Vial)			Gram			
								Milliliter						
Does supplier meet DSCSA definit	ion of manufacturer?	Yes No		GLN:	031722000000				ITEN	I AND PACKING I	NFORMATION	N		
Is product exempt from DSCSA?		INU								_				
If yes, select exemption:				GCP:					Weight Lbs.		ons (US msm	,	Volume (Cubo)	Saleable #
Other exemption - Write in: Is product repackaged?		No		If yes was o	riginal product pur	rchased		Item/Each:		Depth	Width	Height	(Cube)	Pieces
Is product sold by manufacturer's	exclusive distributor?	No		direct from n		Shuseu		nony Laon.	0.2		2	3.34		1
Has FDA granted waiver/exception		No			ce manufacturer f	or repacka	aged product	Box/Carton/Bu	ndle/					
If yes, attach documentation from	n FDA.							Inner Pack:						
		GTIN AND HIBCC PRO						Case:	4	12	8	5	0.28	24
		GTIN AND HIBCC PRO	DUCT INFORMATION					Pallet:						
Saleable Unit of Measure	Saleable Qua	antity HIBCC		GT	N-14		Unit of Use GTIN-14	rallet.						
X Item/Each	1				31722665902		00331722665902	L		1				
Box/Carton/Bundle/Inner Pack	00001122000302						00001722000002		COST INFORMATION		1	WHOLESAL	ER USE ONL	.Y:
X Case	24			303	31722665903									
Pallet						_		Regular Cost			Vendor #:			
		_				-		Invoice Cost (W	VAC) (\$)	\$30.00	Whsl. Code			
						-		As of date:			Fineline Co	ae:		
		-				-		As or date:			1			
[•		Attach copy of SAI	ETY DATA SHEET (SI	OS) or non haza	ard letter, PACKAGE	E INSERT,	LABEL AND PHOTO OF P	RODUCT PACKAG	ING and BARCODE.		•			
*Please provide any additional info	ormation on page 2.		- (-	,			ed Drop Ship Only.		Signature:					
-	-					-								

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

Version 2021 For Design	ated Drop Ship Only Products, Please Use Page 3
MATERIAL H/	AZARD 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? No 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? (If yes, attach SDS with special instructions.) e. Does the product contain DEHP? No Is this product regulated for shipment by DOT? No	SDS Hazard Classification x Organic Inorganic Oxidizer Steroid/Androgen Contact Hazard Does the product have an Aerosol class? If yes, identify NFPA Storage Level: No NFPA Storage Level: Image: Contact Hazard Is the product a NIOSH hazardous drug? 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? Is this product regulated for shipment by IATA2	If yes, indicate which: 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? 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 product shipped utilizing an authorized DOT exception or Special Permit? No (if yes, identify method below)	REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? No If Yes, is it managed with a pharmacy registry? No Website URL: Image: Colspan="2">Image: Colspan="2" REMS: No Image: Colspan="2" Image: Colspan="2" <t< td=""></t<>
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); SP#	Site Enrollment Number assigned by Supplier: NCPDP#: NPI #: Comments Registry: Registry Program Contact Name: Comments
	Comments
Is the Product Controlled Substance? No Controlled Substance Code Listed Chemical (List I or II) No ARCOS Reportable? No If yes, indicate which: 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 Yes	RETURN INSTRUCTIONS Contact tel. # if product received damaged: 1-866-827-3647 Is product returnable for credit: URL/Link to returns policy:
Restricted to retail pharmacy only: No Restricted to hospital, clinics, and physician offices only: No Restricted from US territories? (explain in comments) No Comments:	Special regulations or returns requirements for this product in certain states? If so, which states? Other requirements? Comments?
MISCELLAN	EOUS 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?