

Version 2024

Introduction Type:

Final Version

Date: 9/8/2025

PRODUCT INFORMATION			
Company Name:		MILLA PHARMACEUTICALS INC	Application:
Application Number for NDA/ANDA/BLA; PMA/510(k):		A211081	ANDA
Medical Device Class, if applicable:			
DUNS:	119319487		
Proprietary Name (If Applicable) and Established Name:			
Selling Unit NDC:	81284-0212-10	Unit of Use NDC:	81284-0212-00
UDI		UPC:	381284212106
		CVX Code:	
		MVX Code:	
Description:	Phenylephrine 10mg/mL 5 mL x 10 vials		
Active Ingredient(s):	Phenylephrine		
URL for Additional Product Information:			
Address:	1310 Highway 96E		Address 2:
City:	White Bear Lake		Suite 104A
Key Contact:	Siddharth Nirmal		State:
Phone Number:	+1 302 946 0117		MN
			Email:
			siddharth.nirmal@aforallpharma.com
			Fax:
Product Therapeutic Classification:			

ADDITIONAL PRODUCT INFORMATION		PRODUCT DESCRIPTION INFORMATION	
The product is?	No	Is the Product... Is the Product... Orphan Drug Status	Direct-Ship Only Unit of Use
a legend device?	No		
if yes, enter class #	No		
a product kit?	No		
if yes, list NDCs of component parts reverse numbered?	No	FDA Approval Status	
co-licensed?	No		
latex-free?	Yes	Allergens Present	
preservative-free?	Yes		
correctional institution block?	No		
opioid?	No	Country of Origin	India
Cannabinoid?	No		
If Unit Dose, is item bar coded to unit dose for hospital scanning?	Yes	Is this product covered under the Trade Agreements Act (TAA)?	No
If Unit Dose, indicate NDC here:			

FOR GENERIC DRUG PRODUCTS	
I. Orange Book Rating:	AP1
II. Generic Equivalent to What Brand?:	Biorphen
	Authorized Generic
	*If Authorized Generic, other section fields are not applicable

DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION			
Does supplier meet DSCSA definition of manufacturer?	Yes	GLN:	0381284000000
Is product exempt from DSCSA?	No	GCP:	
If yes, select exemption: Other exemption - Write in:			
Is product repackaged?	No	If yes, was original product purchased direct from mfr?	
Is product sold by manufacturer's exclusive distributor?	Yes	Provide source manufacturer for repackaged product	
Has FDA granted waiver/exception/exemption for product? If yes, attach documentation from FDA.	No		

GTIN AND HIBCC PRODUCT INFORMATION					
Saleable Unit of Measure	RFID tag(Y/N)	Saleable Quantity	HIBCC	GTIN-14	Unit of Use GTIN-14
Item/Each					381284212007
x Box/Carton/Bundle/Inner Pack		1		00381284212106	
x Case		72		20381284212100	
x Pallet		2592			

a. Temperature – Indicate the USP temperature range for this product.			
Temperature Range	Controlled Room – between 20 and 25 C (68° – 77° F)		
Other Temperature Range Requirement (write in)			
Notes			
Is this product to be shipped to customers on ice?	No		
Is this product to be shipped to customers on dry ice?	No		
b. Contact for temperature excursion questions:			
Name:	DL-QA GPI Pharma		
Number:			
Group E-mail:	ga@gpi-pharma.lc		
c. Special regulations for product in any states?	No		
Special returns requirements for this product?	No		
d. Store product (unit of sale) upright?	No		
Protect product (unit of sale) from light?	Yes		
e. Shelf life:	24 Months		
Initial shelf life at launch (if different):	Months		

ORDER INFORMATION	
Unit of Sale	What is the NDC selling unit?
x Bottle	1 box of 10 vials
Box/Carton	(Write-in, e.g. 1 Box of 10 Vials)
Ampule	
Glass	Minimum order quantity?
Tube	Yes
Vial Liquid Sgl	
Vial Liquid Multi	If Yes, how many of which package type?
Vial Powder Sgl	12 Each
Vial Powder Multi	Inner/Carton/Pack
Other: Write In	Case

PHARMACY ORDER / BILL UNIT	
Rec. sell unit to customer?	Rx billing unit to pharmacy:
1 box of 10 vials	Each
(Write-in, e.g. 1 Vial)	Gram
HPCS J-Code:	x Milliliter

ITEM AND PACKING INFORMATION						
	Weight Lbs.	Dimensions (US msmts.)			Volume	Saleable #
		Depth	Width	Height	(Cube)	Pieces
Item/Each:	0.032739	0.874	0.874	1.76	1.3444218	0.1
Box/Carton/Bundle/ Inner Pack:	0.36826	4.528	1.89	2.008	17.184303	1
Case:	28.66006	14.921	12.795	9.646	1841.5583	72
Pallet:	1093.49152	48	40	43.701	83905.92	2592

COST INFORMATION		WHOLEALER USE ONLY:	
Regular Cost		Vendor #:	
Invoice Cost (WAC) (\$)	\$90.00	Whsl. Code #:	
As of date:	6/24/2022	Fineline Code:	

Attach copy of SAFETY DATA SHEET (SDS) or non hazard letter, PACKAGE INSERT, LABEL AND PHOTO OF PRODUCT PACKAGING and BARCODE.

*Please provide any additional information on page 2.

See new p. 3 for Designated Drop Ship Only.

Signature:



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

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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? ☐ No
Is the product a CA Prop 65 reproductive toxicant? ☐ No
Does the product label bear a CA Prop 65 warning? ☐ No

- c. Contact Hazard? ☐ Yes
- d. Does this product require special clean-up instructions?
(If yes, attach SDS with special instructions.) ☐ Yes
- e. Does the product contain DEHP? ☐ No

Is this product regulated for shipment by DOT?
(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? ☐ Yes

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? ☐ Yes

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 Permit; DOT-SP
- ☐ Special Provision (listed in Column 7 of 49 CFR 172.101);
SP#

ADD'L STORAGE INFORMATION

Is the Product...

- Controlled Substance? ☐ No Controlled Substance Code
- Controlled by State(s)? ☐ No Listed Chemical (List I or II) ☐ No
- ARCOS Reportable? ☐ No 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

☐ Yes

Restricted to retail pharmacy only:

☐ No

Restricted to hospital, clinics, and physician offices only:

☐ No

Restricted from US territories? (explain in comments)

☐ No

Comments:

SDS Hazard Classification

- ☐ Organic ☐ Corrosive
- ☐ Inorganic ☐ Oxidizer
- ☐ Steroid/Androgen ☒ Contact Hazard

Does the product have an Aerosol class? If yes,
identify NFPA Storage Level:

NFPA Storage Level:

Is the product a NIOSH hazardous drug?
If yes, indicate which:

☐ No

Hazardous Waste Identification

EPA Hazardous Waste Code:

Waste Characteristics

REMS or REGISTRY RESTRICTIONS

Is there a REMS on this product?

☐ No

If Yes, is it managed with a pharmacy registry?

Website URL:

Med Guide Required ☐

Limited Distribution Requirement ☐

Comments / Details: (For example, iPledge program?)

REMS:

REMS Program Manager Name:

Phone:

Supplier Manages REMS registry exclusively: ☐

Wholesale distributor support: ☐

Provider Name:

DEA #:

Site Enrollment Number assigned
by Supplier:

NCPDP#:

NPI #:

Comments

Registry:

Registry Program Contact Name:

Phone:

Comments

RETURN INSTRUCTIONS

Contact tel. # if product received damaged:

773-541-3317

Is product returnable for credit: ☐ Yes

URL/Link to returns policy:

www.millapharmaceuticals.com

Special regulations or returns requirements for this
product in certain states? ☐ No

If so, which states? Other requirements? Comments:

MISCELLANEOUS NOTES and/or Image of Product Barcode:

Release DATE



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

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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: Fax Number: Fax Number: Phone No.: Site Address:	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: Expedited freight fees billed with each order: Drop Ship service fee billed with each order: Drop Ship miscellaneous fees billed: Comments:	Overnight and Priority Overnight PO Processing Overnight receipt available: ☐ PO Receipt cut off time: Days of week overnight is available: ☐ Monday ☐ Tuesday ☐ Wednesday ☐ Thursday ☐ Friday Priority Overnight receipt available: ☐ PO Receipt Cut off time: Saturday Overnight receipt available: ☐ PO Receipt Cut off time: Order receipt method: Phone: Phone #: Fax: Fax #: EDI: Overnight Fees apply: ☐ Other fees apply: ☐
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:	
Other Data Information Required to Process PO: Patient Procedure Date: Physician Name: Physician/Clinic Phone #: Physician State License #: Physician/Clinic DEA #: Physician/Clinic Specialty:	Return Instructions 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? ☐