



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

Version 2024		Introduction Type: New Item		☒ Final Version		Date: 3/3/2025					
PRODUCT INFORMATION						SPECIAL HANDLING AND STORAGE REQUIREMENTS*					
Company Name: Novadoz Pharmaceuticals LLC				Application: ANDA		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? Is this product to be shipped to customers on dry ice? b. Contact for temperature excursion questions: Name: Number: Group E-mail: c. Special regulations for product in any states? Special returns requirements for this product? d. Store product (unit of sale) upright? Protect product (unit of sale) from light? e. Shelf life: Initial shelf life at launch (if different): <table border="1"><tr><td>24</td><td>Months</td></tr><tr><td>24</td><td>Months</td></tr></table>		24	Months	24	Months
24	Months										
24	Months										
Application Number for NDA/ANDA/BLA; PMA/510(k): 217397				NDA 505(b) Type:							
Medical Device Class, if applicable:											
DUNS: 081109687											
Proprietary Name (If Applicable) and Established Name: Nebivolol Tablets 20 mg, 90											
Selling Unit NDC: 72205-153-90		Unit of Use NDC:		UPC: 372205153905							
UDI		CVX Code:		MVX Code:							
Description: <input "n9"="" and="" m"="" on="" one="" other="" side="" side"="" type="text" value="Blue colored, triangular shaped, unscored tablets debossed with "/>											
Active Ingredient(s): Nebivolol Hydrochloride											
URL for Additional Product Information:											
Address: 20 Duke Road		Address 2: Suite A									
City: Piscataway		State: NJ		Zip: 08854							
Key Contact:		Email: sales@novadozpharma.com									
Phone Number: 908-360-1500		Fax: 732-902-2113									
Product Therapeutic Classification:											
ADDITIONAL PRODUCT INFORMATION			PRODUCT DESCRIPTION INFORMATION								
The product is? a legend device? if yes, enter class # a product kit? if yes, list NDCs of component parts reverse numbered? co-licensed? latex-free? preservative-free? correctional institution block? opioid? Cannabinoid? If Unit Dose, is item bar coded to unit dose for hospital scanning? If Unit Dose, indicate NDC here:			Anti-hypertensive Is the Product... Is the Product... Orphan Drug Status FDA Approval Status Allergens Present Country of Origin India Is this product covered under the Trade Agreements Act (TAA)?								
			Size: 90 Strength: 20 mg Dosage Form: Tablet Product Shape: Triangular Shaped Product Color: Blue colored Product Imprint: <input "n9"="" and="" on="" one="" other="" side="" side"="" type="text" value="M"/>								
FOR GENERIC DRUG PRODUCTS											
I. Orange Book Rating: AB ☐ Authorized Generic *If Authorized Generic, other section fields are not applicable											
II. Generic Equivalent to What Brand?: Bystolic											
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION											
Does supplier meet DSCSA definition of manufacturer? Yes			GLN: 8904159670843								
Is product exempt from DSCSA? No			GCP: 89041596								
If yes, select exemption: Other exemption - Write in:			If yes, was original product purchased direct from mfr?								
Is product repackaged? No			Provide source manufacturer for repackaged product								
Is product sold by manufacturer's exclusive distributor? No											
Has FDA granted waiver/exception/exemption for product? No											
If yes, attach documentation from FDA.											
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			1 Bottle of 90 Tablets		00372205153905	00372205153905					
☒ Box/ Carton/ Bundle/ Inner Pack			NA		NA						
☒ Case			96 Bottles in Case		30372205153906						
☒ Pallet			4224 Bottles in Pallet		50372205153900						
ITEM AND PACKING INFORMATION											
	Weight Lbs.	Dimensions (US msmts.)			Volume (Cube)	Saleable # Pieces					
	Depth	Width	Height								
Item/Each:	0.102	1.53	NA	3.28	52.00	1					
Box/ Carton/ Bundle/ Inner Pack:	NA	NA	NA	NA	NA	NA					
Case:	10.84	14.17	10.43	8.85	1,307.96	96					
Pallet:	516.63	47.24	39.37	41.33	76,867.13	4,224					
COST INFORMATION						WHOLESALE USE ONLY:					
Regular Cost					Vendor #:						
Invoice Cost (WAC) (\$)		\$15.00			Whsl. Code #:						
As of date: 3/3/2025					Fineline Code:						
*Please provide any additional information on page 2.											
Attach copy of SAFETY DATA SHEET (SDS) or non hazard letter, PACKAGE INSERT, LABEL AND PHOTO OF PRODUCT PACKAGING and BARCODE.											
See new p. 3 for Designated Drop Ship Only.											
Signature:											



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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? ☐
- 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?
(If yes, attach SDS with special instructions.) ☐
- e. Does the product contain DEHP? ☐

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? ☐

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? ☐

Is the product restricted for air shipment? If so, indicate restriction:

- ☐ Passenger
☐ Cargo
☐ Passenger & Cargo

Is this a reportable quantity? ☐

RQ Threshold:

Is this a marine pollutant? ☐

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);
SP#

ADD'L STORAGE INFORMATION

Is the Product...

- Controlled Substance? ☐ Controlled Substance Code
- Controlled by State(s)? ☐ Listed Chemical (List I or II)
- ARCOS Reportable? ☐ If yes, indicate which:
- Schedule No. Is it a scheduled listed chemical product?: ☐

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:

MISCELLANEOUS NOTES and/or Image of Product Barcode:

Release DATE

SDS Hazard Classification

- ☐ Organic
☐ Inorganic
☐ Steroid/Androgen
- ☐ Corrosive
☐ Oxidizer
☐ 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:

Hazardous Waste Identification

EPA Hazardous Waste Code:

Waste Characteristics

REMS or REGISTRY RESTRICTIONS

Is there a REMS on this product? ☐

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:

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?



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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? ☐