



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

Version 2024 Introduction Type: New Item ☒ Final Version Date: 6/8/2026

PRODUCT INFORMATION				SPECIAL HANDLING AND STORAGE REQUIREMENTS*			
Company Name: Novadoz Pharmaceuticals, LLC Application: ANDA				a. Temperature – Indicate the USP temperature range for this product.			
Application Number for NDA/ANDA/BLA; PMA/510(k): NDA 505(b) Type:				Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)			
Medical Device Class, if applicable:				Other Temperature Range Requirement (write in) NA			
DUNS: 08-110-9687				Notes			
Proprietary Name (If Applicable) and Established Name: Rivaroxaban Tablets, USP 2.5 mg 60's				Is this product to be shipped to customers on ice? No			
Selling Unit NDC: 72205-416-04 Unit of Use NDC: UPC: 372205416048				Is this product to be shipped to customers on dry ice? No			
UDI CVX Code: MXV Code:				b. Contact for temperature excursion questions:			
Description: Yellow, round, biconvex, fil coated tablets, debossed with "R" on one side and "2.5" on other side.				Name: P. Krishna Reddy			
Active Ingredient(s): Rivaroxaban				Number: 040-30449311			
URL for Additional Product Information:				Group E-mail: krishna_p@msnlab.com			
Address: 20 Duke Road Address 2: Suite A				c. Special regulations for product in any states?			
City: Piscataway State: NJ Zip: 08854				Special returns requirements for this product?			
Key Contact: Email: Sales@novadozpharma.com				d. Store product (unit of sale) upright? No			
Phone Number: 908-360-1500 Fax: 678-608-3778				Protect product (unit of sale) from light? No			
Product Therapeutic Classification:				e. Shelf life:			
				Initial shelf life at launch (if different): 24 Months			
				24 Months			
ADDITIONAL PRODUCT INFORMATION				PRODUCT DESCRIPTION INFORMATION			
The product is?		Is the Product...		Size:			
a legend device? No		Is the Product... Neither		Strength:		60's	
if yes, enter class #		Orphan Drug Status		Dosage Form:		2.5 mg	
a product kit? No						Tablets	
if yes, list NDCs of component parts		FDA Approval Status		Product Shape:		Round	
reverse numbered? No		Allergens Present		Product Color:		Yellow	
co-licensed? No				Product Imprint:		"R" on one side and "2.5" on other side	
latex-free? Yes							
preservative-free? Yes							
correctional institution block? No		Country of Origin					
opioid? No							
Cannabinoid? No							
If Unit Dose, is item bar coded to unit dose for hospital scanning?		Is this product covered under the Trade Agreements Act (TAA)? No					
If Unit Dose, indicate NDC here: 72205-416-04							
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?: XARELTO							
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:				If yes, was original product purchased direct from mfr?			
Other exemption - Write in:				Provide source manufacturer for repackaged product			
Is product repackaged? No							
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	
☐ Yes	Item/Each		Bottle of 60 tablets		00372205416048	00372205416048	
☐ NA	Box/Carton/Bundle/Inner Pack		NA		NA		
☐ Yes	Case		48 Bottles in Case		30372205416049		
☐ Yes	Pallet		4416 Bottles in Pallet		50372205416043		
ITEM AND PACKING INFORMATION							
	Weight Lbs.	Dimensions (US msmts.)			Volume (Cube)	Saleable # Pieces	
		Depth	Width	Height			
Item/Each:	0.066	NA	1.531	3.280	52	1	
Box/Carton/Bundle/Inner Pack:	NA	NA	NA	NA	NA	NA	
Case:	4.85	10.43	7.28	7.68	583.15	48	
Pallet:	486.17	47.24	39.37	36.61	68089	4,416	
COST INFORMATION							
Regular Cost				WHOLESALE USE ONLY:			
Invoice Cost (WAC) (\$) \$25.00				Vendor #:			
As of date: 6/8/2026				Whsl. Code #:			
				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:							



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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?

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?

(if yes, answer a-e below and provide SDS)

a. UN/Identification Number

UN3077

b. Proper Shipping Name

Environmentally hazardous substance, solid, N.O.S

c. DOT Hazard Class

9

d. Packing Group

III

e. Inhalation Hazard?

No

Is this product regulated for shipment by IATA?

(if yes, answer a-e below and provide SDS)

a. UN/Identification Number

UN3077

b. Proper Shipping Name

Environmentally hazardous substance, solid, N.O.S

c. DOT Hazard Class

9

d. Packing Group

III

e. Inhalation Hazard?

Yes

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

No

☐ Passenger

☐ Cargo

☐ Passenger & Cargo

Is this a reportable quantity? ☐ Yes

RQ Threshold: 5 litres or less for

Is this a marine pollutant? ☐ Yes

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?:

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

☐

Inorganic

☐

Steroid/Androgen

☐

Corrosive

☐

Oxidizer

☐

Contact Hazard

Does the product have an Aerosol class? If yes, identify

No

NFPA Storage Level:

NFPA Storage Level:

Is the product a NIOSH hazardous drug?

No

If yes, indicate which:

Group 2 items (non-antineoplastic that meets a hazard criterion)

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:

Supplier Manages REMS registry exclusively:

Wholesale distributor support:

Provider Name:

Site Enrollment Number assigned

by Supplier:

No

Phone:

DEA #:

NCPDP#:

NPI #:

Comments

Registry:

Registry Program Contact Name:

Comments

No

Phone:

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?

MISCELLANEOUS NOTES and/or Image of Product Barcode:

Release DATE



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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:	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: 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:	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?