

Version 2024

Introduction Type: New Item

x Final Version

Date: 2/21/2026

PRODUCT INFORMATION			
Company Name:		Novadoz Pharmaceuticals, LLC	
Application Number for NDA/ANDA/BLA; PMA/510(k):		NDA 505(b) Type:	
		AND A	
Medical Device Class, if applicable:			
DUNS:	08-110-9687		
Proprietary Name (If Applicable)		Established Name:	
		Brivaracetam Tablets 75 mg 60	
Selling Unit NDC:	Unit of Use NDC:	UPC:	
UDI	CVX Code:	MVX Code:	
Description:	Purple,Oval film coated tablet debossed with "B13" on one side and "M" on other side.		
Active Ingredient(s):	Brivaracetam		
URL for Additional Product Information:			
Address:		Address 2:	
City:		Suite A	
Key Contact:		State:	
Phone Number:		Email:	
		Zip:	
		Fax:	
Product Therapeutic Classification:			

ADDITIONAL PRODUCT INFORMATION		PRODUCT DESCRIPTION INFORMATION	
The product is?	No	Is the Product... Direct-Skip Only	
a legend device?	No	Is the Product... Neither	
if yes, enter class #		Orphan Drug Status	
a product kit?	No	FDA Approval Status	
if yes, list NDCs of component parts reverse numbered?	No	Allergens Present	
co-licensed?	No	Country of Origin	INDIA
latex-free?	Yes	If Unit Dose, is item bar coded to unit dose for hospital scanning?	
preservative-free?	Yes	If Unit Dose, indicate NDC here:	72205-270-60
correctional institution block? opioid?	No	Is this product covered under the Trade Agreements Act (TAA)?	No
Cannabinoid?	No		

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?:	BRIVIACT		

DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION			
Does supplier meet DSCSA definition of manufacturer?	Yes	GLN:	
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/exemption/exception 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	Bottle of 60 tablets			00372205270602	00372205270602
Box/Carton/Bundle/Inner Pack	NA			NA	
Case	24 Bottles in Case			30372205270603	
Pallet	4200 Bottles in Pallet			50372205270607	

SPECIAL HANDLING AND STORAGE REQUIREMENTS*					
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): Months Months					

ORDER INFORMATION	
Unit of Sale	What is the NDC selling unit?
x Bottle	1 Bottle of 60 tablets
Box/Carton	(Write-in, e.g. 1 Box of 10 Vials)
Ampule	
Glass	
Tube	
Vial Liquid Sgl	
Vial Liquid Multi	
Vial Powder Sgl	
Vial Power Multi	
Other: Write In	
	Minimum order quantity?
	Yes
	If Yes, how many of which package type?
	24 Each
	Inner/Carton/Pack
	Case

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

ITEM AND PACKING INFORMATION						
	Weight Lbs.	Dimensions (US msmts.)			Volume (Cube)	Saleable # Pieces
		Depth	Width	Height		
Item/Each:	0.108	NA	1.531	3.280	54	1
Box/Carton/Bundle/ Inner Pack:	NA	NA	NA	NA	NA	NA
Case:	3.81	10.04	7.09	4.92	350.22	24
Pallet:	706.33	47.24	39.37	40.35	75044.496	4,200

COST INFORMATION		WHOLEALER USE ONLY:	
Regular Invoice Cost (WAC) (\$)	\$16.00	Vendor #:	
As of date:	2/21/2026	Whsl. Code #:	
		Fineline Code:	



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

Version 2024

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?
(If yes, attach SDS with special instructions.) ☐ No
- 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? ☐ No

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

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? ☐ Yes ☐ No Controlled Substance Code
- Controlled by State(s)? ☐ No ☐ Yes Listed Chemical (List I or II) ☐ No ☐ Yes
- ARCOS Reportable? ☐ No ☐ Yes If yes, indicate which:
- Schedule No. 5 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:

☐ No

NFPA Storage Level:

Is the product a NIOSH hazardous drug?

☐ No

If yes, indicate which:

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:

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: 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?