



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

Version 2024

Introduction Type:

Post Launch Change

x

Final Version

Date:

01.22.2025

PRODUCT INFORMATION

Company Name:

Novadoz Pharmaceuticals LLC

Application Number for NDA/ANDA/BLA; PMA/510(k):

210686

Application:

ANDA

NDA 505(b) Type:

Medical Device Class, if applicable:

DUNS:

081109687

Proprietary Name (If Applicable) and Established Name:

Abiraterone Tablets 250mg

Selling Unit NDC:

72205-030-92

Unit of Use NDC:

UPC:

372205030923

UDI

CVX Code:

MVX Code:

Description:

White to off-white, oval shaped tablets, debossed with "ABR" on one side and "250" on other side.

Active Ingredient(s):

Abiraterone acetate USP

URL for Additional Product Information:

Address:

20 Duke Road

City:

Piscataway

Key Contact:

Phone Number:

908-360-1500

Product Therapeutic Classification:

State:

NJ

Address 2:

Suite A

Email:

sales@novadozpharma.com

Fax:

732-902-2113

Zip:

08854

ADDITIONAL PRODUCT 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:

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)?

No

Direct-Ship Only

Neither

PRODUCT DESCRIPTION INFORMATION

Size:

120

Strength:

250mg

Dosage Form:

Tablets

Product Shape:

Oval

Product Color:

White to Off-White

Product Imprint:

"ABR" on one side and "250" on other side

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?

No

Is this product to be shipped to customers on dry ice?

No

b. Contact for temperature excursion questions:

Name:

P. Krishna Reddy

Number:

040-30449311

Group E-mail:

krishna_p@msnlabs.com

c. Special regulations for product in any states?

Special returns requirements for this product?

No

d. Store product (unit of sale) upright?

Protect product (unit of sale) from light?

Yes

e. Shelf life:

Initial shelf life at launch (if different):

24

 Months

ORDER INFORMATION

Unit of Sale

x

 Bottle

Box/ Carton

Ampule

Glass

Tube

Vial Liquid Sgl

Vial Liquid Multi

Vial Powder Sgl

Vial Powder Multi

Other: Write In

What is the NDC selling unit?

1 bottle of 120 Tablets

(Write-in, e.g. 1 Box of 10 Vials)

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?

1 bottle

(Write-in, e.g. 1 Vial)

HCPCS J-Code:

Rx billing unit to pharmacy:

Each

Gram

Milliliter

ITEM AND PACKING INFORMATION

Item/Each:	Weight Lbs.	Dimensions (US msmts.)			Volume (Cube)	Saleable # Pieces
		Depth	Width	Height		
Box/ Carton/ Bundle/ Inner Pack:	0.24	2.19	2.19	4.01	19.179706	1
Case:	NA	NA	NA	NA	NA	NA
Pallet:	6.86	13.58	9.05	5.90	725.1041	24
	451.28	47.24	39.37	38.38	71380.613	1,152

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	1		00372205030923	00372205030923
x	Box/ Carton/ Bundle/ Inner Pack	NA		NA	
Yes	Case	24		30372205030924	
Yes	Pallet	1152			

COST INFORMATION

Regular Cost

Invoice Cost (WAC) (\$)

As of date:

1/27/2025

Vendor #:

Whsl. Code #:

Fineline Code:

\$150.00

WHOLESALE USE ONLY:

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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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?

No

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?

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?

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 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:

No

☐ 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)

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

Restricted to retail pharmacy only:

Restricted to hospital, clinics, and physician offices only:

Restricted from US territories? (explain in comments)

Comments:

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?

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:

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

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: 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? ☐