



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

Version 2021 Introduction Type: New Item Final Version Date: 7/22/2024

PRODUCT INFORMATION **SPECIAL HANDLING AND STORAGE REQUIREMENTS***

Company Name: Camber Pharmaceuticals, Inc. Application: ANDA
 Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): 207418
 Medical Device Class, if applicable:
 DUNS: 11-856-3719
 Proprietary Name (If Applicable) and Established Name: Oxycodone Hydrochloride Tablets, USP 30 mg
 Selling Unit NDC: 31722-918-01 Unit of Use NDC: UPC: 331722918015
 UDI: CVX Code: MVX Code:
 Description: Oxycodone Hydrochloride Tablets, USP 30 mg
 Active Ingredient(s): Oxycodone hydrochloride, USP
 URL for Additional Product Information: www.camberpharma.com
 Address: 800 Centennial Ave, Suite 1 Address 2:
 City: Piscataway State: NJ Zip: 08854
 Key Contact: Customer Service Email: customerservice@camberpharma.com
 Phone Number: 1-866-827-3647 Fax: 732-562-8788
 Product Therapeutic Classification: Opioid agonist

a. Temperature – Indicate the USP temperature range for this product.
 Temperature Range:
 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: Soma Raju
 Number: 732-529-0423
 Group E-mail: somaraju@heterousa.com

c. Special regulations for product in any states?
 Special returns requirements for this product? *Yes *Yes

d. Store product (unit of sale) upright? No
 Protect product (unit of sale) from light? No

e. Shelf life:
 Initial shelf life at launch (if different): Months

| ADDITIONAL PRODUCT INFORMATION | | PRODUCT DESCRIPTION INFORMATION | |
|--|------------------------------|---|--|
| The product is a legend device? if yes, enter class # a product kit? | <input type="checkbox"/> No | Is the Product... Direct-Ship Only | <input type="checkbox"/> |
| if yes, list NDCs of component parts reverse numbered? | <input type="checkbox"/> No | Is the Product... Orphan Drug Status | <input type="checkbox"/> Neither |
| co-licensed? | <input type="checkbox"/> No | FDA Approval Status | <input type="text"/> |
| latex-free? | <input type="checkbox"/> No | Allergens Present | <input type="text" value="Lactose, Dairy, Alcohol, Animal, Sugar, Rennet, Casein, Whey"/> |
| preservative-free? | <input type="checkbox"/> Yes | Country of Origin | <input type="text" value="USA"/> |
| correctional institution block? opioid? | <input type="checkbox"/> No | Is this product covered under the Trade Agreements Act (TAA)? | <input type="checkbox"/> Yes |
| Cannabinoid? | <input type="checkbox"/> No | Size: | <input type="text" value="100 ct"/> |
| If Unit Dose, is item bar coded to unit dose for hospital scanning? | <input type="checkbox"/> | Strength: | <input type="text" value="30 mg"/> |
| If Unit Dose, indicate NDC here: | <input type="text"/> | Dosage Form: | <input type="text" value="Tablet"/> |
| | | Product Shape: | <input type="text" value="Round, flat faced, beveled edge"/> |
| | | Product Color: | <input type="text" value="Light yellow"/> |
| | | Product Imprint: | <input type="text" value="Debossed with 'T' and '189' with functional scoreline on one side and plain on the other side"/> |

ORDER INFORMATION

| Unit of Sale | What is the NDC selling unit? |
|--|---|
| <input checked="" type="checkbox"/> Bottle | <input type="text" value="1 Bottle of 100 Tablets"/> |
| <input type="checkbox"/> Box/Carton | <input type="text" value="(Write-in, e.g. 1 Box of 10 Vials)"/> |
| <input type="checkbox"/> Ampule | |
| <input type="checkbox"/> Glass | Minimum order quantity? <input type="checkbox"/> Yes |
| <input type="checkbox"/> Tube | |
| <input type="checkbox"/> Vial Liquid Sgl | |
| <input type="checkbox"/> Vial Liquid Multi | |
| <input type="checkbox"/> Vial Powder Sgl | If Yes, how many of which package type? |
| <input type="checkbox"/> Vial Powder Multi | <input type="text" value="24"/> Each |
| <input type="checkbox"/> Other: Write In | <input type="text"/> Inner/ Carton/ Pack |
| | <input type="text"/> Case |

FOR GENERIC DRUG PRODUCTS

I. Orange Book Rating: Authorized Generic *If Authorized Generic, other section fields are not applicable
 II. Generic Equivalent to What Brand?:

PHARMACY ORDER / BILL UNIT

Rec. sell unit to customer? Rx billing unit to pharmacy:
 (Write-in, e.g. 1 Vial) Each
 Gram
 Milliliter

DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION

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

ITEM AND PACKING INFORMATION

| Item/Each: | Weight Lbs. | Dimensions (US msmts.) | | | Volume (Cube) | Saleable # Pieces |
|--------------------------------|-------------|------------------------|-------|--------|---------------|-------------------|
| | | Depth | Width | Height | | |
| Box/Carton/Bundle/ Inner Pack: | 0.07 | 1.56 | 1.56 | 2.94 | 7.16 | 1 |
| Case: | 2.14 | 10.2 | 7 | 3.4 | 242.76 | 24 |
| Pallet: | | | | | | |

GTIN AND HIBCC PRODUCT INFORMATION

| Saleable Unit of Measure | Saleable Quantity | HIBCC | GTIN-14 | Unit of Use GTIN-14 |
|---|---------------------------------|----------------------|---|----------------------|
| <input checked="" type="checkbox"/> Item/Each | <input type="text" value="1"/> | <input type="text"/> | <input type="text" value="00331722918015"/> | <input type="text"/> |
| <input type="checkbox"/> Box/Carton/Bundle/Inner Pack | <input type="text"/> | <input type="text"/> | <input type="text"/> | <input type="text"/> |
| <input checked="" type="checkbox"/> Case | <input type="text" value="24"/> | <input type="text"/> | <input type="text" value="10331722918012"/> | <input type="text"/> |
| <input type="checkbox"/> Pallet | <input type="text"/> | <input type="text"/> | <input type="text"/> | <input type="text"/> |

COST INFORMATION **WHOLESALE USE ONLY:**

Regular Cost
 Invoice Cost (WAC) (\$)
 As of date:
 Vendor #:
 Whsl. Code #:
 Finline Code:

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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?
 (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) No

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) No

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: 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#

SDS Hazard Classification

Organic
 Inorganic
 Steroid/Androgen

Corrosive
 Oxidizer
 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? Yes
 No
 If Yes, is it managed with a pharmacy registry? No
 Website URL: <https://opioidanalgesicrems.com/home.html>

Med Guide Required Yes
 Limited Distribution Requirement No
 Comments / Details: (For example, iPledge program?)

REMS: Yes

REMS Program Manager Name: Prathima Arrabelly Phone: (631) 881-4614 Ext. 1412
 Supplier Manages REMS registry exclusively:
 Wholesale distributor support:
 Provider Name: DEA #:
 Site Enrollment Number assigned by Supplier: NCPDP#:
 NPI #:

Comments

Registry: No

Registry Program Contact Name: Phone:
 Comments

ADD'L STORAGE INFORMATION

Is the Product...
 Controlled Substance? Yes Controlled Substance Code: 9143
 Controlled by State(s)? Yes Listed Chemical (List I or II) No
 ARCOS Reportable? Yes If yes, indicate which:
 Schedule No. 2 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:

RETURN INSTRUCTIONS

Contact tel. # if product received damaged: 1-866-827-3647

Is product returnable for credit: Yes

URL/Link to returns policy: contact - customerservice@camberpharma.com

Special regulations or returns requirements for this product in certain states? Yes

If so, which states? Other requirements? Comments?

DEA Form 222 or its electronic equivalent is required for all returns in all states.

MISCELLANEOUS NOTES and/or Image of Product Barcode:

*Storage of this product must abide by the federally mandated DEA requirements outlined in 21 CFR Part 1301.72.

