



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

Version 2024		Introduction Type: New Item		x Final Version		Date: 7/22/2025			
PRODUCT INFORMATION						SPECIAL HANDLING AND STORAGE REQUIREMENTS*			
Company Name: Camber Pharmaceuticals, Inc. Application Number for NDA/ANDA/BLA; PMA/510(k): 209591 Medical Device Class, if applicable: DUNS: 11-856-3719 Proprietary Name (If Applicable) and Established Name: Amlodipine and Olmesartan Medoxomil Tablets, USP 10 mg/20 mg Selling Unit NDC: 31722-447-90 Unit of Use NDC: 31722-447-90 UPC: 331722447904 CVX Code: MVX Code: Description: Amlodipine and Olmesartan Medoxomil Tablets, USP 10 mg/20 mg Active Ingredient(s): Amlodipine besylate, USP and olmesartan medoxomil, USP URL for Additional Product Information: www.camberpharma.com Address: 800 Centennial Ave, Suite 1 City: Piscataway Key Contact: Customer Service Phone Number: 1-866-827-3647 State: NJ Address 2: Zip: 08854 Email: customerservice@camberpharma.com Fax: 732-562-8788 Product Therapeutic Classification: Dihydropyridine calcium channel blocker and angiotensin II receptor blocker						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: 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? No d. Store product (unit of sale) upright? No Protect product (unit of sale) from light? No e. Shelf life: 24 Months Initial shelf life at launch (if different): Months			
ADDITIONAL PRODUCT INFORMATION				PRODUCT DESCRIPTION INFORMATION					
The product is? a legend device? No if yes, enter class # a product kit? No if yes, list NDCs of component parts reverse numbered? co-licensed? No latex-free? Yes preservative-free? No correctional institution block? No opioid? No Cannabinoid? No If Unit Dose, is item bar coded to unit dose for hospital scanning? If Unit Dose, indicate NDC here:				Is the Product... Direct-Ship Only Is the Product... Unit of Use Orphan Drug Status FDA Approval Status Allergens Present Alcohol, Opadry II Yellow 85F520064.IH, Opadry II White 85F28751.IH, Opadry II Beige 85F570009.IH, Opadry II Brown 85F565953.IH Country of Origin India Is this product covered under the Trade Agreements Act (TAA)? No				Size: 90 ct Strength: 10 mg/20 mg Dosage Form: Film coated tablet Product Shape: Round, biconvex Product Color: Grayish orange Product Imprint: Debossed with 'H' on one side and 'A 10' on the other side	
FOR GENERIC DRUG PRODUCTS						ORDER INFORMATION			
I. Orange Book Rating: AB II. Generic Equivalent to What Brand?: Azor Authorized Generic *If Authorized Generic, other section fields are not applicable						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 90 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? Rx billing unit to pharmacy: Each Gram Milliliter (Write-in, e.g. 1 Vial) HCP/CS J-Code:									
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION									
Does supplier meet DSCSA definition of manufacturer? Yes 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: 0331722498975 GCP: If yes, was original product purchased direct from mfr? Provide source manufacturer for repackaged product									
GTIN AND HIBCC PRODUCT INFORMATION									
Saleable Unit of Measure RFID tag(Y/N) Saleable Quantity HIBCC GTIN-14 Unit of Use GTIN-14 x Item/Each N 1 00331722447904 00331722447904 x Box/ Carton/ Bundle/ Inner Pack N 24 20331722447908 Case Pallet									
COST INFORMATION				WHOLESALE USE ONLY:					
Regular Cost Invoice Cost (WAC) (\$) \$45.00 As of date: 7/14/2025				Vendor #: Whsl. Code #: 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?

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?
(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

No

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)

No

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

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 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

No

Limited Distribution Requirement

Comments / Details: (For example, iPledge program?)

REMS:

No

REMS Program Manager Name:

Phone:

Supplier Manages REMS registry exclusively:

Wholesale distributor support:

Provider Name:

Site Enrollment Number assigned

by Supplier:

DEA #:

NCPDP#:

NPI #:

Comments

Registry:

No

Registry Program Contact Name:

Phone:

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?

No

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? ☐