

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

| Version 2021 | | | | | | Introduction | Туре: | Post Launch Change | [[| x Final Version | | | Date: | 6/6/2 | 2024 | |
|--|---|---------------------|--------------------------|-------------------|-----------------|---|------------------------------|---|---|--------------------------------------|------------------------|-------------------|-----------------------------|------------|------------|--|
| | | | PRODUCT INFORMA | TION | | | | | | SPECIAL HAN | IDLING AND STOP | AGE REQUI | REMENTS* | | | |
| Company Name: Camber Pharmaceuticals, Inc. | | | | Application: ANDA | | a. Temperature – Indicate the USP temperature range for this product. | | | | | | | | | | |
| Application Number for NDA/AN | A/ANDA/BLA (drug); PMA/510(k)(med device): 203835 | | | | | | | Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F) | | | | | | | | |
| Medical Device Class, if applicable: | | | | | | | | | | | | | | | | |
| DUNS: | 11-856-3719 | | | | | | | | | Other Temperature Range | Requirement | | | | | |
| Proprietary Name (If Applicable) a | | Losarta | n Potassium Tablets, USP | | | | | | | (write in) | | | | | | |
| Selling Unit NDC: | 31722-702-10 | | Unit of Use NDC | : | | UPC: | 331722 | 702102 | | Notes | | | | | | |
| UDI | | | CVX Code: | | | MVX Code: | | | | | | | | | | |
| Description: | Losartan Potassium Tablets, | USP 100 r | ng | | | | | | | Is this product to be shippe | | | | No | | |
| | | | | | | | Is this product to be shippe | ed to customers on | dry ice? | | No | | | | | |
| Active Ingredient(s): Losartan potassium, USP | | | | | | h Contact for t | emperature excursion q | unctions | | | | | | | | |
| URL for Additional Product Inform | www.can | nberpharm | a com | | | | | | | Name: | uestions: | Soma Raju | | | | |
| Address: | 800 Centennial Ave, Suite 1 | in or prior in | | | | Address 2: | | | | Number: | | 732-529-042 | 23 | | | |
| City: | Piscataway State: | | | State: | NJ | Zip: | 08854 | Group E-mail: | | | somaraju@heterousa.com | | | | | |
| Key Contact: | Customer Service | | | | customerservice | | | | | | | | | | | |
| Phone Number: | 1-866-827-3647 | I-866-827-3647 Fax: | | | 732-562-8788 | | | c. Special regu | | | No | | | | | |
| Product Therapeutic Classification | n: Angiotensin II receptor blocker (ARB) | | | | | | | | Special returns requirements for this product? | | | | No | | | |
| | | | | | | | | | | | | | | | | |
| | ADDITIONAL PRO | DUCT INF | | | | PRODUCT | DESCRIP | TION INFORMATION | - | ct (unit of sale) upright? | | | | No | | |
| The product is? | | | Is the Product | Direct-Ship C | Only | | | | | Protect product (unit of | sale) from light? | | | No | | |
| a legend device? | No | _ | Is the Product | Neither | | Size: | 1 | 1000 ct | e. Shelf life: | | <i></i> | | | 24 | Months | |
| if yes, enter class # | | _ | Orphan Drug Status | | | | _ | | | Initial shelf life at launch | (if different): | | | | Months | |
| a product kit? if yes, list NDCs of | No | _ | FDA Approval Status | | | Strength: 100 mg | | | ORDER INFORMATION | | | | | | | |
| component parts | | | PDA Approval Status | | | | F | Film coated tablet | | | ORDER INFORM | | | | | |
| reverse numbered? | No | | | | | Dosage For | m: ' | init coaled tablet | | Unit of Sale | | What is the | NDC selling | unit? | | |
| co-licensed? | No | _ | Allergens Present | | | | | | | x Bottle | | 1 Bottle of 1 | 000 Tablets | | | |
| latex-free? | Yes | | | ctose | | Product Sha | T inc | Tear drop | | Box/Carton | | (Write-in, e. | g. 1 Box of 1 | 0 Vials) | | |
| preservative-free? | Yes | | La | ciose | | FIGURE SI | ape. | | | Ampule | | | | | | |
| correctional institution block? | No | | | | | Product Co | lor: V | White to off-white | | Glass | | Minimum o | rder quantity | ? | Yes | |
| opioid? | No | _ | | | | | | | | Tube | | | | | | |
| Cannabinoid? | No | | Country of Origin | India | | Product Imp | | Debossed with 'H' on one side and '145' on the other side | | Vial Liquid Sgl | | | | | | |
| If Unit Dose, is item bar coded to un hospital scanning? | nit dose for | | Is this product covered | under the | | | | | | Vial Liquid Multi Vial Powder Sgl | | If Yes, how 12 | many of wh Each | ch package | type? | |
| If Unit Dose, indicate NDC here: | | | Trade Agreements Act (| | No | | | | Vial Powder Sgl 12 Each Vial Powder Multi Inner/Cartor | | | /Pack | | | | |
| If Unit Dose, indicate NDC nere: | | | | | | | Other: Write In Case | | | | | | | | | |
| | | | FOR GENERIC DRUG PR | ODUCTS | | • | | | | | | | | | | |
| | | | | | | | | | | | | | | | | |
| | | | | | A | uthorized Generic | | orized Generic, other | PHARMACY ORDER / BILL UNIT | | | | | | | |
| | AB | | | | | section fields are not applicable | | | Rec. sell unit to customer? Rx I | | | | x billing unit to pharmacy: | | | |
| II. Generic Equivalent to What Bra | nd?: Cozaar | | | | | | | | | | | | Each | | | |
| | DDU | | CHAIN SECURITY ACT | | MATION | | | | (Write-in, e.g. 1 | Vial) | | | Gram Milliliter | | | |
| | DRU | G SUPPLI | CHAIN SECORITY ACT | DSCSA) INFOR | | | | | - | | | | Milliter | | | |
| Does supplier meet DSCSA definit | tion of manufacturer? | | Yes | | GLN: | 0331722498975 | | | | ITE | M AND PACKING I | FORMATION | ١ | | | |
| Is product exempt from DSCSA? | | | No | | | | | | | | | | | | | |
| If yes, select exemption: | | | | | GCP: | | | | | | Dimensi | ons (US msn | nts.) | Volume | Saleable # | |
| Other exemption - Write in: | | | | | | | | | · [| Weight Lbs. | Depth | Width | Height | (Cube) | Pieces | |
| Is product repackaged? | | | No | | If yes, was o | riginal product | | | Item/Each: | 0.8 | 3 | 3 | 5.75 | 51.75 | 1 | |
| Is product sold by manufacturer's | exclusive distributor? | | Yes | | purchased d | lirect from mfr? | | | | | 3 | 3 | 5.75 | 51.75 | ' | |
| Has FDA granted waiver/exception | | | No | | Provide sou | rce manufacturer f | for repact | kaged product | Box/Carton/Bu | ndle/ | | | | | | |
| If yes, attach documentation from | n FDA. | | | | | | | | Inner Pack: | | | | | | | |
| | | GTIN | AND HIBCC PRODUCT I | | | | | | Case: | 10.7 | 12.5 | 9.5 | 7 | 831.25 | 12 | |
| | | GTIN | AND HIBCC PRODUCT I | NFORMATION | | | | | Pallet: | | | | | | | |
| Saleable Unit of Measure | Saleable Qu | uantitv | HIBCC | | GT | ïN-14 | | Unit of Use GTIN-14 | i unct. | | | | | | | |
| X Item/Each | 1 | | | | | 331722702102 | | | - | | | 1 | | | | |
| Box/Carton/Bundle/Inner Pack | | | | | | | | | | COST INFORMATION | | 1 | NHOLESALI | ER USE ONL | Y: | |
| X Case | 12 | | | | 203 | 331722702106 | | | | | | | | | | |
| Pallet | | | | | | | _ | | Regular Cost | | | Vendor #: | | | | |
| | | | | | | | _ | | Invoice Cost (V | VAC) (\$) | \$257.23 | Whsl. Code | | | | |
| | | | | | | | - | | As of date: | 11/25/2015 | | Fineline Co | ue: | | | |
| | | | | | | | - | | As or date: | 11/20/2010 | | | | | | |
| | | | | | | | | | | | | | | | | |
| 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 infe | ormation on page 2. | | | | , | | | ated Drop Ship Only. | | Signature: | | | | | | |
| | | | | | | • | | | | | | - | | | | |

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

| Version 2021 For Designate | ed Drop Ship Only Products, Please Use Page 3 | | | | | |
|---|---|--|--|--|--|--|
| MATERIAL HAZ | ZARD CLASSIFICATION and TRANSPORTATION | | | | | |
| Is this product (check all that apply): a. Cytotoxic? b. CA Prop. 65 Carcinogen or Reproductive Toxicant? Is the product a CA Prop 65 carcinogen? Is the product a CA Prop 65 reproductive toxicant? Does the product label bear a CA Prop 65 warning? No | x Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard | | | | | |
| 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) | Does the product have an Aerosol class? If yes, identify NFPA Storage Level: No NFPA Storage Level: Image: Storage Level: Is the product a NIOSH hazardous drug? No If yes, indicate which: Image: Storage Level: | | | | | |
| 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 | Hazardous Waste Identification EPA Hazardous Waste Code: Waste Characteristics | | | | | |
| In the product organization of any ment of a manual of a manu | REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? If Yes, is it managed with a pharmacy registry? No Website URL: Image: Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2">Colspan="2"Colsp | | | | | |
| Is the product restricted for air shipment? If so, indicate restriction: No Passenger Cargo Passenger & Cargo La this e scontrible question? | Med Guide Required Limited Distribution Requirement Comments / Details: (For example, iPledge program?) | | | | | |
| 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 | REMS: No REMS Program Manager Name: Phone: Supplier Manages REMS registry exclusively: Phone: Wholesale distributor support: Provider Name: Provider Name: Image: Comments | | | | | |
| Special Provision (listed in Column 7 of 49 CFR 172.101); SP# | Registry: No | | | | | |
| ADD'L STORAGE INFORMATION Is the Product | Registry Program Contact Name: Phone: Comments | | | | | |
| Is the Frouduct No Controlled Substance? Controlled Substance? No Listed Chemical (List I or II) ARCOS Reportable? No If yes, indicate which: Schedule No. Is it a scheduled listed chemical product?: No | RETURN INSTRUCTIONS Contact tel. # if product received damaged: 1-866-827-3647 Is product returnable for credit: Yes | | | | | |
| | URL/Link to returns policy: | | | | | |
| 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: | contact - customerservice@camberpharma.com Special regulations or returns requirements for this product in certain states? If so, which states? Other requirements? Comments? | | | | | |
| | | | | | | |
| MISCELLANEC | DUS NOTES and/or Image of Product Barcode: | | | | | |
| | | | | | | |



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| Version 2021 | FOR DESIGNATED DROP SHIP PRODUCT ONLY - | not a designated drop ship, do not complete. | | | | | |
|--|---|--|---|--|--|--|--|
| Order Method fo | r 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: | 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 Ships same day for next day receipt: Ships for second day receipt: Ships regular ground for 3-10 days receipt: | Days | | | | |
| 1 3 | Name:Phone: | - | _ | | | | |
| Expedited Freight Charge | ges or Other Designated Drop Ship Fees: | Overnight and Priority Overnight PO Processing | | | | | |
| Expedited freight fees billed with each order: | | Overnight receipt available: | | | | | |
| Drop Ship service fee billed with each order: | | PO Receipt cut off time: | | | | | |
| Drop Ship miscellaneous fees billed: Comments: | | | londay uesday /ednesday hursday riday | | | | |
| | | Priority Overnight receipt available: | | | | | |
| Class | s of Trade Restriction: | PO Receipt Cut off time: | | | | | |
| No restriction: Select YES if sold to retail pha Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician of Restricted from US territories? (explain in cor Comments: | | Saturday Overnight receipt available: PO Receipt Cut off time: PO Receipt Cut off time: Phone: Order receipt method: Phone: Fax: EDI: Overnight Fees apply: Other fees apply: | | | | | |
| Other Data Info | rmation 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? | | | | | |
| Mi | scellaneous Notes: | | | | | | |
| | | | | | | | |
| | | ADDITIONAL INFORMATION | | | | | |
| | | Is product order for scheduled patient procedure? Is product order for restocking purposes? | | | | | |