



Date: 10/23/2025

SPECIAL HANDLING AND STORAGE INFORMATION*						
Company Name: Camber Pharmaceuticals, Inc.						
Application Number for NDA/ANDA/BLA; PMA/510(k): 214956		Application: ANDA				
Medical Device Class, if applicable:		NDA 505(b) Type: NOT APPLICABLE				
DUNS: 11-856-3719						
Proprietary Name (If Applicable) and Established Name: Gabapentin Capsules, USP 100 mg						
Selling Unit NDC: 31722-148-05		Unit of Use NDC:				
UDI		UPC: 331722148054				
CVX Code:		MVX Code:				
Description: Gabapentin Capsules, USP 100 mg						
Active Ingredient(s): Gabapentin, USP						
URL for Additional Product Information: www.camberpharma.com						
Address: 800 Centennial Ave, Suite 1		Address 2:				
City: Piscataway		State: NJ				
Key Contact: Customer Service		Zip: 08854				
Phone Number: 1-866-827-3647		Email: customerservice@camberpharma.com				
Fax: 732-562-8788						
Product Therapeutic Classification: Anticonvulsant						
ADDITIONAL PRODUCT 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? No co-licensed? No latex-free? Yes preservative-free? Yes correctional institution block? No opioid? No Cannabinoid? No		Is the Product... Direct-Ship Only Is the Product... Neither Orphan Drug Status FDA Approval Status Allergens Present Corn, Alcohol, Animal, Wheat Country of Origin India If Unit Dose, is item bar coded to unit dose for hospital scanning? If Unit Dose, indicate NDC here:				
PRODUCT DESCRIPTION INFORMATION						
Size: 500 ct						
Strength: 100 mg						
Dosage Form: Hard gelatin capsule						
Product Shape: Capsule						
Product Color: White opaque cap and white opaque body						
Product Imprint: Imprinted with 'A' on cap and '469' on body in black ink						
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?: Neurontin						
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION						
Does supplier meet DSCSA definition of manufacturer? Yes		GLN: 0331722498975 shipments to non-controlled substance states				
Is product exempt from DSCSA? No		0865624000382 shipments to controlled substance states				
If yes, select exemption: Other exemption - Write in:		GCP:				
Is product repackaged? No		If yes, was original product purchased direct from mfr?				
Is product sold by manufacturer's exclusive distributor? Yes		Provide source manufacturer for repackaged product				
Has FDA granted waiver/exception/exemption for product? No						
If yes, attach documentation from FDA.						
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		00331722148054		
x Box/Carton/Bundle/Inner Pack						
x Case	N	12		20331722148058		
x Pallet						
ITEM AND PACKING INFORMATION						
Item/Each:	Weight Lbs.	Dimensions (US msmts.) Depth	Width	Height	Volume (Cube)	Saleable # Pieces
Box/Carton/Bundle/ Inner Pack:	0.36	2.75	2.75	6	45.38	1
Case:	4.9	11.5	8.75	7.5	754.69	12
Pallet:						
COST INFORMATION						
Regular Cost Invoice Cost (WAC) (\$)	\$20.00	Vendor #: Whsl. Code #: Fineline Code:				
As of date:	10/1/2025					
WHOLESALE USE ONLY:						
Signature:						

*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:

Signature:



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?
- 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?

- c. Contact Hazard?
- d. Does this product require special clean-up instructions?
(If yes, attach SDS with special instructions.)
- e. Does the product contain DEHP?

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?

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?

Is the product restricted for air shipment? If so, indicate restriction:

- Passenger
 Cargo
 Passenger & Cargo

Is this a reportable quantity?

RQ Threshold:

Is this a marine pollutant?

Is this product shipped utilizing an authorized DOT exception or Special Permit?

- (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? Controlled Substance Code
- Controlled by State(s)? Listed Chemical (List I or II)
- ARCOS Reportable? If yes, indicate which:
- Schedule No. Is it a scheduled listed chemical product?:

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?

If yes, indicate which:

Hazardous Waste Identification

EPA Hazardous Waste Code:

Waste Characteristics

REMS or REGISTRY RESTRICTIONS

Is there a REMS on this product?

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:

Phone:

DEA #:

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?

This product is classified as a Schedule V controlled substance in Alabama, Kentucky, Montana, North Dakota, Tennessee, Utah, Virginia, and West Virginia.

MISCELLANEOUS NOTES and/or Image of Product Barcode:

Release DATE



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

Version 2024

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? ☐