

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

				Introduction Type:	New Item		1 Final Version			Date:	12/2	/2024
		PRODUCT INFORMAT	ION				SPECIAL HAN	DLING AND STOP	AGE REQUI	REMENTS*		
Company Name:	Camber Pharmaceuticals, Inc.			Application:	ANDA	a. Temperature –	Indicate the USP tempe	erature range for t	his product.			
Application Number for NDA/AN	DA/BLA; PMA/510(k):	209267		NDA 505(b) Type:	NOT APPLICABLE		nperature Range	Controlled Room		and 25 C (68	° – 77° F)	
Medical Device Class, if applical	ble:											
DUNS:	11-856-3719					Oth	er Temperature Range I	Requirement				
Proprietary Name (If Applicable) a		Erlotinib Tablets 150 mg					(write in)					
Selling Unit NDC:	31722-265-30	Unit of Use NDC:	31722-265-30		722265300	Not	es					
UDI		CVX Code:		MVX Code:								
Description:	Erlotinib Tablets 150 mg					ls ti	his product to be shipped	to customers on i	ce?		No	
						ls ti	his product to be shipped	to customers on o	try ice?		No	
Active Ingredient(s):	Erlotinib hyd	rochloride										
UDI for Additional Draduct Inform	unuu combo	pharma.com				b. Contact for tem Nar	perature excursion qu	estions:	Soma Raju			
URL for Additional Product Inform Address:	800 Centennial Ave, Suite 1	pharma.com		Address 2:			mber:		732-529-042	2		
City:	Piscataway		State:	NJ Zi	08854		oup E-mail:			eterousa.cor	n	
Key Contact:	Customer Service		Email:	customerservice@can							_	
Phone Number:	1-866-827-3647		Fax:	732-562-8788		c. Special regulation	ions for product in any	states?			No	
Product Therapeutic Classificatio	n: Kinase inhib	tor				Spe	ecial returns requirement	s for this product?			No	
	1					_						_
	ADDITIONAL PRODU	ICT INFORMATION		PRODUCT DESC	RIPTION INFORMATION	d. Store product (	unit of sale) upright?				No	
The product is?		Is the Product	Direct-Ship Only			Pro	etect product (unit of sa	le) from light?			No	1
a legend device?	No	Is the Product	Unit of Use	Size:	30 ct	e. Shelf life:					24	Months
if yes, enter class #		Orphan Drug Status		0.20.		Init	ial shelf life at launch (	f different):				Months
a product kit?	No			Strength:	150 mg			ORDER INFORM				
if yes, list NDCs of component parts		FDA Approval Status			Film-coated tablet			ORDER INFORM	IATION			
reverse numbered?	No			Dosage Form:	Film-coaled tablet	Uni	it of Sale		What is the	NDC selling	unit?	
co-licensed?	No	Allergens Present					x Bottle		1 Bottle of 3			
latex-free?	Yes			Des des si Olivera	Round, biconvex		Box/Carton			g. 1 Box of 1	0 Vials)	
preservative-free?	Yes	Dairy, L	Lactose	Product Shape:			Ampule					
correctional institution block?	No			Product Color:	White		Glass		Minimum o	der quantity	?	Yes
opioid?	No						Tube					
Cannabinoid?	No	Country of Origin	India	Product Imprint:	Debossed with 'H' on one side and '22' on the other side		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u hospital scanning?	unit dose for	Is this product covered ur	ader the				Vial Liquid Multi Vial Powder Sgl			Each	ch package	type?
If Unit Dose, indicate NDC here:		Trade Agreements Act (T					Vial Powder Multi			Inner/Cartor	/Pack	
		3	,				Other: Write In			Case		
							Other: write in			Case		
		FOR GENERIC DRUG PRO	DDUCTS				Other: white in			Case		
		FOR GENERIC DRUG PRC	DDUCTS				Other: White In			Case		
		FOR GENERIC DRUG PRO			uthorized Generic, other			ARMACY ORDER		Case		
I. Orange Book Rating:	АВ	FOR GENERIC DRUG PRO			uthorized Generic, other ion fields are not applicable	Rec. sell unit to cr	PH	ARMACY ORDER		1	acy:	
I. Orange Book Rating: II. Generic Equivalent to What Bra		FOR GENERIC DRUG PRO				Rec. sell unit to cr	PH	ARMACY ORDER	/ BILL UNIT	1	acy:	
	nd?: Tarceva		At			(Write-in, e.g. 1 Vi	PH ustomer?	ARMACY ORDER	/ BILL UNIT	<b>nit to pharm</b> Each Gram	acy:	
	nd?: Tarceva	FOR GENERIC DRUG PRO	At				PH ustomer?	ARMACY ORDER	/ BILL UNIT	nit to pharm	acy:	
II. Generic Equivalent to What Bra	nd?: Tarceva	SUPPLY CHAIN SECURITY ACT (D	At SSCSA) INFORMATION	sec		(Write-in, e.g. 1 Vi	PH ustomer? al)	]	/ BILL UNIT Rx billing u	<b>nit to pharm</b> a Each Gram Milliliter	acy:	
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## **HDA** Standard Pharmaceutical Product and Medical Device Information (Rx Product Only)

Version 2024 For Desig	nated 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? C. Contact Hazard? (If yes, attach SDS with special instructions.) e. Does the product contain DEHP? No	X       Organic       Corrosive         Inorganic       Oxidizer         Steroid/Androgen       Contact Hazard         Does the product have an Aerosol class? If yes, identify NFPA Storage Level:       No         NFPA Storage Level:       Image: Contact Hazard			
Is this product regulated for shipment by DOT? 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	Is the product a NIOSH hazardous drug? If yes, indicate which: Hazardous Waste Identification			
d. Packing Group e. Inhalation Hazard?	EPA Hazardous Waste Code: Waste Characteristics			
Is this product regulated for shipment by IATA?  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	REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? No			
d. Packing Group         e. Inhalation Hazard?         Is the product restricted for air shipment? If so, indicate restriction:         No	If Yes, is it managed with a pharmacy registry?       Website URL:       Med Guide Required       No			
Passenger         Cargo         Passenger & Cargo         Is this a reportable quantity?	Limited Distribution Requirement       Comments / Details: (For example, iPledge program?)       REMS:			
RQ Threshold:       No         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)	REMS Program Manager Name:     Phone:       Supplier Manages REMS registry exclusively:     Phone:       Wholesale distributor support:     Provider Name:       Provider Name:     DEA #:       Site Enrollment Number assigned     NCPDP#:       by Supplier:     NPI #:			
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101);				
SP#ADD'L STORAGE INFORMATION	Registry:     No       Registry Program Contact Name:     Phone:       Comments			
Is the Product Controlled Substance? No Controlled Substance Code	RETURN INSTRUCTIONS			
Controlled by State(s)?         No         Listed Chemical (List I or II)         No           ARCOS Reportable?         No         If yes, indicate which:         It         It <td< td=""><td>Contact tel. # if product received damaged: Is product returnable for credit: Yes</td></td<>	Contact tel. # if product received damaged: Is product returnable for credit: Yes			
CLASS OF TRADE RESTRICTION: No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Yes	URL/Link to returns policy: contact - customerservice@camberpharma.com			
Restricted to retail pharmacy only:     No       Restricted to hospital, clinics, and physician offices only:     No       Restricted from US territories? (explain in comments)     No	Special regulations or returns requirements for this product in certain states?       No         If so, which states? Other requirements? Comments?			
Comments:				
MISCELLA	NEOUS NOTES and/or Image of Product Barcode:			



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

Version 2024	FOR DESIGNATED DROP SHIP PRODUCT ONLY - if r	ot a designated drop ship, do not complete.	
Order Method for Des	signated 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 Expedited Freight Charges o		Purchase order daily receipt cut off time by supplier         Cut off time:         Shipping lead time of PO:         Hours         Da         Ships same day for next day receipt:         Ships for second day receipt:         Ships regular ground for 3-10 days receipt:         Overnight and Priority Overnight PO Processing	ays
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:		Days of week overnight is available: Monday	
Comments:	s only:	Days of week overnight is available:       Tuesday         Tuesday       Wednessi         Priority Overnight receipt available:       Friday         PO Receipt Cut off time:       PO Receipt Cut off time:         Order receipt method:       Phone:         Fax:       EDI:         Overnight Fees apply:       Policity	/ day
Other Data Informati	ion 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?	
Miscell	aneous Notes:		
		ADDITIONAL INFORMATION	
		Is product order for scheduled patient procedure? Is product order for restocking purposes?	