

Standard Pharmaceutical Product Information (Rx Product Only)

© August 2014						Introdu	ction Type:	Post Launch Change		Final Version			Date:		
			PRODUCT II	IFORMATION						SPECIAL HANDL	ING AND ST	ORAGE REQI	JIREMENTS	*	
Company Name:	Camber Pharmaceut	icals					Application:	ANDA	a. Temperature – Indie	ate the USP temper	ature range f	or this produ	ct.		
Application Number for ND			levice):	2	04343					ature Range				en 20 and 25	C (68° – 77° F
DUNS:	82-667-4775		•						Other Tr	emperature Range Re	quirement	-			
Proprietary Name (If Applica		Name: Du	loxetine Delayed Releas	e Capsules 30MG 30	ICT					rite in)	quironnoni				1
Selling Unit NDC:	31722-582-30		Individual Ur	it NDC:	31722-582-30	1	UPC: 3317225	82308	Ì.	,					4
UDI	NA		CVX Code	:		MVX Co	ode: NA		Is this p	roduct to be shipped t	o customers	on ice?		No	_
Description: Opaque white/opaque blue capsules imprinted with 'H'/191'									Is this product to be shipped to customers on dry ice? No						
Active Ingredient(s): Duloxetine							b. Contact for temperature excursion questions: Name: Soma Raju								
URL for Additional Product I	Information [.]	www.camberpharr	ma com						Number			732-529-042	3		
Address:	1031 Centennial Avenue				Address 2:			Group			somaraju@heterousa.com				
City:	Piscataway				State: NJ Zip: 08854						, 0				
Key Contact:	Customer Service				Email: customerservice@camberpharma.com				c. Special regulations	for product in any s	tates?			No	_
Phone Number:	732-529-0430				Fax: 732-562-8788			Special returns requirements for this product? No					-		
Product Therapeutic Classifi	fication:														
				_	_				d. Store product (unit					No	-
ADDITIONA	AL PRODUCT INFORM	IATION			Р	PRODUCT DE	SCRIPTION INFO	ORMATION	Protect	product (unit of sale	e) from light?			No	-
Is the Product									e. Shelf life:					24	Months
a legend device?		No			Size:	30			Initial s	helf life at launch (if	different):				Months
reverse numbered?		No									ORDER INFO	DMATION			
co-licensed? Is the Product		No Direct-Ship Only	0		Strength:	30	MG			Ļ		RMATION			
Is the Product		Unit of Use							Unit of	Sale		What is the	NDC selling	unit?	
is the routet		01111 01 000			Dosage Form:	cap	osule		x	Bottle		1 bottle of 30			
If their Deeper is items have a de		ital a consistent								Box/Carton			g. 1 Box of 1	0 Vials)	
If Unit Dose, is item bar code	ed to unit dose for nosp	oital scanning?			Product Shape		osule			Ampule			-		
If Unit Dose NDC, indicate N	NDC here:				Froduct Shape	e. ca	Jaule			Glass		Minimum or	der quantity	?	Yes
		les all a			Product Color	: whi	ite/blue			Tube Vial Liquid Sgl					
Country of Origin		India							Vial Liquid Ogi						
Is this product covered under the Trade Agreements Act (TAA)?				Product Imprint: H'/'191'			Vial Powder Sql 12 Each			ype.					
										Vial Power Multi			Inner/Carton	/Pack	
•										Other: Write In	_		Case		
			FOR GENERIC	RUG PRODUCTS											
							*16 Author	ized Generic, other section		DHAR		ER / BILL UNI	Ŧ		
	[Autro	orized Generic		not applicable	-						
I. Orange Book Rating:	AB	O multi alta						nor applicable	Rec. sell unit to custo	mer?	-	Rx billing u	nit to pharm	acy:	
II. Generic Equivalent to What	at Brand?:	Cymbalta							(Write-in, e.g. 1 Vial)				Each Gram		
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION (VITIB-In, e.g. 1 Viai) Gram															
Does supplier meet DSCSA		turer?	Yes	G	LN:	033172200	0000			ITEM AI	ND PACKING	INFORMATI	ON		
Is product exempt from DSC If yes, select exemption:	JON!		NU								Dime	nsions (US m	smts.)	Volume	
Other exemption - Write in:	:									Weight Lbs.	Depth	Height	Width	(Cube)	# Pieces:
Is product repackaged?			No		Yes, was origina	al product pu	rchased direct		Item:	0.05		3	1.4		1
Is product sold by manufact			No		om mfr?					3.00		Ŭ			
Has FDA granted waiver/exc	ception/exemption for	product?	No	If	yes, attach docu	umentation fr	om FDA.		Box/Carton/Bundle/ Inner Pack:						
			GTIN PRODUC						Case:						
				Saleable					0436.	0.85	6.7	3	5		12
			Level	Unit			Quantity	GTIN-14	Pallet:						
Serialized?	Yes		X Item		X 2D	Lin		00331722582308							
If not, when?			X Box/Carton/Bundle/Inn	er Pack X	X 2D	Lin		10331722582305	UPC:	Case:					
Items aggregated?	Yes		X Case		X 2D	Lin		30331722582309		Carton:					
			Pallet		2D	Lin									
					2D 2D	Lin			COST	INFORMATION			WHOLESAL	ER USE ONL	r:
					2D 2D	Line			Regular Cost			Vendor #:			
					2D 2D	Lin			Invoice Cost (WAC) (\$	`	\$14.72	Whsl. Code	<i>#</i> ·		
									Federal Excise Tax Pe		φ14.73	Fineline Co			
-									As of date:	5	1				
										·		1			
			Attach copy of SAF	ETY DATA SHEET (SDS) or non haza	ard letter, PAC	CKAGE INSERT,	LABEL AND PHOTO OF PR	ODUCT PACKAGING and E	ARCODE.					
*Please provide any addition	nal information on pag	je 2.						d Drop Ship Only.	Signatu						
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Standard Pharmaceutical Product Information (Page 2)

	jnated Drop Ship Only Products, Please Use Page 3					
MATERIAL	HAZARD CLASSIFICATION and TRANSPORTATION					
Is this product (check all that apply):						
a. Cytotoxic? No	SDS Hazard Classification					
b. CA Prop. 65 Carcinogen or Reproductive Toxicant?						
Is the product a CA Prop 65 carcinogen? No	Organic Corrosive					
Is the product a CA Prop 65 reproductive toxicant?	Inorganic Oxidizer					
Does the product label bear a CA Prop 65 warning?	Steroid/Androgen Contact Hazard					
c. Contact Hazard? No	Aerosol Class; Identify NFPA Storage Level:					
d. Does this product require special clean-up instructions? No						
(If yes, attach SDS with special instructions.)	Is the product a NIOSH hazardous drug?					
e. Does the product contain DEHP? No	If yes, indicate which:					
Is this product regulated for shipment by DOT or IATA? No						
(if yes, answer a-e below and provide SDS)						
a. UN/Identification Number	Hazardous Waste Identification					
b. Proper Shipping Name						
c. DOT Hazard Class	EPA Hazardous Waste Code: NA					
d. Packing Group						
e. Inhalation Hazard?						
Is the product restricted for air shipment? If so, indicate restriction:	REMS or REGISTRY RESTRICTIONS					
Passenger	Is there a REMS on this product? No					
Cargo	If Yes, is it managed with a pharmacy registry?					
Passenger & Cargo	Website URL:					
Is this a reportable quantity? No						
RQ Threshold:	Comments / Details: (For example, iPledge program?)					
Is this a marine pollutant? No						
Is this product shipped utilizing an authorized DOT exception or Special Permit?						
No (if yes, identify method below)	REMS:					
Limited Quantity	REMS Program Manager Name: Phone:					
Consumer Commodity, ORM-D	Supplier Manages REMS registry exclusively: No					
Small Quantity (49 CFR 173.4)	Wholesale distributor support: No					
Special Permit; DOT-SP	Provider Name:					
Special Provision (listed in Column 7 of 49 CFR 172.101);	Site Enrollment Number assigned DEA #: No					
SP#	by Supplier: PCPDP #: No					
	NPI #: No					
ADD'L STORAGE INFORMATION						
Is the Product	Comments					
Controlled Substance? No	Continents					
	Projekter No.					
	Registry: No					
ARCOS Reportable? No	Registry Program Contact Name: Phone: Phone:					
Schedule No. (inc. N for non-narcotic)	Comments					
Controlled Substance Code						
Listed Chemical (List I or II) No	RETURN INSTRUCTIONS					
If yes, indicate which:						
Is it a scheduled listed chemical product?: No	Contact tel. # if product received damaged: 732-529-0430					
CLASS OF TRADE RESTRICTION:	Is product returnable for credit: Yes					
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	Special regulations or returns requirements for this product in certain states? No					
Restricted to hospital, clinics, and physician offices only: No	If so, which states? Other requirements? Comments?					
Restricted from US territories? (explain in comments) No	In so, which states? Other requirements? Comments?					
Comments:						
MISCELLA	ANEOUS NOTES and/or Image of Product Barcode:					



Standard Pharmaceutical Product Information (Page 3)

FOR DESIGNATED DROP SHIP PRODUCT ONLY - if	
Order Method for Designated Drop Ship Product	Standard Order Receipt and Processing
Purchase orders may be accepted by: a. EDI b. Autofax Fax Number: c. Fax Fax Number: d. Phone only Phone No.: e. Supplier Web Site only Site Address: Minimum Order Quantity:	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:
Contracted 3PL company / contact #: Name:	
Phone:	
Expedited Freight Charges 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:	Days of week overnight is available: Tuesday Wednesday Thursday Friday
	Priority Overnight receipt available:
Class of Trade Restriction:	PO Receipt Cut off time:
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:	Saturday Overnight receipt available: PO Receipt Cut off time: Order receipt method: Phone: Fax: EDI: Overnight Fees apply: Overnight Fees apply:
Other Data Information Required to Process PO:	Return Instructions
Patient Procedure Date: Physician Name: Physician/Clinic Phone # Physician/Clinic DEA #: Physician/Clinic Specialty: Miscellaneous Notes:	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?
	ADDITIONAL INFORMATION
	Is product order for scheduled patient procedure?
	Is product order for restocking purposes?