

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

Version 2024						Introduction T	уре:	New Item		x Final Version			Date:	3/10/	2025	
			PRODUCT INFORMA	TION						SPECIAL HAN	DLING AND STOR	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	IDA/BLA; PMA/510	0(k):	207022			NDA 505(b) Type:	NOT	T APPLICABLE		Temperature Range	Controlled Room -		and 25 C (68	° – 77° F)		
Medical Device Class, if applical	ble:															
DUNS:	11-856-3719									Other Temperature Range F	Requirement		permitted to 1	5°C to		
Proprietary Name (If Applicable) a		ame:	Mycophenolate Mofetil Capsules,							(write in)		30°C (59°F	to 86°F)			
Selling Unit NDC:	31722-878-05		Unit of Use NDC:			UPC:	3317228780	050		Notes						
UDI			CVX Code:			MVX Code:										
Description:	Mycophenolate M	Mofetil Capsules	s, USP 250 mg							Is this product to be shipped				No		
										No						
Active Ingredient(s): Mycophenolate mofetil, USP									h Cambant fan							
URL for Additional Product Information: www.camberpharma.com								b. Contact for temperature excursion questions: Name: Soma Raju								
Address:	800 Centennial A		priama.com		1	Address 2:			1	Number:		732-529-042	23			
City:	Piscataway		Sta				NJ Zip: 08854							omaraju@heterousa.com		
Key Contact:	Customer Service					customerservice@camberpharma.com										
Phone Number:	1-866-827-3647					Fax: 732-562-8788			c. Special regulations for product in any states?			No				
Product Therapeutic Classificatio	n:	Antimetaboli	te immunosuppressant							Special returns requirement	s for this product?			No		
	ADDITI	IONAL PRODU	ICT INFORMATION			PRODUCT	DESCRIPTIO	N INFORMATION	d. Store produ	uct (unit of sale) upright?				No		
The product is?			Is the Product	Direct-Ship C	Only					Protect product (unit of sa	le) from light?			No		
a legend device?		No	Is the Product	Neither		Size:	500 0	ct	e. Shelf life:					24	Months	
if yes, enter class #		NI.	Orphan Drug Status				050			Initial shelf life at launch (i	f different):				Months	
a product kit? if yes, list NDCs of		No	FDA Approval Status			Strength:	250 r	mg			ORDER INFORM	MATION				
component parts			T DA Approvai Status				Hard	gelatin capsule			ONDER IN ON	.,				
reverse numbered?		No				Dosage Forn	n:	goldin odpodio		Unit of Sale		What is the	NDC selling	unit?		
co-licensed?		No	Allergens Present							x Bottle		1 Bottle of 5	00 Capsules			
latex-free?		Yes	Dye	, Corn		Product Sha	Caps	sule		Box/Carton		(Write-in, e.	.g. 1 Box of 1	0 Vials)		
preservative-free?		Yes		, 00111		1 Todact Ona				Ampule						
correctional institution block?		No				Product Cold	or: Blue	cap and brown body		Glass		Minimum o	rder quantity	?	Yes	
opioid?		No	Occupation of October	India			Lance of	Catada 205 B B an ann		Tube						
Cannabinoid? If Unit Dose, is item bar coded to u	unit dose for	No	Country of Origin	IIIuia		Product Imp		inted with 'H' on cap M1' on body		Vial Liquid Sgl Vial Liquid Multi		If Voc. how	many of whi	ch nackago (wno2	
hospital scanning?	unit dose for		Is this product covered of	inder the			and	iii oii boay		Vial Powder Sal			Each	cii package i	ype:	
If Unit Dose, indicate NDC here:			Trade Agreements Act (No					Vial Powder Multi			Inner/Carton	/Pack		
		1								Other: Write In			Case			
			FOR GENERIC DRUG PR	ODUCTS												
										D.I.	ADMAQY ODDED	/ DULL LINUT				
				_	Au	thorized Generic		ed Generic, other			ARMACY ORDER					
I. Orange Book Rating:	AB					section fields are not applicable			Rec. sell unit to customer? Rx billing unit to pharmacy:				acy:			
II. Generic Equivalent to What Brand?: Cellcept							(Write-in, e.g. 1 Vial) Each									
		DRUG S	SUPPLY CHAIN SECURITY ACT	(DSCSA) INFOR	RMATION				HCPCS J-Cod	e:			Milliliter			
				,]		-			
Does supplier meet DSCSA defini	ition of manufactu	irer?	Yes		GLN:	0331722498975				ITEN	AND PACKING I	NFORMATIO	N			
Is product exempt from DSCSA?			No													
If yes, select exemption:					GCP:					Weight Lbs.		ons (US msr	•	Volume	Saleable #	
Other exemption - Write in:			No		1	data da a control					Depth	Width	Height	(Cube)	Pieces	
Is product repackaged? Is product sold by manufacturer's	avalucius distribu	utor?	Yes	_	direct from n	riginal product purd	cnased		Item/Each:	0.60	3.41	3.41	6.1	70.93	1	
Has FDA granted waiver/exceptio			No	+		ce manufacturer fo	r repackage	d product	Box/Carton/B	undle/						
If yes, attach documentation from									Inner Pack:							
									Case:	8.35	14.5	11	7.25	1156.38	12	
			GTIN AND HIBCC PRODUCT I	NFORMATION						0.55	14.0		7.20	1100.00		
Saleable Unit of Measure	DEID : 0//AD	0-1	HIBCC		0.71	N-14	11-1	t of Use GTIN-14	Pallet:							
Saleable Unit of Measure	RFID tag(Y/N)	Quantity	HIBCC		GII	N-14	Uni	t of Use GTIN-14								
x Item/Each	N	Quantity 1			003	31722878050										
Box/Carton/Bundle/Inner Pack										COST INFORMATION			WHOLESALI	ER USE ONL	Y:	
X Case	N	12			203	31722878054										
Pallet									Regular Cost			Vendor #:				
									Invoice Cost (WAC) (\$)	\$90.00	Whsl. Code				
							-		As of date:	1/27/2025		Fineline Co	de:			
							-		As of date:	1/2//2020		1				
			Attach copy of SAFETY D	ATA SHEET (SD	S) or non haza	ard letter, PACKAGE	INSERT, LAI	BEL AND PHOTO OF F	PRODUCT PACKA	GING and BARCODE.						
*Please provide any additional inf			• • •	· -		See new n 3 for										



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For Designated Drop Ship Only Products, Please Use Page 3

MATERIAL HAZ	ARD CLASSIFICATION and TRANSPORTATION						
Is this product (check all that apply):							
a. Cytotoxic?	SDS Hazard Classification						
	ODO Hazaru Classification						
b. CA Prop. 65 Carcinogen or Reproductive Toxicant?							
Is the product a CA Prop 65 carcinogen?	x 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?	Does the product have an Aerosol class? If yes, No						
d. Does this product require special clean-up instructions?	identify NFPA Storage Level:						
(If yes, attach SDS with special instructions.)	NFPA Storage Level:						
e. Does the product contain DEHP?							
Is this product regulated for shipment by DOT?	Is the product a NIOSH hazardous drug?						
(if yes, answer a-e below and provide SDS)	If yes, indicate which: Group 2 items (non-antineoplastic that meets a hazard criterion)						
a. UN/Identification Number							
b. Proper Shipping Name							
c. DOT Hazard Class	Hazardous Waste Identification						
d. Packing Group							
e. Inhalation Hazard?	EPA Hazardous Waste Code: Waste Characteristics						
Is this product regulated for shipment by IATA?	DEUG . DEGLOTAV ARGERIQUO						
(if yes, answer a-e below and provide SDS)	REMS or REGISTRY RESTRICTIONS						
a. UN/Identification Number							
b. Proper Shipping Name	Is there a REMS on this product?						
c. DOT Hazard Class	If Yes, is it managed with a pharmacy registry?						
d. Packing Group	Website URL: www.MycophenolateREMS.com						
e. Inhalation Hazard?							
Is the product restricted for air shipment? If so, indicate restriction:	Med Guide Required Yes						
Passenger	Limited Distribution Requirement No						
Cargo	Comments / Details: (For example, iPledge program?)						
	Confinents / Details. (For example, iFredge program?)						
Passenger & Cargo							
Is this a reportable quantity? No	REMS: Yes						
RQ Threshold:	REMS Program Manager Name: Allison Prezioso Phone: 732-529-0430 ext. 465						
Is this a marine pollutant? No	Supplier Manages REMS registry exclusively: No						
Is this product shipped utilizing an authorized DOT exception or Special Permit?	Wholesale distributor support: No						
No (if yes, identify method below)	Provider Name: DEA #:						
Limited Quantity	Site Enrollment Number assigned NCPDP#:						
Consumer Commodity, ORM-D	by Supplier: NPI #:						
Small Quantity (49 CFR 173.4)	Бу Саррио.						
Special Permit; DOT-SP	Comments						
Special Provision (listed in Column 7 of 49 CFR 172.101);	Continents						
SP#	Registry: Yes						
	Registry Program Contact Name: REMS Call Center Phone: 1-800-617-8191						
ADD'L STORAGE INFORMATION	Comments Mycophenolate Mofetil REMS is a shared REMS program.						
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:	Contact tel. # if product received damaged: 1-866-827-3647						
Schedule No. Is it a scheduled listed chemical product?: No							
	is product returnable for credit.						
CLASS OF TRADE RESTRICTION:	URL/Link to returns policy:						
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Yes	contact - customerservice@camberpharma.com						
Restricted to retail pharmacy only: No	Out the land of the control of the c						
	Special regulations or returns requirements for this product in certain states?						
Restricted to hospital, clinics, and physician offices only: No	110						
Restricted from US territories? (explain in comments)	If so, which states? Other requirements? Comments?						
Comments:							
MISCELLANEC	OUS NOTES and/or Image of Product Barcode:						
	kin or mucous membranes of the powder contained in mycophenolate mofetil capsules. Follow applicable special handling and disposal						
procedures in "OSHA Hazardous Drugs." http://www.osha.gov/SLTC/hazardousdrugs/index.htm							



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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 Fax Number:	Purchase order daily receipt cut off time by supplier Cut off time:							
c. Fax d. Phone only Phone No.:	Shipping lead time of PO: Hours Days							
e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Name: Phone:	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:	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: Monday 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: Phone: Fax: EDI: Overnight Fees apply: Other 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:	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?							