

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

Version 2020						Introduction Type:	New Item		Final Version	n		Date:	3/19	9/2021
			PRODUCT INFORMAT	ION					SPECIAL	HANDLING AND STO	RAGE REQU	REMENTS*		
Company Name: Amneal Pharmaceuticals LLC Application: ANDA Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): 210043									a. Temperature – Indicate the USP temperature range for this product. Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)					
DUNS:	827748190	mirao ro(k)(mea aevi		2.00.					Other Temperature R		. 5000000000000000000000000000000000000	7 and 20 0 (0	,	
Proprietary Name (If Applicable)		ame: Methyli	PREDNISolone Acetate Injec	table Suspension.	. USP 80ma	ı/mL			(write in)	inge Requirement				
Selling Unit NDC:	70121-1574-1	,	Unit of Use NDC:		,		2115741-0		Notes					
UDI			CVX Code:			MVX Code:								
Description:	Is this product to be shipped to customers on ice? Is this product to be shipped to customers on dry ice? No													
Active Ingredient(s):	h Contact for	temperature excursi		,			_							
URL for Additional Product Infor	mation:	b. Contact for	Name:	ni questions.										
Address:	400 Crossing Blvd. Third Floor Address 2:								Number: 877-835-5472 option 1					
City:	Bridgewater				Group E-mail:									
Key Contact: Phone Number:	866-525-7270				Email: Fax:	866-525-7271		a Charial sam	ulatiana fan muaduat i				No	
Product Therapeutic Classification		Corticosteroid			гах.	800-323-7271		c. Special reg	ulations for product i	ements for this product)		No	-
Troduct merapeutic classification	JII.	Corticosteroid							Opeciai retarris requii	ments for this product	•			-
	ADDITIO	d. Store produ	uct (unit of sale) uprig	ht?			Yes							
The product is?			Is the Product	Direct-Ship Only	,				Protect product (uni	of sale) from light?			No	-
a legend device?		No	Is the Product			Size:	1mL Single-Dose Vial	e. Shelf life:	•	-			24	Months
if yes, enter class #			Orphan Drug Status			*	00		Initial shelf life at lau	nch (if different):				Months
a product kit? if yes, list NDCs of		No	FDA Approval Status	80mg/mL	ORDER INFORMATION									
component parts			1 DA Appiovai otatas			B 5	Injectable Suspension			3113 <u>2</u> 111111 311				
reverse numbered?		No	•			Dosage Form:			Unit of Sale		_	NDC selling		
co-licensed?		No	Allergens Present						Bottle			1mL Single-E		
latex-free? preservative-free?		Yes Yes				Product Shape:			x Box/Carton Ampule		(Write-in, e	e.g. 1 Box of 1	10 Vials)	
correctional institution block?		No					white to off-white		Glass		Minimum o	order quantit	v?	Yes
opioid?		No				Product Color:	homogeneous		Tube				•	
Cannabinoid?		No	Country of Origin	India		Product Imprint:			Vial Liquid S					
If Unit Dose, is item bar coded to	unit dose for hospita	ıl	In this was don't account to	den de e					Vial Liquid I		If Yes, how 240		nich package	type?
scanning? If Unit Dose, indicate NDC here:			Is this product covered un Trade Agreements Act (T.		0				Vial Powder Vial Power		240	Each Inner/Cartor	n/Pack	
III STINCESCO, INGIGARO (125 Note:				, <u></u>	<u> </u>				Other: Write			Case	ivi don	
			FOR GENERIC DRUG PRO	DUCTS								_		
										DUADMACY ORDE	D / DILL LINIT			
	AB			- <u>L</u>	Autric		norized Generic, other section are not applicable							
I. Orange Book Rating: II. Generic Equivalent to What Br		DEPO MEDROL						Rec. Sell ullit	to customer?		KX billing t	unit to pharm Each	nacy:	
,								(Write-in, e.g.	1 Vial)			Gram		
		DRUG SUPPLY	Y CHAIN SECURITY ACT (D	SCSA) INFORM	ATION							Milliliter		
Does supplier meet DSCSA defin	nition of manufactu	rer?	Yes	GLN:						ITEM AND PACKING	INFORMATIO	N		
Is product exempt from DSCSA?			No	_										
If yes, select exemption:									Weight L	he	sions (US ms		Volume	# Pieces:
Other exemption - Write in:			No	lf Van	aa aslada	and were decent wound become		Item/Each:	1	Depth	Width	Height	(Cube)	
Is product repackaged? Is product sold by manufacturer!	s exclusive distribu	utor?	No		from mfr?	nal product purchased		item/Each.	14.74 g	m 1.1875"	1.5"	1.8125"		1
Has FDA granted waiver/exception			No	If yes	, attach doo	cumentation from FDA.		Box/Carton/B	undle/					
				-				Inner Pack:						
		GIIN	AND HIBCC PRODUCT IN	FORMATION				Case:	8.3 lb	16.0625"	8.25"	8.625"		240
Saleable Unit of Measure		Quantity	HIBCC		GTIN-	14	Unit of Use GTIN-14	Pallet:	621 lb	40.813"	33"	65.375"		70
X Item/Each		1			00370	121157410			62110	40.613	33	05.375		70
Box/Carton/Bundle/Inner Pack		240			E0070	121157415			COST INFORMA	ION		WHOLESAL	ER USE ON	I V.
X Case									COSTINFORMA	TON		WHOLESAL	LER USE ON	LT:
T GHOT	7							Regular Cost			Vendor #:			
								Invoice Cost (WAC) (\$)	\$16.3	2 Whsl. Code			
									mimicooc		Fineline Co	ode:		
								As of date:	7/7/2020					
			Attach copy of SAFFTY DAT	A SHEET (SDS)	or non haza	rd letter, PACKAGE INSE	RT, LABEL AND PHOTO OF	PRODUCT PACK	AGING and BARCODI		ı			
*Please provide any additional in	formation on page					See new p. 3 for Desig			Signature:					



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For Designated Drop Ship Only Products, Please Use Page 3 MATERIAL HAZARD CLASSIFICATION and TRANSPORTATION Is this product (check all that apply): a. Cytotoxic? SDS Hazard Classification No 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? No Oxidizer Inorganic Does the product label bear a CA Prop 65 warning? No Steroid/Androgen Contact Hazard Yes Aerosol Class; Identify NFPA Storage Level: c. Contact Hazard? d. Does this product require special clean-up instructions? No Is the product a NIOSH hazardous drug? (If yes, attach SDS with special instructions.) No e. Does the product contain DEHP? No If yes, indicate which: Is this product regulated for shipment by DOT? No (if yes, answer a-e below and provide SDS) **Hazardous Waste Identificatio** a. UN/Identification Number b. Proper Shipping Name EPA Hazardous Waste Code: Waste Characteristics c. DOT Hazard Class d. Packing Group e. Inhalation Hazard? Yes Is this product regulated for shipment by IATA? REMS or REGISTRY RESTRICTIONS (if yes, answer a-e below and provide SDS) Is there a REMS on this product? No a. UN/Identification Number If Yes, is it managed with a pharmacy registry? Website URL: b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard? Med Guide Required Is the product restricted for air shipment? If so, indicate restriction: Limited Distribution Requirement x Passenger Comments / Details: (For example, iPledge program?) Cargo Passenger & Cargo Is this a reportable quantity? REMS: RQ Threshold: REMS Program Manager Name: Phone Is this a marine pollutant? Supplier Manages REMS registry exclusively: Is this product shipped utilizing an authorized DOT exception or Special Permit? Wholesale distributor support: No (if yes, identify method below) Provider Name: DEA #: Limited Quantity Site Enrollment Number assigned PCPDP# Consumer Commodity, ORM-D NPI#: by Supplier: Small Quantity (49 CFR 173.4) Special Permit: DOT-SP Comments Special Provision (listed in Column 7 of 49 CFR 172.101); SP# Registry: Phone: Registry Program Contact Name: ADD'L STORAGE INFORMATION Comments Is the Product... RETURN INSTRUCTIONS Controlled Substance? No Controlled Substance Code Controlled by State(s)? No Listed Chemical (List I or II) No 866-525-7270 ARCOS Reportable? No If yes, indicate which: Contact tel. # if product received damaged: 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 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: Restricted from US territories? (explain in comments) No If so, which states? Other requirements? Comments? Comments: MISCELLANEOUS NOTES and/or Image of Product Barcode



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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 c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity: 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:						
Supplier's Customer Service Number: Contracted 3PL company / contact #: Phone: Expedited Freight Charges or Other Designated Drop Ship Fees:	Ships regular ground for 3-10 days receipt: 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						
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: 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 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?						
Miscellaneous Notes:							
	ADDITIONAL INFORMATION						
	Is product order for scheduled patient procedure? Is product order for restocking purposes?						