

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

Version 2021					Introduction Type:	New Item		Final Version			Date:	6/23/	2023
		PRODUCT INFOR	MATION					SPECIAL HAN	IDLING AND STOP	RAGE REQUI	REMENTS*		
Company Name: SOLA Pharmaceuticals Application: ANDA							a. Temperature – Indicate the USP temperature range for this product.						
Application Number for NDA/AN		)(med device):	2104	36				perature Range	Controlled Room		and 25 C (68	8° – 77° F)	
Medical Device Class, if applicat		· · · · ·						<u> </u>					
DUNS:	080121345						Othe	r Temperature Range	Requirement				
Proprietary Name (If Applicable) a	nd Established Name:	Dimethyl Fumarate 120mg						(write in)					
Selling Unit NDC:	70512-852-14	Unit of Use N	DC:			120852149	Note	6					
UDI		CVX Code:			MVX Code:								
Description:	Dimethyl Fumarate 120mg,	, 14ct; Capsule, delayed release; Or	al					s product to be shippe				No	
							Is thi	s product to be shippe	d to customers on o	dry ice?		No	
Active Ingredient(s): Dimethyl Fumarate													
								erature excursion qu	estions:				
URL for Additional Product Inform Address:	655 Highlandia Dr.				Address 2:		Nam Num			866-747-736	5		
City:	Baton Rouge			State:		<b>p:</b> 70810		p E-mail:		info@sola			
Key Contact:	Baton ribago			Email:	info@solameds.us			p =		moesona	incus.us		
Phone Number:	866-747-7365			Fax:	800-754-9550		c. Special regulation	ns for product in any	states?			No	
Product Therapeutic Classification	n: Immuno	omodulator Agent					Spec	ial returns requiremen	ts for this product?			No	
		-											
	ADDITIONAL PR	RODUCT INFORMATION			PRODUCT DESC	CRIPTION INFORMATION	d. Store product (u	nit of sale) upright?				No	
The product is?		Is the Product	Direct-Ship Onl	ly			Prot	ect product (unit of s	ale) from light?			Yes	
a legend device?	No	Is the Product	Unit of Use		Size:	14ct	e. Shelf life:					48	Months
if yes, enter class #		Orphan Drug Statu	5		0120.		Initia	I shelf life at launch	if different):				Months
a product kit?	No				Strength:	120mg							
if yes, list NDCs of component parts		FDA Approval State	IS		•	Canaula, delayed valesses				MATION			
reverse numbered?	No				Dosage Form:	Capsule, delayed release; Oral	Unit	of Sale		What is the	NDC selling	unit?	
co-licensed?	No	Allergens Present				olui				1 bottle of 1	-	unit:	
latex-free?	Yes					Capsule		Box/Carton			g. 1 Box of 1	0 Vials)	
preservative-free?	Yes				Product Shape:			Ampule		· · ·	5	- /	
correctional institution block?	No				Product Color:	Green cap, white body		Glass		Minimum o	rder quantity	?	Yes
opioid?	No				riodder obior.			Tube					
Cannabinoid?	No	Country of Origin	Greece		Product Imprint:	"120" in black ink		Vial Liquid Sgl					_
If Unit Dose, is item bar coded to u hospital scanning?	unit dose for No	In this product environ	ad under the					Vial Liquid Multi Vial Powder Sol		If Yes, how	Each	ch package f	ype?
If Unit Dose, indicate NDC here:	NO	Is this product cover Trade Agreements A		/es				Vial Powder Sqi Vial Power Multi		24	Each Inner/Cartor	Pack	
in onit bose, indicate NDC here.		Trade Agreements /		63				Other: Write In			Case	I/I dok	
		FOR GENERIC DRUG	PRODUCTS										
							-						
				Au	thorized Generic *If A	Authorized Generic, other		Pł	ARMACY ORDER	R / BILL UNIT			
I. Orange Book Rating:	AB				sec	tion fields are not applicable	Rec. sell unit to cus	stomer?		Rx billing u	nit to pharm	acy:	
II. Generic Equivalent to What Brand?: Tecfidera®			1 bottle				x Each						
-							(Write-in, e.g. 1 Via	)	_		Gram		
	DF	RUG SUPPLY CHAIN SECURITY A	CT (DSCSA) INFORM	IATION							Milliliter		
Doos supplier most DSCSA defini	tion of monufacturor?	Yes		21.11.	0370512000004			ITEN	I AND PACKING I		N		
Does supplier meet DSCSA defini Is product exempt from DSCSA?		No		GLN:	0370312000004			1160	AND FACKING I		•		
If yes, select exemption:				GCP:	0370512		1		Dimensi	ions (US msn	nte )	Volume	Saleable #
If yes, select exemption: Other exemption - Write in:				JUP:	0370512		1	Weight Lbs.	Depth	Width	Height	(Cube)	Saleable # Pieces
Is product repackaged?		No	14	f ves, was o	riginal product purchas	ed	Item/Each:		1				
Is product sold by manufacturer's	exclusive distributor?	No		lirect from n			10111240111	0.05	1.9685	1.9685	3.54331	13.730299	1
Has FDA granted waiver/exception		No			ce manufacturer for rep	ackaged product	Box/Carton/Bundle	1					
If yes, attach documentation from	m FDA.						Inner Pack:						
							Case:	1.2	12.5984	5.51181	8.66142	601.44889	24
		GTIN AND HIBCC PRODU	TINFORMATION				<b>D</b> -11-4						
Saleable Unit of Measure	Saleable G	Quantity HIBCC		CT.	N-14	Unit of Use GTIN-14	Pallet:	493	39.3701	39.3701	47.2441	73228.581	1440
x Item/Each	Saleable G				70512852146	00370512852146							
Box/Carton/Bundle/Inner Pack							C	OST INFORMATION			WHOLESAL	ER USE ONL	Y:
X Case	24			503	70512852141								
Pallet							Regular Cost			Vendor #:			
							Invoice Cost (WAC)	(\$)	\$75.00	Whsl. Code			
							A second states	5/1/2023		Fineline Co	de:		
							As of date:	5/1/2023		-			
		Attach copy of SAFET	DATA SHEET (SDS)	) or non haza	ard letter. PACKAGE INS	ERT, LABEL AND PHOTO OF F	PRODUCT PACKAGING	and BARCODE					
*Please provide any additional inf	ormation on page 2.	, and topy of OATET		, 5		ignated Drop Ship Only.		ature:					
						5 ···· -··· -··· -··· /·	0.9.	· · · · · ·					

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

Version 2021 Fo	or Designat	ed Drop Ship Only Products, Please Use Page 3			
MAT	ERIAL HAZ	ZARD 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)		DS Hazard Classification Corrosive Oxidizer Contact Hazard No No No			
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?	No	Hazardous Waste Code:	Waste Characteristics		
(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?	No	REMS of Is there a REMS on this product? If Yes, is it managed with a pharmacy registry? Website URL:	No No		
Is the product restricted for air shipment? If so, indicate restriction:           Passenger           Cargo           Passenger & Cargo	No	Med Guide Required Limited Distribution Requirement Comments / Details: (For example, iPledge program?)	No No		
Is this a reportable quantity? No RQ Threshold: 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) Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101);	REMS: REMS Program Manager Name: Supplier Manages REMS registry exclusively: Wholesale distributor support: Provider Name: Site Enrollment Number assigned by Supplier: Comments	No	Phone:		
SP#		Registry:	No		
ADD'L STORAGE INFORMATION		Registry Program Contact Name: Comments		Phone:	
Controlled Substance? No Controlled Substance Code		F	ETURN INSTRUCTIONS		
Controlled by State(s)? No Listed Chemical (List I or II) ARCOS Reportable? No If yes, indicate which: Schedule No. Is it a scheduled listed chemical product?: CLASS OF TRADE RESTRICTION:	No No	Contact tel. # if product received damaged: Is product returnable for credit: URL/Link to returns policy:	866-747-7365 Yes		
	Yes	URL/LINK to returns policy:			
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:	Special regulations or returns requirements for this product in certain states? If so, which states? Other requirements? Comments?	No			
MIS	CELLANE	OUS NOTES and/or Image of Product Barcode:			



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

Version 2021	FOR DESIGNATED DROP SHIP PRODUCT ONLY - in	f not a designated drop ship, do not complete.				
Order Method	l 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 #:	Fax Number:         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 Ch	arges or Other Designated Drop Ship Fees:	Overnight and Priority Overnight PO Processing				
Expedited freight fees billed with each ord Drop Ship service fee billed with each ord Drop Ship miscellaneous fees billed: Comments:		Overnight receipt available:       Image: Comparison of the second				
CI	ass of Trade Restriction:	PO Receipt Cut off time:				
	oharmacy, hospitals, clinics and physician offices	Saturday Overnight receipt available:       PO Receipt Cut off time:         Order receipt method:       Phone:         Fax:       EDI:         Overnight Fees apply:       Other fees apply:				
Other Data Ir	nformation 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?				