

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

Version 2021						Introduction Typ	e: Open Stock		Final Version			Date:	7/13/	2023
			PRODUCT INFORMA	TION					SPECIAL HAN	IDLING AND STO	RAGE REQUI	REMENTS*		
Company Name:	Xiromed LLC					Application	1: ANDA	a. Temperature – In	dicate the USP temp	erature range for	this product.			
Application Number for NDA/AN		/510(k)(med devic	e):	090	114				perature Range	Controlled Room	- between 20	and 25 C (68	° – 77° F)	
Medical Device Class, if applicat			,						Ŭ					
DUNS:	080228637							Othe	r Temperature Range	Requirement				
Proprietary Name (If Applicable) a		e: Syeda							(write in)					
Selling Unit NDC:	70700-115-85		Unit of Use NDC				70700115855	Note	s					
UDI			CVX Code:			MVX Code:								
Description:	Drospirenone + EE,	3mg-0.03mg - 3 x 2	8 Pack					Is thi	s product to be shippe	d to customers on	ice?		No	
								Is thi	s product to be shippe	d to customers on	dry ice?		No	
Active Ingredient(s):	C	Prospirenone And E	thinyl Estradiol											
UDI for Additional Draduat Inform								b. Contact for temp Nam	erature excursion qu	estions:				
URL for Additional Product Inform Address:	180 Park Ave					Address 2: S	uite 101	Num						
City:	Florham Park				State:		Zip: 07932		ip E-mail:					
Key Contact:	David Hernandez				Email:	david.hernandez								
Phone Number:	973-324-0200				Fax:	862-286-0932		c. Special regulatio	ons for product in any	states?			No	
Product Therapeutic Classification	n:							Spec	ial returns requiremen	ts for this product?	,		No	
	ADDITION	IAL PRODUCT INF	ORMATION			PRODUCT DE	SCRIPTION INFORMATION	d. Store product (u	nit of sale) upright?				Yes	
The product is?			Is the Product	Direct-Ship O	nly				ect product (unit of s	ale) from light?			Yes	
a legend device?	N	lo	Is the Product	Neither		Size:	3 x 28 Pack	e. Shelf life:					24	Months
if yes, enter class #			Orphan Drug Status					Initia	I shelf life at launch	if different):				Months
a product kit?						Strength:	3mg/0.03mg			ORDER INFOR	ΜΑΤΙΟΝ			
if yes, list NDCs of component parts			FDA Approval Status				Tablets			ORDER IN OR	MATION			
reverse numbered?	N	lo				Dosage Form:	Tubleta	Unit	of Sale		What is the	NDC selling	unit?	
co-licensed?		lo	Allergens Present						Bottle		1 Box of 84,	3 blisters of	28	
latex-free?			_			Product Shape	Round	X	Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free?						i roudet onape.			Ampule					
correctional institution block?						Product Color:	21 yellow active; 7 white in		Glass		Minimum or	der quantity	?	Yes
opioid?	_			On alla					Tube					
Cannabinoid? If Unit Dose, is item bar coded to u	unit doop for		Country of Origin	Spain		Product Imprint	t: 21 active debossed SZ and		Vial Liquid Sgl Vial Liquid Multi		If Yoo how	many of whi	ch package t	huno2
hospital scanning?	Init dose for		Is this product covered	under the					Vial Powder Sql			Each	ch package i	type?
If Unit Dose, indicate NDC here:	-		Trade Agreements Act (Yes				Vial Power Multi			Inner/Carton	/Pack	
									Other: Write In			Case		
			FOR GENERIC DRUG PR	RODUCTS										
					Au		f Authorized Generic, other			ARMACY ORDER				
I. Orange Book Rating:	AB					St	ection fields are not applicable	Rec. sell unit to cus		-	Rx billing u		acy:	
II. Generic Equivalent to What Brand?: Yasmin					1 Pack 3X28			X	x Each Gram					
		DRUG SUPPL	Y CHAIN SECURITY ACT	(DSCSA) INFOR	MATION			(Write-in, e.g. 1 Vial	1)			Milliliter		
				()				-				winniter		
Does supplier meet DSCSA definit	tion of manufacturer	?	Yes		GLN:	037070000007			ITE	AND PACKING	INFORMATIO	N		
Is product exempt from DSCSA?			No											
If yes, select exemption:					GCP:				Weight Lbs.		ions (US msn		Volume	Saleable #
Other exemption - Write in:									troigin LDS.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?		-	No			riginal product purcha	ised	Item/Each:	0.08	4.0	2.0	1.0	8.0	1
Is product sold by manufacturer's Has FDA granted waiver/exception			No		direct from n	rce manufacturer for r	anackaged product	Box/Carton/Bundle	1					
If yes, attach documentation from					FIOVILE SOUL		epackaged product	Inner Pack:					0	
								Case:	6.9	10.7	8.4	8.5	763.98	80
		GTIN	AND HIBCC PRODUCT	INFORMATION					0.9	10.7	0.4	0.0	/03.90	00
								Pallet:					0	
Saleable Unit of Measure	Sale	eable Quantity	HIBCC			IN-14 370700115855	Unit of Use GTIN-14							
X Item/Each Box/Carton/Bundle/Inner Pack	_	1			003	010100110055			OST INFORMATION				ER USE ONL	v
X Case	-	80			203	370700115859						MOLLOAL		
Pallet	-	00			200			Regular Cost			Vendor #:			
								Invoice Cost (WAC)) (\$)		Whsl. Code	#:		
											Fineline Co	de:		
								As of date:			_			
μ					<u></u>						1			
*Diseas manida and different a	-		Attach copy of SAFETY D	A I A SHEET (SD	S) or non haza		SERT, LABEL AND PHOTO OF							
*Please provide any additional inf	ormation on page 2.					See new p. 3 for De	signated Drop Ship Only.	Sign	ature:					

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

Version 2021 For Design	ated Drop Ship Only Products, Please Use Page 3					
MATERIAL HA	AZARD 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? Is the product a CA Prop 65 reproductive toxicant? Does the product label bear a CA Prop 65 warning? No	SDS Hazard Classification Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard					
c. Contact Hazard? No d. Does this product require special clean-up instructions? No (If yes, attach SDS with special instructions.) e. Does the product contain DEHP? No Is this product regulated for shipment by DOT? No	Does the product have an Aerosol class? If yes, identify NFPA Storage Level: NFPA Storage Level: Is the product a NIOSH hazardous drug? Yes					
(if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group	If yes, indicate which: Group 2 items (non-antineoplastic that meets a hazard criterion) Hazardous Waste Identification					
e. Inhalation Hazard? No	EPA Hazardous Waste Code: Waste Characteristics					
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 d. Packing Group	REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? No If Yes, is it managed with a pharmacy registry? Website URL:					
e. Inhalation Hazard? No Is the product restricted for air shipment? If so, indicate restriction: No Passenger Cargo Passenger & Cargo	Med Guide Required Limited Distribution Requirement Comments / Details: (For example, iPledge program?)					
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 Perovision (listed in Column 7 of 49 CFR 172.101);	REMS: Phone: REMS Program Manager Name: Phone: Supplier Manages REMS registry exclusively: Phone: Wholesale distributor support: Provider Name: Site Enrollment Number assigned DEA #: by Supplier: NPI #:					
Special Provision (listed in Column 7 of 49 CFR 172, 101); SP#	Registry:					
ADD'L STORAGE INFORMATION Is the Product	Registry Program Contact Name: Phone: Phone: Comments					
Controlled Substance? No Controlled Substance Code 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?: No	RETURN INSTRUCTIONS Contact tel. # if product received damaged: 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:	Special regulations or returns requirements for this product in certain states? If so, which states? Other requirements? Comments?					
MISCELLAN	EOUS NOTES and/or Image of Product Barcode:					



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Version 2021 FOR DESIGNATED DROP SHIP PRODUCT ONLY - i	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: c. Fax Fax Number: d. Phone only Phone No.: e. Supplier Web Site only Site Address: Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Name: Phone:	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 Charges or Other Designated Drop Ship Fees:	Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each order: Drop Ship service fee billed with each order: Drop Ship miscellaneous fees billed: Comments:	Overnight receipt available: Image: Comparison of the second
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: Fax #: EDI: Other fees apply:
Other Data Information Required to Process PO:	Return Instructions
Patient Procedure Date:	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?