

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

Version 2020						Introduction Type:	Post Launch Change]		Final Version			Date:	10/8/	2021
			PRODUCT INFORMAT	ION						SPECIAL HAN	DLING AND STOR	RAGE REQUI	REMENTS*		
Company Name: Xiromed LLC Application: ANDA						a. Temperature – Indicate the USP temperature range for this product.									
Application Number for NDA/AN					0124			Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)							
DUNS:	080228637							_	Other Ten	nperature Range R	equirement				
Proprietary Name (If Applicable) a		ne: FLUOR	OURACIL					1		e in)	- 1				
Selling Unit NDC:	70700-188-22		Unit of Use NDC:			UPC: 37070	0188224	11	Notes						
UDI			CVX Code:			MVX Code:									
Description:	Fluorouracil Injection	on, USP 2.5g/50mL	1x50mL Multiple-dose vial					1	Is this pro	duct to be shipped	to customers on ic	ce?		No	
-		-	•							duct to be shipped				No	•
Active Ingredient(s): FLUOROURACIL						11									
								b. Contact fo		ure excursion que	stions:				
URL for Additional Product Inform						A 1 1 0		41	Name:			Steven Yeur			
Address:	180 Park Ave Florham Park				State:	Address 2: Suite NJ Zip:			Number:			844-947-663			
City: Key Contact:	David Hernandez				Email:	david.hernandez@x		-11	Group E-	maii:		steven.yeu	ing@xirom	ea.com	
Phone Number:	844-947-6633				Fax:	862-286-0932	aromea.com	c Special red	ulations fo	or product in any	states?			No	
Product Therapeutic Classificatio						002 200 0002				turns requirements				No	
Troduct merapeano olassineano	,,,,								Opeciai ic	itamo requiremento	rior and product:				
	ADDITIO	NAL PRODUCT INF	FORMATION			PRODUCT DESC	RIPTION INFORMATION	d. Store prod	uct (unit o	f sale) upright?					
The product is?			Is the Product	Direct-Ship On	ılv					roduct (unit of sal	e) from light?			Yes	
a legend device?		No	Is the Product	Neither	y		1 x 50mL multiple-dose	e. Shelf life:	i ioteot p	roduct (dilit or sal	c) iroin iigiit.			24	Months
if yes, enter class #		140	Orphan Drug Status	11010101		Size:	vial	0.0.0	Initial she	elf life at launch (if	different):				Months
a product kit?		No				Strength:	2.5GM/50ML (50MG/ML)			•	,				
if yes, list NDCs of			FDA Approval Status			Strength:					ORDER INFOR	MATION			
component parts						Dosage Form:	INJECTABLE								
reverse numbered?		No				2 coago : c			Unit of Sa			What is the		unit?	
co-licensed?		No	Allergens Present							Bottle		1 Box contai			
latex-free? preservative-free?		Yes	Container closure is no rubber la		itural	Product Shape:				Box/Carton		(Write-in, e.	g. 1 Box of 10) Vials)	
correctional institution block?		Yes	rubber ia	atex.						Ampule Glass		Minimum or	dor augntitu	2	Yes
opioid?		No				Product Color:				Tube		William Or	uer quaritity	٠.	162
Cannabinoid?		No	Country of Origin	India						Vial Liquid Sgl					
If Unit Dose, is item bar coded to u			,g			Product Imprint:				Vial Liquid Multi		If Yes, how	many of which	ch package ty	ype?
hospital scanning?			Is this product covered un	der the						Vial Powder Sql			Each		
If Unit Dose, indicate NDC here:			Trade Agreements Act (TA	λA)?	No					Vial Power Multi			Inner/Carton	/Pack	
								<u>J</u>		Other: Write In			Case		
			FOR GENERIC DRUG PRO	DDUCTS											
					Autho		thorized Generic, other section	PHARMACY ORDER / BILL UNIT							
I. Orange Book Rating:					fields are not applicable			Rec. sell unit to customer?			Rx billing unit to pharmacy:				
II. Generic Equivalent to What Bra	and?:								ox containir	ng 1 Vial		x	Each		
		DRIIC SUBBI	Y CHAIN SECURITY ACT (I	DECEAL INFOR	MATION			(Write-in, e.g	. 1 Vial)				Gram Milliliter		
		DRUG SUFFL	T CHAIN SECURITT ACT (L	JSCSA) INFOR	WATION								Milliliter		
Does supplier meet DSCSA defini	ition of manufacture	er?	Yes	GLN	l:	0370700000007				ITEM	AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			No	-											
If yes, select exemption:				-				-			Dimonsi	ions (US msm	ite \	Volume	
Other exemption - Write in:										Weight Lbs.	Depth	Width	Height	(Cube)	# Pieces:
Is product repackaged?			No	If Ye	s. was origin	nal product purchased		Item/Each:		0.040					
Is product sold by manufacturer's	s exclusive distribut	or?	No		ct from mfr?					0.243	1.969	1.969	3.15	12.212427	1
Has FDA granted waiver/exceptio	n/exemption for pro	duct?	No	If ye	s, attach doc	umentation from FDA.		Box/Carton/E	Bundle/					0	
								Inner Pack:						Ü	
		GTI	N AND HIBCC PRODUCT IN	IFORMATION				Case:		11.464	10.433	8.465	6.89	608.49273	40
Onlankin Hair of Manager								11		-					
Saleable Unit of Measure	1	Quantity	HIBCC		GTIN-	14 700188224	Unit of Use GTIN-14	Pallet:						0	
x Item/Each Box/Carton/Bundle/Inner Pack		1			00370	700100224									
x Case		40			20370	700188228			COST	INFORMATION			WHOLESAL	ER USE ONL'	Y:
Pallet		40			20070										
	7							Regular Cost	:	ſ		Vendor #:			
								Invoice Cost	(WAC) (\$)		\$29.70	Whsl. Code			
								11	-			Fineline Cod	de:		
								As of date:	L						
<u> </u>				au=== :	a) ,		EDT ABEL AND BUG	II				1			
			Attach copy of SAFETY DA	IA SHEET (SD	5) or non haz	ard letter, PACKAGE INS	ERT, LABEL AND PHOTO OF	PRODUCT PACK	AGING and	BARCODE.					



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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								
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?	Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard								
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?	No No Aerosol Class; Identify NFPA Storage Level: Is the product a NIOSH hazardous drug? If yes, indicate which: Yes Group 1 items (antineoplastic)								
(if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class	Hazardous Waste Identification EPA Hazardous Waste Code: Waste Characteristics								
d. Packing Group e. Inhalation Hazard?	No DELICATION DE CARRIENTO NO								
Is this product regulated for shipment by IATA? (if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class	No REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? No If Yes, is it managed with a pharmacy registry? Website URL:								
d. Packing Group e. Inhalation Hazard? Is the product restricted for air shipment? If so, indicate restriction: Passenger Cargo	Med Guide Required Limited Distribution Requirement Comments / Details: (For example, iPledge program?)								
Passenger & Cargo Is this a reportable quantity? Yes RQ Threshold: 500 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)	REMS: REMS Program Manager Name: Supplier Manages REMS registry exclusively: Wholesale distributor support: Provider Name: Site Enrollment Number assigned by Supplier: PROVIDENTIAL PROPERS TO BE A #: PCPDP#: NPI #:								
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101); SP#	Comments Registry:								
ADD'L STORAGE INFORMATION Is the Product	Registry Program Contact Name: Phone: Comments								
Controlled Substance? Controlled by State(s)? ARCOS Reportable? Schedule No. No Listed Chemical (List I or II) If yes, indicate which: Is it a scheduled listed chemical product?: CLASS OF TRADE RESTRICTION:	No No No Understand the state of the state o								
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?								
	_LANEOUS 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	r Designated Drop Ship Product	Standard Order Receipt and Processing						
Purchase orders may be accepted by:		Purchase order daily receipt cut off time by supplier						
a. EDI		Cut off time:	311					
b. Autofax	Fax Number:							
c. Fax	Fax Number:	Shipping lead time of PO: Hours	Days					
d. Phone only	Phone No.:							
e. Supplier Web Site only	Site Address:	Ships same day for next day receipt:						
Minimum Order Quantity: Supplier's Customer Service Number:		Ships for second day receipt: Ships regular ground for 3-10 days receipt:						
	lame:	Ships regular ground for 3-10 days receipt.						
	Phone:							
Expedited Freight Charge	es or Other Designated Drop Ship Fees:	Overnight and Priority Overnight PO Pro	cessing					
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:		Days of week overnight is available:	Monday					
Comments:		Days of week overliight is available.	Tuesday					
Gommonio.			Wednesday					
			Thursday					
			Friday					
		Priority Overnight receipt available:						
Class	of Trade Restriction:	PO Receipt Cut off time:						
No restriction: Select YES if sold to retail phar		Saturday Overnight receipt available:						
Restricted to retail pharmacy only:	mady, neophale, climed and physician chiese	PO Receipt Cut off time:						
Restricted to hospital, clinics, and physician of	ffices only:	Order receipt method: Phone: Phone #:						
Restricted from US territories? (explain in com		Fax: Fax #:						
Comments:		EDI:	•					
		Overnight Fees apply:						
		Other fees apply:						
Other Data Infor	mation Required to Process PO:	Return Instructions						
Patient Procedure Date:		Contact # if product is received damaged:						
Physician Name:		Is product returnable for credit:						
Physician/Clinic Phone #		URL/Link to returns policy:						
Physician State License # Physician/Clinic DEA #:		Special regulations or returns requirements for this product in certa	ain states?					
Physician/Clinic Specialty:		If so, which states? Other requirements? Comments?	iii states:					
_ · · · · · · · · · · · · · · · · · · ·	scellaneous Notes:	·						
		ADDITIONAL INFORMATION						
		Is product order for scheduled patient procedure?						
		Is product order for restocking purposes?						