

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

Version 2020				Introduction Type:	Post Launch Change		Final Version			Date:	7/10/	2024
		PRODUCT INFORMATION					SPECIAL HAN	NDLING AND STO	RAGE REQUI	REMENTS*		
Company Name:	Xiromed LLC			Application:	ANDA	a. Temperature - Indicate the USP temperature range for this product. Temperature Range Controlled Room - between 20 and 25 C (68° - 77° F)						
Application Number for NDA/AN	DA/BLA (drug); PMA/510(k)(med dev	ice): 210	123		•							
DUNS:	080228637						emperature Range I	Requirement				
Proprietary Name (If Applicable) a Selling Unit NDC:	Ind Established Name: FLUC 70700-186-23	ROURACIL Unit of Use NDC:	1	UPC: 37070	0186237	(v Notes	rite in)					
	10100-180-23	CVX Code:		MVX Code:	10106237	Notes						
Description:	Fluorouracil Injection, USP 500mg/10					Is this r	roduct to be shipped	d to customers on i	ce?		No	
Description.		-					roduct to be shipped				No	•
Active Ingredient(s):	FLUOROURACIL											
URL for Additional Product Inform	action:					b. Contact for temper Name:	ature excursion qu	estions:	Xiromed Qua	ality		
Address:	180 Park Ave		1	Address 2: Suite	101	Numbe	r:		844-947-663			
City:	Florham Park		State:	NJ Zip:		Group	E-mail:		US-Quality	-Xiromed@	xiromed.co	<u>om</u>
Key Contact:	Xiromed Regulatory		Email: Fax:	usregulatory@xiron	ned.com						Ν.	
Phone Number: Product Therapeutic Classificatio	844-947-6633		Fax:	862-286-0932		c. Special regulations	returns requirement				No No	
Froduct merapeutic classificatio	n.					Special	returns requirement	ts for this product?			NU	
	ADDITIONAL PRODUCT IN	FORMATION		PRODUCT DESC	RIPTION INFORMATION	d. Store product (unit	of sale) upright?					
The product is?		Is the Product Direct-Ship Or	nly			Protect	product (unit of sa	ale) from light?			Yes	
a legend device?	No	Is the Product Neither		Size:	10 x 10mL Single-dose	e. Shelf life:					24	Months
if yes, enter class # a product kit?	No	Orphan Drug Status			vials 500MG/10ML (50MG/ML)	Initial s	helf life at launch ((if different):				Months
if yes, list NDCs of	NO	FDA Approval Status		Strength:	COUNCY TOWNE (COUNCYME)			ORDER INFOR	MATION			
component parts				Dosage Form:	INJECTABLE							
reverse numbered? co-licensed?	No No	Allergens Present				Unit of	Sale Bottle		What is the 1 Box contai		unit?	
latex-free?	Yes	Container closure is not made with na	atural			x	Box/Carton			q. 1 Box of 1) Vials)	
preservative-free?	Yes	rubber latex.		Product Shape:			Ampule			-	,	
correctional institution block?	Yes			Product Color:			Glass		Minimum or	der quantity	?	Yes
opioid? Cannabinoid?	No No	Country of Origin India					Tube Vial Liquid Sgl					
If Unit Dose, is item bar coded to u				Product Imprint:			Vial Liquid Multi		If Yes, how	many of whi	ch package ty	ype?
hospital scanning?		Is this product covered under the					Vial Powder Sql		24	Each		
If Unit Dose, indicate NDC here:		Trade Agreements Act (TAA)?	No				Vial Power Multi Other: Write In			Inner/Carton Case	/Pack	
		FOR GENERIC DRUG PRODUCTS		+						ouse		
			Auth		thorized Generic, other section			HARMACY ORDER				
I. Orange Book Rating:	AP		fields are not applicable			Rec. sell unit to custo		-	Rx billing ur		acy:	
II. Generic Equivalent to What Bra	nd?:					1 Box contain (Write-in, e.g. 1 Vial)	ing 10 Vials		x	Each Gram		
-	DRUG SUPF	PLY CHAIN SECURITY ACT (DSCSA) INFOR	MATION			(vinio ini, o.g. i viai)				Milliliter		
			_					M AND PACKING				
Does supplier meet DSCSA defini Is product exempt from DSCSA?	tion of manufacturer?	Yes GLM	N:	037070000007			II EI	M AND PACKING	INFORMATION	N		
If yes, select exemption:								Dimens	ions (US msm	uts)	Volume	
Other exemption - Write in:							Weight Lbs.	Depth	Width	Height	(Cube)	# Pieces:
Is product repackaged?				nal product purchased		Item/Each:	0.617	5.197	2.126	2.559	28.273935	1
Is product sold by manufacturer's Has FDA granted waiver/exception			ct from mfr?			Box/Carton/Bundle/			-			
Has FDA granted waiver/exception	n/exemption for product?		es, attach dot	cumentation from FDA.		Inner Pack:					0	
	G	TIN AND HIBCC PRODUCT INFORMATION				Case:	16.535	11.024	9.134	8.465	852.36807	24
	0		0711				10.000	11.024	5.104	0.400	002.00001	24
Saleable Unit of Measure	Quantity 1	HIBCC	GTIN- 00370	14 0700186237	Unit of Use GTIN-14	Pallet:					0	
Box/Carton/Bundle/Inner Pack			00010									
X Case	24		20370	0700186231		CO	ST INFORMATION			WHOLESAL	ER USE ONL	Y:
Pallet	4					Regular Cost		-	Vendor #:			
	┥ ┣━━━┥					Invoice Cost (WAC) (5)	\$60.00	Whsl. Code	#:		
							-		Fineline Cod			
						As of date:						
<u> </u>		Attach copy of SAFETY DATA SHEET (SE	S) or non bor						1			
*Please provide any additional inf	formation on page 2.	Auach copy of SAFETT DATA SHEET (SL	or non haz	See new p. 3 for Desig		PRODUCT PACKAGING a						
					,	orginati						



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Version 2020 For Designated Drop Ship Only Products, Please Use Page 3								
MATERIAL 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? 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	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 d. Packing Group	Hazardous Waste Identification EPA Hazardous Waste Code: Waste Characteristics							
e. Inhalation Hazard? No								
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: Passenger Cargo Passenger & Cargo	Med Guide Required No Limited Distribution Requirement No Comments / Details: (For example, iPledge program?) No							
Is this a reportable quantity? Yes RQ Threshold: 500 Is this a marine pollutant? <u>No</u> 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: Supplier Manages REMS registry exclusively: Phone: Wholesale distributor support: Provider Name: Provider Name: DEA #: Site Enrollment Number assigned PCPDP#: by Supplier: NPI #:							
SP#	Registry:							
ADD'L STORAGE INFORMATION	Registry Program Contact Name: Phone: Comments							
Is the Product Controlled Substance? No Controlled Substance Code	RETURN INSTRUCTIONS							
No Listed Chemical (List I or II) No ARCOS Reportable? No If yes, indicate which: If yes, indicate which: Schedule No. Is it a scheduled listed chemical product?: No CLASS OF TRADE RESTRICTION:	Contact tel. # if product received damaged: Is product returnable for credit: 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?							
Comments:								
	DUS NOTES and/or Image of Product Barcode:							



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Version 2020	FOR DESIGNATED DROP SHIP PRODUCT ONLY - i	f not a designated drop ship, do not complete.				
Order Meth	od 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	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: Ships same day for next day receipt:	Days			
Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Expedited Freight (Name: Phone: Charges or Other Designated Drop Ship Fees:	Ships for second day receipt: Ships regular ground for 3-10 days receipt: Overnight and Priority Overnight PO Processing				
Expedited freight fees billed with each o	rder:	Overnight receipt available:				
Drop Ship service fee billed with each o	rder:	PO Receipt cut off time:	_			
Drop Ship miscellaneous fees billed: Comments:			londay luesday Vednesday hursday iriday			
		Priority Overnight receipt available:				
No restriction: Select YES if sold to retain Restricted to retail pharmacy only: Restricted to hospital, clinics, and physic Restricted from US territories? (explain Comments:	in comments)	PO Receipt Cut off time: Image: Saturday Overnight receipt available: PO Receipt Cut off time: PO Receipt Cut off time: Order receipt method: Phone: Fax: Fax #: EDI: Image: Saturday Overnight Fees apply: Other fees apply: Image: Saturday Overnight 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?				