

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

Version 2020						Introduction Type:	Post Launch Change	]	Final Version			Date:	10/8/	2021
			PRODUCT INFORMA	TION					SPECIAL HA	NDLING 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	NDA/BLA (drug); PN	IA/510(k)(med devi	ice):	210	124			]	Temperature Range	Controlled Room	- between 20	and 25 C (68°	– 77° F)	
DUNS:	080228637							_	Other Temperature Range	Requirement				
Proprietary Name (If Applicable)		me: FLUOI	ROURACIL						(write in)					
Selling Unit NDC:	70700-189-22		Unit of Use NDC:				0189221		Notes					
UDI			CVX Code:			MVX Code:		!						
Description:	Fluorouracil Inject	ion, USP 5g/100mL	1x100mL Multiple-dose vial						Is this product to be shippe				No No	
Active Ingredient(s):		FLUOROURACIL						1	Is this product to be shippe	a to customers on a	ry ice?		INO	
								b. Contact for	temperature excursion qu	estions:				
URL for Additional Product Inform									Name:		Steven Yeun			
Address:	180 Park Ave				State:	Address 2: Suite		<b>!</b>	Number:		844-947-663			
City: Key Contact:	Florham Park David Hernandez				Email:	david.hernandez@x	07932	1	Group E-mail:		steven.yeu	ng@xirome	ea.com_	
Phone Number:	844-947-6633				Fax:	862-286-0932	iromeu.com	c. Special reg	ulations for product in any	states?			No	
Product Therapeutic Classification								]	Special returns requiremen				No	
									.,					
	ADDITIC	NAL PRODUCT IN	IFORMATION			PRODUCT DESC	RIPTION INFORMATION	d. Store produ	ct (unit of sale) upright?					
The product is?			Is the Product	Direct-Ship O	nly			11	Protect product (unit of s	ale) from light?			Yes	
a legend device?		No	Is the Product	Neither		Size:	1 x 100mL multiple-dose	e. Shelf life:					24	Months
if yes, enter class #			Orphan Drug Status			0.20.	vial		Initial shelf life at launch	(if different):				Months
a product kit?		No	FD 4 4			Strength:	5GM/100ML (50MG/ML)			ORDER INFOR	MATION			
if yes, list NDCs of component parts			FDA Approval Status				INJECTABLE			ORDER IN OR	MATION			
reverse numbered?		No				Dosage Form:	III O E O I / I D E E		Unit of Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present						Bottle		1 Box contai	ning 1 Vial		
latex-free?		Yes	Container closure is n		atural	Product Shape:			x Box/Carton		(Write-in, e.	J. 1 Box of 10	Vials)	
preservative-free?		Yes	rubber	latex.					Ampule				_	.,
correctional institution block? opioid?	,	Yes No				Product Color:			Glass Tube		Minimum or	der quantity		Yes
Cannabinoid?		No	Country of Origin	India					Vial Liquid Sgl					
If Unit Dose, is item bar coded to	unit dose for		,g			Product Imprint:			Vial Liquid Multi		If Yes, how	nany of which	ch package ty	/pe?
hospital scanning?			Is this product covered u						Vial Powder Sql			Each		
If Unit Dose, indicate NDC here:			Trade Agreements Act (	AA)?	No				Vial Power Multi			Inner/Carton	/Pack	
								<u> </u>	Other: Write In			Case		
			FOR GENERIC DRUG PF	ODUCIS										
					Auth	orized Generic *If Aut	horized Generic, other section		Р	HARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AP						are not applicable	Rec. sell unit to customer? Rx billing unit to pharmacy:						
II. Generic Equivalent to What Bra									x containing 1 Vial		X	Each	,.	
-								(Write-in, e.g.	1 Vial)	<del></del>		Gram		
		DRUG SUPP	LY CHAIN SECURITY ACT	(DSCSA) INFOR	MATION							Milliliter		
Does supplier meet DSCSA defin	nition of manufactur	or?	Yes	GL	d.	0370700000007			ITE	M AND PACKING I	NEORMATIO	ı		
Is product exempt from DSCSA?			No	_	٠.	037070000007				III AIRD I AORINO I	III OIIIIATIOI			
If yes, select exemption:				<del>_</del>				1		Dimensi	ons (US msm	ts )	Volume	
Other exemption - Write in:									Weight Lbs.	Depth	Width	Height	(Cube)	# Pieces:
Is product repackaged?			No	If Y	es, was origi	nal product purchased		Item/Each:	0.309	2.165	2.165	4.528	21.223755	1
Is product sold by manufacturer's	a and the street of the section of	tor?	No	_	ct from mfr?					2.103	2.103	4.520	21.223733	'
				If v	es, attach do	cumentation from FDA.		Box/Carton/B	undle/				0	
Has FDA granted waiver/exception		oduct?	No	_ ",										
				_				Inner Pack:						20
			No	_				Inner Pack: Case:	7.937	11.22	9.055	5.315	539.98859	20
				_	GTIN-	14	Unit of Use GTIN-14			11.22	9.055	5.315		20
Has FDA granted waiver/exception  Saleable Unit of Measure  Item/Each		GT	IN AND HIBCC PRODUCT	_		.14 1700189221	Unit of Use GTIN-14	Case:		11.22	9.055	5.315	539.98859	20
Saleable Unit of Measure  X Item/Each Bow/Carton/Bundle/Inner Pack		Quantity 1	IN AND HIBCC PRODUCT	_	00370	7700189221	Unit of Use GTIN-14	Case:	7.937	11.22			0	
Has FDA granted waiver/exception  Saleable Unit of Measure  x Item/Each Box/Carton/Bundle/Inner Pack Case		GT Quantity	IN AND HIBCC PRODUCT	_	00370		Unit of Use GTIN-14	Case:		11.22				
Saleable Unit of Measure  X Item/Each Bow/Carton/Bundle/Inner Pack		Quantity 1	IN AND HIBCC PRODUCT	_	00370	7700189221	Unit of Use GTIN-14	Case:	7.937	11.22			0	
Has FDA granted waiver/exception  Saleable Unit of Measure  x Item/Each Box/Carton/Bundle/Inner Pack Case		Quantity 1	IN AND HIBCC PRODUCT	_	00370	7700189221	Unit of Use GTIN-14	Case:	7.937  COST INFORMATION	11.22 \$59.40	Vendor #:	WHOLESALI	0	
Has FDA granted waiver/exception  Saleable Unit of Measure  x Item/Each Box/Carton/Bundle/Inner Pack Case		Quantity 1	IN AND HIBCC PRODUCT	_ ′	00370	7700189221	Unit of Use GTIN-14	Case: Pallet:  Regular Cost Invoice Cost (	7.937  COST INFORMATION		Vendor #:	WHOLESALI	0	
Has FDA granted waiver/exception  Saleable Unit of Measure  x Item/Each Box/Carton/Bundle/Inner Pack Case		Quantity 1	IN AND HIBCC PRODUCT	_ ′	00370	7700189221	Unit of Use GTIN-14	Case: Pallet:  Regular Cost	7.937  COST INFORMATION		Vendor #: Whsl. Code	WHOLESALI	0	
Has FDA granted waiver/exception  Saleable Unit of Measure  x Item/Each Box/Carton/Bundle/Inner Pack Case		Quantity 1	HIBCC	NFORMATION	20370	)700189221 )700189225	Unit of Use GTIN-14  ERT, LABEL AND PHOTO OF	Case: Pallet:  Regular Cost Invoice Cost (	7.937  COST INFORMATION  WAC) (\$)		Vendor #: Whsl. Code	WHOLESALI	0	



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Version 2020

For Designated Drop Ship Only Products, Please Use Page 3

	AL HAZARD CLASSIFICATION and TRANSPORTATION
· · ·	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?	No Organic Corrosive Oxidizer 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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#### Version 2020

#### 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 Process	sing
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?	