

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

Version 2021						Introduction Ty	pe:	New Item			Final Version			Date:	3/15/	5/2024
			PRODUCT INFORMA	TION							SPECIAL HAN	IDLING AND STOR	AGE REQUI	REMENTS*		
Company Name: Xiromed LLC Application: ANDA							a. Temperature – Indicate the USP temperature range for this product.									
Application Number for NDA/AN		A/510(k)(med	device):	218	8326		-				ture Range	Controlled Room		and 25 C (68	3° – 77° F)	
Medical Device Class, if applical		, , , , , , , , , , , , , , , , , , ,	,								9					
DUNS:	790387927									Other Te	mperature Range I	Requirement	None			
Proprietary Name (If Applicable) a	and Established Nar	me: G	UANFACINE TABLETS, USP							(wri	te in)					
Selling Unit NDC:	70700-301-01		Unit of Use NDC:				370700301	012		Notes						
UDI			CVX Code:			MVX Code:										
Description:	1 mg: white, oval, f	lat-faced, beve	led-edge tablet with "XI" on one s	ide and "130" or	n other side.					Is this pro	oduct to be shipped	d to customers on i	ce?		No	1
										Is this pro	oduct to be shipped	d to customers on o	ry ice?		No	
Active Ingredient(s):		GUANFACINE	Tablets USP													
									b. Contact for temperature excursion questions:							
URL for Additional Product Inform Address:					Address 2:			Name: Number:				Xiromed Quality 862-895-6230				
City:	180 Park Avenuue Florham Park	State:			NJ <b>Zip:</b> 07932			Number: Group E-mail:				US-Quality-Xiromed@xiromed.com				
Key Contact:	Xiromed Regulator	v			Email:	-			Group E-mail:				U3-Quality	-xii oineu(	axii oiiieu.co	.0111
Phone Number:	(845) 649-7130	y			Fax:	usregulatory@xiromed.com 862-286-0932			c. Special re	gulations f	or product in any	states?			No	7
Product Therapeutic Classificatio	. ,	Attention Defic	it Hyperactivity Disorder (ADHD),	hypertension						-		ts for this product?			No	-
Troduct Therapeatic Glassificatio	,	/ MOTHER POINT	in Thypolada vity Biodradi (18118),	пурополого						Opcolai I	ctumo requiremen	is for this product:			140	7
	ADDITIO	NAL PRODUC	T INFORMATION			PRODUCT DE	ESCRIP <u>TI</u>	ON INFORMATION	d. Store prod	luct (unit o	f sale) upright?				Yes	1
The product is?			Is the Product	Direct-Ship C	Only						product (unit of sa	ale) from light?			No	i
a legend device?		No	Is the Product	Neither	,		11m	m	e. Shelf life:	FIOLECT	Ji Guaci (unit Oi Sa	ac, nom ngm:			24	Months
if yes, enter class #		INO	Orphan Drug Status			Size:			0. 0	Initial sh	elf life at launch (	if different):			24	Months
a product kit?		No				24	1 m	g				,				2
if yes, list NDCs of			FDA Approval Status			Strength:						ORDER INFORM	IATION			
component parts						Dosage Form:	Tab	ets								
reverse numbered?		No								Unit of S			What is the			
co-licensed?		No	Allergens Present								Bottle		1 Bottle con			
latex-free?		Yes	ı	No		Product Shape	e: Ova	I			Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free? correctional institution block?		Yes					whit				Ampule Glass		Minimum o	dor augntiti	.2	Yes
opioid?		Yes No				Product Color	: Will	е			Tube		Wilnimum o	der quantity	y r	res
Cannabinoid?		No	Country of Origin	India			"XI"	on one side and			Vial Liquid Sgl					
If Unit Dose, is item bar coded to u			,g			Product Imprii		)" on other side			Vial Liquid Multi		If Yes, how	many of wh	ich package t	type?
hospital scanning?		No	Is this product covered u	nder the							Vial Powder Sql			Each		31
If Unit Dose, indicate NDC here:			Trade Agreements Act (	ΓAA)?	No						Vial Power Multi			Inner/Cartor	n/Pack	
											Other: Write In			Case		
			FOR GENERIC DRUG PR	ODUCTS												
													/ B			
				_	Au			ed Generic, other				IARMACY ORDER				
I. Orange Book Rating: AB				section fields are not applicable			Rec. sell unit to customer?				Rx billing unit to pharmacy:					
II. Generic Equivalent to What Brand?: TENEX							1 Bottle with 100 Tablets (Write-in, e.g. 1 Vial)				X Each Gram					
		DRIIG SI	JPPLY CHAIN SECURITY ACT (	DSCSA) INFOR	MATION				(vvrite-in, e.g	. 1 Viai)				Milliliter		
		DINOC O	THE CHAIN CESSION AST	DOGGA) IIII GI	(IIIATTON									willinger		
Does supplier meet DSCSA defini	ition of manufacture	er?	Yes		GLN:	00370700000007					ITEN	AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			No													
If yes, select exemption:	İ				GCP:							Dimensi	ons (US msn	nts.)	Volume	Saleable #
Other exemption - Write in:											Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		If yes, was o	riginal product purch	nased		Item/Each:		0.125	1.913	1.52	2.694	7.8335054	1
Is product sold by manufacturer's			No		direct from n						0.120	1.510	1.02	2.004	7.0000004	
Has FDA granted waiver/exceptio		oduct?	No		Provide sour	ce manufacturer for	repackage	ed product	Box/Carton/E	Bundle/					0	
If yes, attach documentation from	m FDA.								Inner Pack:							
			GTIN AND HIBCC PRODUCT I	NEODMATION					Case:		3	11.0625	7.625	3.625	305.77441	24
			GTIN AND HIBCC PRODUCT II	VIFORWATION					Pallet:							
Saleable Unit of Measure	Sa	aleable Quantity	HIBCC		GTI	IN-14	Ur	it of Use GTIN-14	r allet.		650	48	40	52	99840	6000
x Item/Each		1	1.11200			370700301012	Ü.									
Box/Carton/Bundle/Inner Pack							_			cos	TINFORMATION			WHOLESAL	ER USE ONL	LY:
X Case		24			103	70700301019										
Pallet									Regular Cos				Vendor #:			
									Invoice Cost	(WAC) (\$)		\$40.00	Whsl. Code			
									A				Fineline Co	de:		
									As of date:				1			
	_															
			Attach copy of SAFETY DA	ITA SHEET (SE	)S) or non hear	ard letter DACKACE !!	NSEDT	AREL AND PHOTO OF P		AGING and	1 BADCODE					



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

# Version 2021 For Designated Drop Ship Only Products, Please Use Page 3 MATERIAL HAZARD CLASSIFICATION and TRANSPORTATION Is this product (check all that apply): a. Cytotoxic? b. CA Prop. 65 Carcinogen or Reproductive Toxicant?

a. Cytotoxic? 218326 No	SDS Hazard Classification						
b. CA Prop. 65 Carcinogen or Reproductive Toxicant?							
Is the product a CA Prop 65 carcinogen?	Organic Corrosive						
Is the product a CA Prop 65 reproductive toxicant?	Inorganic Oxidizer						
Does the product label bear a CA Prop 65 warning?	Steroid/Androgen Contact Hazard						
•							
c. Contact Hazard?	Does the product have an Aerosol class? If yes,						
d. Does this product require special clean-up instructions?	identify NFPA Storage Level:						
(If yes, attach SDS with special instructions.)	NFPA Storage Level:						
e. Does the product contain DEHP?							
Is this product regulated for shipment by DOT?	Is the product a NIOSH hazardous drug? No						
(if yes, answer a-e below and provide SDS)	If yes, indicate which:						
a. UN/Identification Number							
b. Proper Shipping Name							
c. DOT Hazard Class	Hazardous Waste Identification						
d. Packing Group							
e. Inhalation Hazard?	EPA Hazardous Waste Code: Waste Characteristics						
Is this product regulated for shipment by IATA?							
(if yes, answer a-e below and provide SDS)	REMS or REGISTRY RESTRICTIONS						
a. UN/Identification Number							
b. Proper Shipping Name	Is there a REMS on this product?						
c. DOT Hazard Class	If Yes, is it managed with a pharmacy registry?						
d. Packing Group	Website URL:						
e. Inhalation Hazard?							
Is the product restricted for air shipment? If so, indicate restriction:	Med Guide Required						
Passenger	Limited Distribution Requirement						
Cargo	Comments / Details: (For example, iPledge program?)						
Passenger & Cargo							
Is this a reportable quantity? No	REMS:						
RQ Threshold:	REMS Program Manager Name: Phone:						
Is this a marine pollutant? No	Supplier Manages REMS registry exclusively:						
Is this product shipped utilizing an authorized DOT exception or Special Permit?	Wholesale distributor support:						
No (if yes, identify method below)	Provider Name: DEA #:						
Limited Quantity	Site Enrollment Number assigned  NCPDP#:						
Consumer Commodity, ORM-D	by Supplier: NPI #:						
Small Quantity (49 CFR 173.4) Special Permit; DOT-SP	Comments						
Special Provision (listed in Column 7 of 49 CFR 172.101);	Comments						
SP#	Registry:						
3F#	Registry Program Contact Name: Phone:						
ADD'L STORAGE INFORMATION	Comments						
	Commons						
Is the Product	DETURN INSTRUCTIONS						
Controlled Substance? No Controlled Substance Code Controlled by State(s)? No Listed Chemical (List I or II) No	RETURN INSTRUCTIONS						
Controlled by State(s)? No Listed Chemical (List I or II) No ARCOS Reportable? No If yes, indicate which:	Contact tel. # if product received damaged:						
Schedule No. Is it a scheduled listed chemical product?: No							
	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							
Restricted to retail pharmacy only:	Special regulations or returns requirements for this						
Restricted to hospital, clinics, and physician offices only:	product in certain states?						
Restricted from US territories? (explain in comments)	If so, which states? Other requirements? Comments?						
Comments:							
NIGOSI I AUS	NA VATEO - No long of Double December 1						
MISCELLANEC	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 - 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:	Purchase order daily receipt cut off time by supplier Cut off time:
c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #:  Fax Number: Site Address: Site Address: Name:	Shipping lead time of PO:  Ships same day for next day receipt: Ships for second day receipt: Ships regular ground for 3-10 days receipt:
Phone:	
Expedited Freight Charges or Other Designated Drop Ship Fees:  Expedited freight fees billed with each order:	Overnight and Priority Overnight PO Processing  Overnight receipt available:
Drop Ship service fee billed with each order:	PO Receipt cut off time:
Drop Ship miscellaneous fees billed:  Comments:	Days of week overnight is available:  Monday  Tuesday  Wednesday  Thursday  Friday
	Priority Overnight receipt available:
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:  Phone: Fax: EDI:  Overnight Fees apply: Other 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:  Miscellaneous Notes:	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?
	ADDITIONAL INFORMATION
	Is product order for scheduled patient procedure? Is product order for restocking purposes?