

Ganirelix Formulation

Version 6.4	Revision Date: 16.10.2020	-	S Number: 195-00017	Date of last issue: 23.03.2020 Date of first issue: 15.10.2014
SECTION	1. PRODUCT AND CO	MPA		TION
Produ	uct name	:	Ganirelix Formu	llation
Manı	ufacturer or supplier's	deta	ils	
Comp	bany	:	Organon & Co.	
Addre	ess	:	Rua Treze de M Campinas, São	laio, 1161 Paulo, Brazil B-2220
Telep	bhone	:	551-430-6000	
Emer	gency telephone	:	215-631-6999	
E-ma	il address	:	EHSSTEWARD	@organon.com
Reco	ommended use of the c	hem	ical and restrict	ions on use
Reco	mmended use	:	Pharmaceutical	
Repro	Classification in accor oductive toxicity ific target organ toxicity -	:	Category 1B	BR 14725 Standard ne marrow, Liver, Adrenal gland, spleen, Ovar
•	ated exposure		g, - (
GHS	label elements in acco	rdai	nce with ABNT N	IBR 14725 Standard
Haza	rd pictograms	:		
Signa	al Word	:	Danger	
Haza	rd Statements	:	unborn child. H372 Causes d	amage fertility. Suspected of damaging the amage to organs (Bone marrow, Liver, Adrena Dvary) through prolonged or repeated exposur
Preca	autionary Statements	:	Prevention:	
			P201 Obtain sp	ecial instructions before use.

P264 Wash skin thoroughly after handling.

P270 Do not eat, drink or smoke when using this product.

P280 Wear protective gloves/ protective clothing/ eye protection/ face protection.

Response:

P308 + P313 IF exposed or concerned: Get medical advice/ attention.



Ganirelix Formulation

0.4	10.10.2020	22193-00017	Date of hist issue. 13.10.2014	
Version	Revision Date:	SDS Number:	Date of last issue: 23.03.2020	
6.4	16.10.2020	22195-00017	Date of first issue: 15.10.2014	

Storage:

P405 Store locked up.

Other hazards which do not result in classification

None known.

SECTION 3. COMPOSITION/INFORMATION ON INGREDIENTS

: Mixture

Substance / Mixture

Components

Chemical name	CAS-No.	Classification	Concentration (% w/w)
Ganirelix	124904-93-4	Reproductive toxicity, Category 1B Specific target organ toxicity - repeated exposure (Oral) (Bone marrow, Liver, Adrenal gland, spleen, Ovary), Category 1	>= 0,01 -< 0,1

SECTION 4. FIRST AID MEASURES

General advice	:	In the case of accident or if you feel unwell, seek medical advice immediately. When symptoms persist or in all cases of doubt seek medical advice.
If inhaled	:	If inhaled, remove to fresh air. Get medical attention.
In case of skin contact	:	In case of contact, immediately flush skin with soap and plenty of water. Remove contaminated clothing and shoes. Get medical attention. Wash clothing before reuse. Thoroughly clean shoes before reuse.
In case of eye contact	:	Flush eyes with water as a precaution. Get medical attention if irritation develops and persists.
If swallowed	:	If swallowed, DO NOT induce vomiting. Get medical attention. Rinse mouth thoroughly with water.
Most important symptoms and effects, both acute and delayed	:	May damage fertility. Suspected of damaging the unborn child. Causes damage to organs through prolonged or repeated exposure.
Protection of first-aiders	:	First Aid responders should pay attention to self-protection, and use the recommended personal protective equipment when the potential for exposure exists (see section 8).
Notes to physician	:	Treat symptomatically and supportively.

SECTION 5. FIRE-FIGHTING MEASURES



Ganirelix Formulation

Versi 6.4	on	Revision Date: 16.10.2020		9S Number: 195-00017	Date of last issue: 23.03.2020 Date of first issue: 15.10.2014	
S	Suitable extinguishing media		:	Water spray Alcohol-resistant foam Carbon dioxide (CO2) Dry chemical		
	Unsuita media	ble extinguishing	:	None known.		
	Specific fighting	hazards during fire	:	Exposure to combustion products may be a hazard to health.		
	Hazardo ucts	ous combustion prod-	:	No hazardous cor	nbustion products are known	
	Specific extinguishing meth- ods		:	Use extinguishing measures that are appropriate to local cir- cumstances and the surrounding environment. Use water spray to cool unopened containers. Remove undamaged containers from fire area if it is safe to do so. Evacuate area.		
	Special for fire-f	protective equipment fighters	:	In the event of fire, wear self-contained breathing apparatus. Use personal protective equipment.		
SECI	SECTION 6. ACCIDENTAL RELE		ASE	EMEASURES		
t	tive equ	al precautions, protec- ipment and emer- procedures	:		ective equipment. ing advice (see section 7) and personal ent recommendations (see section 8).	
E	Environmental precautions		:	Avoid release to the environment. Prevent further leakage or spillage if safe to do so. Prevent spreading over a wide area (e.g., by containment oil barriers). Retain and dispose of contaminated wash water. Local authorities should be advised if significant spillages cannot be contained.		
		s and materials for ment and cleaning up	:	For large spills, pr containment to ke can be pumped, s container. Clean up remainir absorbent. Local or national r disposal of this ma employed in the c determine which r Sections 13 and 1	absorbent material. Tovide diking or other appropriate ep material from spreading. If diked material atore recovered material in appropriate ing materials from spill with suitable regulations may apply to releases and aterial, as well as those materials and items leanup of releases. You will need to regulations are applicable. 5 of this SDS provide information regarding tional requirements.	

SECTION 7. HANDLING AND STORAGE

Technical measures	:	See Engineering measures under EXPOSURE CONTROLS/PERSONAL PROTECTION section.
Local/Total ventilation	:	If sufficient ventilation is unavailable, use with local exhaust ventilation.



Ganirelix Formulation

Version 6.4	Revision Date: 16.10.2020	SDS Number: 22195-00017	Date of last issue: 23.03.2020 Date of first issue: 15.10.2014			
Advice on safe handling		 Do not get on skin or clothing. Do not breathe mist or vapors. Do not swallow. Avoid contact with eyes. Wash skin thoroughly after handling. Handle in accordance with good industrial hygiene and safet practice, based on the results of the workplace exposure assessment Keep container tightly closed. Do not eat, drink or smoke when using this product. Take care to prevent spills, waste and minimize release to th environment. 				
Hygiene measures		flushing syste place. When using d Wash contam The effective engineering c appropriate de industrial hygi	 If exposure to chemical is likely during typical use, provide ey flushing systems and safety showers close to the working place. When using do not eat, drink or smoke. Wash contaminated clothing before re-use. The effective operation of a facility should include review of engineering controls, proper personal protective equipment, appropriate degowning and decontamination procedures, industrial hygiene monitoring, medical surveillance and the use of administrative controls. 			
Cond	ditions for safe storage	Store locked u Keep tightly c	 Keep in properly labeled containers. Store locked up. Keep tightly closed. Store in generating with the particular patients regulations. 			
Materials to avoid		 Store in accordance with the particular national regulations. Do not store with the following product types: Strong oxidizing agents Organic peroxides Explosives Gases 				

SECTION 8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Ingredients with workplace control parameters						
Components	CAS-No.	Value type (Form of exposure)	Control parame- ters / Permissible concentration	Basis		
Ganirelix	124904-93-4	TŴA	0.2 µg/m3 (OEB 5)	Internal		
		Wipe limit	2 µg/100 cm ²	Internal		

Engineering measures	 Use closed processing systems or containment technologies to control at source (e.g., glove boxes/isolators) and to prevent leakage of compounds into the workplace. All engineering controls should be implemented by facility design and operated in accordance with GMP principles to protect products, workers, and the environment. No open handling permitted. Totally enclosed processes and materials transport systems are required. Operations require the use of appropriate containment
	Operations require the use of appropriate containment technology designed to prevent leakage of compounds into

Ganirelix Formulation



Version 6.4	Revision Date: 16.10.2020	SDS Number: 22195-00017	Date of last issue: 23.03.2020 Date of first issue: 15.10.2014				
		the workpla	ce.				
Pers	onal protective equip	ment					
Resp	iratory protection	: No persona required.	No personal respiratory protective equipment normally				
Hand	l protection						
M	aterial	: Chemical-re	Chemical-resistant gloves				
Remarks Eye protection Skin and body protection		 Wear safety If the work of mists or aei Wear a face potential for aerosols. Work unifor Additional b task being p disposable 	buble gloving. y glasses with side shields or goggles. environment or activity involves dusty conditions, rosols, wear the appropriate goggles. eshield or other full face protection if there is a direct contact to the face with dusts, mists, or m or laboratory coat. body garments should be used based upon the performed (e.g., sleevelets, apron, gauntlets, suits) to avoid exposed skin surfaces. riate degowning techniques to remove potentially ed clothing.				

SECTION 9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance	:	Aqueous solution
Color	:	No data available
Odor	:	No data available
Odor Threshold	:	No data available
рН	:	5
Melting point/freezing point	:	No data available
Initial boiling point and boiling range	:	100 °C
Flash point	:	No data available
Evaporation rate	:	No data available
Flammability (solid, gas)	:	Not applicable
Flammability (liquids)	:	No data available
Upper explosion limit / Upper flammability limit	:	No data available
Lower explosion limit / Lower flammability limit	:	No data available
Vapor pressure	:	23 hPa (20 °C)



Ganirelix Formulation

Versi 6.4	ion	Revision Date: 16.10.2020		S Number: 95-00017	Date of last issue: 23.03.2020 Date of first issue: 15.10.2014				
	Deleting								
	Relative	e vapor density		No data available	3				
	Relative	e density	:	1					
	Solubility(ies) Water solubility		:	completely miscil	ble				
	Partition coefficient: n- octanol/water		:	: No data available					
	Autoignition temperature		:	No data available	9				
	Decomposition temperature		:	No data available	9				
,	Viscosity Viscosity, kinematic		:	: No data available					
	Explosive properties		:	Not explosive					
	Oxidizing properties		:	The substance of	r mixture is not classified as oxidizing.				
	Molecular weight		:	No data available					
	Particle size			No data available					

SECTION 10. STABILITY AND REACTIVITY

Reactivity Chemical stability Possibility of hazardous reac- tions	:	Not classified as a reactivity hazard. Stable under normal conditions. Can react with strong oxidizing agents.
Conditions to avoid Incompatible materials Hazardous decomposition products	:	None known. Oxidizing agents No hazardous decomposition products are known.

SECTION 11. TOXICOLOGICAL INFORMATION

Information on likely routes of exposure	:	Inhalation Skin contact
		Ingestion Eye contact

Acute toxicity

Not classified based on available information.

Components:

Ganirelix:

Acute toxicity (other routes of : LD50 (Rat): 40 mg/kg administration)

Skin corrosion/irritation

Not classified based on available information.



Ganirelix Formulation

rsion	Revision Date: 16.10.2020	-	9S Number: 195-00017	Date of last issue: 23.03.2020 Date of first issue: 15.10.2014
Serio	us eye damage/eye i	irritati	on	
	assified based on ava			
Comp	oonents:			
Ganir	elix:			
Specie		:	Rabbit	
Result Metho		:	Mild eye irritation Draize Test	
Metho		•	Dialze Test	
Respi	ratory or skin sensi	tizatio	n	
Skin s	sensitization			
Not cl	assified based on ava	ailable	information.	
-	ratory sensitization			
	assified based on ava	ailable	information.	
<u>Comp</u>	oonents:			
Conir	elix:			
Ganir				
Test T		:	Maximization Tes	st
Test T Specie Result	cell mutagenicity	:	Guinea pig negative	st
Test T Specie Result Germ Not cla	cell mutagenicity assified based on ava	: : ailable	Guinea pig negative	st
Test T Specie Result Germ Not cla <u>Comp</u>	es t cell mutagenicity assified based on ava ponents:	: : ailable	Guinea pig negative	st
Test T Specie Result Germ Not cla <u>Comp</u> Ganir	es t cell mutagenicity assified based on ava ponents:	i ailable	Guinea pig negative information.	st se mutation assay
Test T Specie Result Germ Not cla <u>Comp</u> Ganir	es t cell mutagenicity assified based on ava <u>conents:</u> elix:	ailable	Guinea pig negative information. Test Type: revers Test system: Sal	
Test T Specie Result Germ Not cla <u>Comp</u> Ganir	es t cell mutagenicity assified based on ava <u>conents:</u> elix:	ailable	Guinea pig negative information. Test Type: revers	se mutation assay
Test T Specie Result Germ Not cla <u>Comp</u> Ganir	es t cell mutagenicity assified based on ava <u>conents:</u> elix:	ailable	Guinea pig negative information. Test Type: revers Test system: Sal Result: negative Test Type: revers	se mutation assay monella typhimurium se mutation assay
Test T Specie Result Germ Not cla <u>Comp</u> Ganir	es t cell mutagenicity assified based on ava <u>conents:</u> elix:	ailable	Guinea pig negative information. Test Type: revers Test system: Sal Result: negative Test Type: revers Test system: Esc	se mutation assay monella typhimurium se mutation assay
Test T Specie Result Germ Not cla <u>Comp</u> Ganir	es t cell mutagenicity assified based on ava <u>conents:</u> elix:	ailable	Guinea pig negative information. Test Type: revers Test system: Sal Result: negative Test Type: revers Test system: Esc Result: negative	se mutation assay monella typhimurium se mutation assay cherichia coli
Test T Specie Result Germ Not cla <u>Comp</u> Ganir	es t cell mutagenicity assified based on ava <u>conents:</u> elix:	ailable	Guinea pig negative information. Test Type: revers Test system: Sal Result: negative Test Type: revers Test system: Esc Result: negative Test Type: in vitr	se mutation assay monella typhimurium se mutation assay cherichia coli
Test T Specie Result Germ Not cla <u>Comp</u> Ganir	es t cell mutagenicity assified based on ava <u>conents:</u> elix:	ailable	Guinea pig negative information. Test Type: revers Test system: Sal Result: negative Test Type: revers Test system: Esc Result: negative Test Type: in vitr	se mutation assay monella typhimurium se mutation assay cherichia coli
Test T Specia Result Mot cla Comp Ganir Genot	t cell mutagenicity assified based on ava <u>conents:</u> elix: coxicity in vitro	ailable	Guinea pig negative information. Test Type: revers Test system: Sal Result: negative Test Type: revers Test system: Esc Result: negative Test Type: in vitr Test Type: in vitr Test system: Chi Result: negative	se mutation assay monella typhimurium se mutation assay cherichia coli o test nese hamster ovary cells
Test T Specia Result Mot cla Comp Ganir Genot	es t cell mutagenicity assified based on ava <u>conents:</u> elix:	ailable :	Guinea pig negative information. Test Type: revers Test system: Sal Result: negative Test Type: revers Test system: Esc Result: negative Test Type: in vitr Test system: Chi Result: negative Test Type: In vivo Species: Mouse	se mutation assay monella typhimurium se mutation assay cherichia coli o test nese hamster ovary cells o micronucleus test
Test T Specia Result Mot cla Comp Ganir Genot	t cell mutagenicity assified based on ava <u>conents:</u> elix: coxicity in vitro	ailable :	Guinea pig negative information. Test Type: revers Test system: Sal Result: negative Test Type: revers Test system: Esc Result: negative Test Type: in vitr Test system: Chi Result: negative Test Type: In vivr Species: Mouse Application Route	se mutation assay monella typhimurium se mutation assay cherichia coli o test nese hamster ovary cells o micronucleus test
Test T Specie Result Or Cla Comp Ganir Genot	t cell mutagenicity assified based on ava <u>conents:</u> elix: coxicity in vitro	ailable	Guinea pig negative information. Test Type: revers Test system: Sal Result: negative Test Type: revers Test system: Esc Result: negative Test Type: in vitr Test Type: in vitr Test system: Chi Result: negative Test Type: In vivo Species: Mouse Application Route Result: negative	se mutation assay monella typhimurium se mutation assay cherichia coli o test nese hamster ovary cells o micronucleus test

Not classified based on available information.



Ganirelix Formulation

rsion L	Revision Date: 16.10.2020		S Number: 195-00017	Date of last issue: 23.03.2020 Date of first issue: 15.10.2014
-	ductive toxicity amage fertility. Suspect	ed o	of damaging the un	born child.
Comp	onents:			
Ganire	elix:			
Effects	s on fertility	:	Species: Rat Application Route Duration of Single Fertility: LOAEL: Result: Effects on	e Treatment: 13 Weeks),1 μg/kg
			Species: Rat, fem Application Route Duration of Single Fertility: LOAEL:	ale : Subcutaneous e Treatment: 8 Weeks
			Test Type: Fertilit Species: Monkey Application Route Fertility: NOAEL: Result: Effects on	: Subcutaneous 0,02 mg/kg body weight
Effects	on fetal development	:	Species: Rat, fem Application Route	:: Subcutaneous city.: LOAEL: 10 μg/kg
			Species: Rabbit, Application Route	:: Subcutaneous city.: LOAEL: 30 μg/kg
Reproc sessm	ductive toxicity - As- ent	:	fertility, based on	adverse effects on sexual function and animal experiments., Some evidence of n development, based on animal

STOT-repeated exposure

Causes damage to organs (Bone marrow, Liver, Adrenal gland, spleen, Ovary) through prolonged or repeated exposure.

Components:

Ganirelix:

Routes of exposure	:	Ingestion
Target Organs	:	Bone marrow, Liver, Adrenal gland, spleen, Ovary

Ganirelix Formulation



Versio 6.4	n Revision Date: 16.10.2020		DS Number: 2195-00017	Date of last issue: 23.03.2020 Date of first issue: 15.10.2014
Δ	ssessment	:	Causes damage exposure.	to organs through prolonged or repeated
F	epeated dose toxicity			
<u>c</u>	components:			
S N L A E	anirelix: pecies IOAEL OAEL opplication Route xposure time arget Organs	:	Rat 0,02 mg/kg 2 mg/kg Subcutaneous 6 Months Bone marrow	
L A E	pecies OAEL pplication Route xposure time arget Organs		Mouse, female 0,3 mg/kg Subcutaneous 3 Months Liver, Adrenal gla	nd, spleen, Ovary
L A E	pecies OAEL pplication Route xposure time arget Organs		Mouse, male 3 mg/kg Subcutaneous 3 Months Liver, Adrenal gla	nd, spleen
N A E	pecies IOAEL pplication Route xposure time temarks	· · · ·	Monkey 2,5 mg/kg Subcutaneous 6 Months No significant adv	verse effects were reported

Aspiration toxicity

Not classified based on available information.

:

Experience with human exposure

Components:

Ganirelix:

Inhalation

Symptoms: The most common side effects are:, vaginal bleeding, Headache, Abdominal pain, Nausea, ectopic pregnancy, miscarriage

SECTION 12. ECOLOGICAL INFORMATION

Components:

Ganirelix:

Ecotoxicology Assessment

Acute aquatic toxicity : No data available



Ganirelix Formulation

Versic 6.4	on Revision Date: 16.10.2020		Number: 5-00017	Date of last issue: 23.03.2020 Date of first issue: 15.10.2014
C	Chronic aquatic toxicity	: N	lo data available	
	Persistence and degradabil No data available	ity		
	Bioaccumulative potential No data available			
	lobility in soil Io data available			
-	Other adverse effects Io data available			
SECT	ION 13. DISPOSAL CONSI	DERA	TIONS	
D	Disposal methods			
	Vaste from residues Contaminated packaging	: E	mpty containers	ordance with local regulations. should be taken to an approved waste ecycling or disposal.

If not otherwise specified: Dispose of as unused product.

SECTION 14. TRANSPORT INFORMATION

International Regulations

UNRTDG

Not regulated as a dangerous good

IATA-DGR

Not regulated as a dangerous good

IMDG-Code

Not regulated as a dangerous good

Transport in bulk according to Annex II of MARPOL 73/78 and the IBC Code

Not applicable for product as supplied.

Domestic regulation

ANTT

Not regulated as a dangerous good

SECTION 15. REGULATORY INFORMATION

Safety, health and environmental regulations/legisl mixture	ation specific for the substance of	or
National List of Carcinogenic Agents for Humans - (LINACH)	: Not applicable	
Brazil. List of chemicals controlled by the Federal Police	: Not applicable	



Ganirelix Formulation

Version 6.4	Revision Date: 16.10.2020		S Number: 95-00017	Date of last issue: 23.03.2020 Date of first issue: 15.10.2014
Intern	national Regulations			
	•		•	e following inventories:
AICS		: 1	not determined	
DSL		: 1	not determined	
IECS	С	: 1	not determined	

SECTION 16. OTHER INFORMATION

Further information

Sources of key data used to	:	Internal technical data, data from raw material SDSs, OECD
compile the Material Safety		eChem Portal search results and European Chemicals Agen-
Data Sheet		cy, http://echa.europa.eu/

Full text of other abbreviations

AIIC - Australian Inventory of Industrial Chemicals; ANTT - National Agency for Transport by Land of Brazil; ASTM - American Society for the Testing of Materials; bw - Body weight; CMR -Carcinogen, Mutagen or Reproductive Toxicant; DIN - Standard of the German Institute for Standardisation; DSL - Domestic Substances List (Canada); ECx - Concentration associated with x% response; ELx - Loading rate associated with x% response; EmS - Emergency Schedule; ENCS - Existing and New Chemical Substances (Japan); ErCx - Concentration associated with x% growth rate response; ERG - Emergency Response Guide; GHS - Globally Harmonized System; GLP - Good Laboratory Practice; IARC - International Agency for Research on Cancer; IATA - International Air Transport Association; IBC - International Code for the Construction and Equipment of Ships carrying Dangerous Chemicals in Bulk; IC50 - Half maximal inhibitory concentration; ICAO - International Civil Aviation Organization; IECSC - Inventory of Existing Chemical Substances in China; IMDG - International Maritime Dangerous Goods; IMO - International Maritime Organization; ISHL - Industrial Safety and Health Law (Japan); ISO - International Organisation for Standardization; KECI - Korea Existing Chemicals Inventory; LC50 - Lethal Concentration to 50 % of a test population; LD50 - Lethal Dose to 50% of a test population (Median Lethal Dose); MARPOL - International Convention for the Prevention of Pollution from Ships; n.o.s. - Not Otherwise Specified; Nch - Chilean Norm; NO(A)EC - No Observed (Adverse) Effect Concentration; NO(A)EL - No Observed (Adverse) Effect Level; NOELR - No Observable Effect Loading Rate; NOM - Official Mexican Norm; NTP - National Toxicology Program; NZIoC - New Zealand Inventory of Chemicals; OECD - Organization for Economic Co-operation and Development; OPPTS - Office of Chemical Safety and Pollution Prevention; PBT - Persistent, Bioaccumulative and Toxic substance; PICCS - Philippines Inventory of Chemicals and Chemical Substances; (Q)SAR - (Quantitative) Structure Activity Relationship; REACH - Regulation (EC) No 1907/2006 of the European Parliament and of the Council concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals; SADT - Self-Accelerating Decomposition Temperature; SDS - Safety Data Sheet; TCSI - Taiwan Chemical Substance Inventory; TDG - Transportation of Dangerous Goods; TSCA - Toxic Substances Control Act (United States); UN - United Nations; UNRTDG - United Nations Recommendations on the Transport of Dangerous Goods; vPvB - Very Persistent and Very Bioaccumulative; WHMIS - Workplace Hazardous Materials Information System

The information provided in this Safety Data Sheet is correct to the best of our knowledge, information and belief at the date of its publication. The information is designed only as a guidance for safe handling, use, processing, storage, transportation, disposal and release and



Ganirelix Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 23.03.2020
6.4	16.10.2020	22195-00017	Date of first issue: 15.10.2014

shall not be considered a warranty or quality specification of any type. The information provided relates only to the specific material identified at the top of this SDS and may not be valid when the SDS material is used in combination with any other materials or in any process, unless specified in the text. Material users should review the information and recommendations in the specific context of their intended manner of handling, use, processing and storage, including an assessment of the appropriateness of the SDS material in the user's end product, if applicable.

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