THYMUS- thymus (suis) liquid BioActive Nutrional, Inc.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Drug Facts:

ACTIVE INGREDIENTS:

Thymus (Suis) 8X.

INDICATIONS:

Regulation and stimulation of the Thymus Gland. For temporary relief of symptoms due to lowered immune response, viral infections, or influenza.

WARNINGS:

If pregnant or breast-feeding, ask a health care professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Do not use if tamper evident seal is broken or missing.

Store in a cool, dry place.

KEEP OUT OF REACH OF CHILDREN:

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

DIRECTIONS:

Three sprays per day. Hold in mouth for three minutes.

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INACTIVE INGREDIENTS:

Citric Acid, Demineralized Water, Ethanol 0.005%, Glycerin, Sodium Benzoate.

QUESTIONS:

Distributed by:

Bio Active Nutritional, Inc.

Melbourne, FL 32935

bioactivenutritional.com

PACKAGE LABEL DISPLAY:

BIOActive **Nutritional**

THYMUS

HOMEOPATHIC

1 FL OZ (30 ml)



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LOT:

Distributed by: BioActive Nutritional, Inc. 1803 N. Wickham Rd. Melbourne, FL 32935 bioactivenutritional.com

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THYMUS

thymus (suis) liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:43857-0194

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name			
Ingredient Name	Basis of Strength	Strength	
SUS SCROFA THYMUS (UNII: 7B69B0BD62) (SUS SCROFA THYMUS - UNII: 7B69B0BD62)	SUS SCROFA THYMUS	8 [hp_X] in 1 mL	

Inactive Ingredients				
Ingredient Name	Strength			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				
GLYCERIN (UNII: PDC6A3C0OX)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
WATER (UNII: 059QF0KO0R)				
ALCOHOL (UNII: 3K9958V90M)				

ı	P	ackaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:43857- 0194-1	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	12/06/2016	

keting End Date

Labeler - BioActive Nutrional, Inc. (624980496)

Registrant - Apotheca Company (844330915)

Establishment			
Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43857-0194), api manufacture(43857-0194), label(43857-0194), pack(43857-0194)

Revised: 7/2021 BioActive Nutrional, Inc.