

**GABA- gaba, liquid**  
**Deseret Biologicals, 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.*

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**Drug Facts**

**ACTIVE INGREDIENTS:**

Gaba 6X, 12X, 30X, 200X, 12C, 30C, 60C, 200C.

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**INDICATIONS:**

For temporary relief of symptoms related to Gaba sensitivity including hypertension, anger, hostility, Seasonal Affective Disorder, sleep difficulties, fatigue, depression, food sensitivities and sleep disorders.\*\*

\*\*These statements are based upon homeopathic principles. They have not been reviewed by the Food and Drug Administration.

**WARNINGS:**

Keep out of reach of children. In case of overdose, contact physician or Poison Control Center right away.

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

Tamper seal: "Sealed for Your Protection." Do not use if seal is broken or missing.

**DIRECTIONS:**

1-10 drops under the tongue, 3 times a day or as directed by a health professional. Consult a physician for use in children under 12 years of age.

**INACTIVE INGREDIENTS:**

Demineralized water, 25% Ethanol.

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## QUESTIONS:

Dist. By: Deseret Biologicals, Inc.

469 Parkland Drive

Sandy, UT 84070

www.desbio.com

## PACKAGE LABEL DISPLAY:

### DESBIO

NDC 43742-0178-1

### HOMEOPATHIC

### GABA

1 FL OZ (30 ml)

#### WARNINGS:

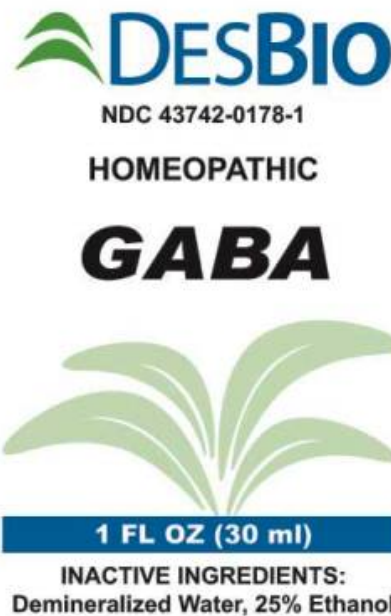
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#### LOT:

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## GABA

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### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43742-0178
Route of Administration	ORAL		

<b>Active Ingredient/Active Moiety</b>				
<b>Ingredient Name</b>			<b>Basis of Strength</b>	<b>Strength</b>
.GAMMA.-AMINOBUTYRIC ACID (UNII: 2ACZ6IPC6I) (.GAMMA.-AMINOBUTYRIC ACID - UNII:2ACZ6IPC6I)			.GAMMA.-AMINOBUTYRIC ACID	6 [hp_X] in 1 mL
<b>Inactive Ingredients</b>				
<b>Ingredient Name</b>			<b>Strength</b>	
WATER (UNII: 059QF0KO0R)				
ALCOHOL (UNII: 3K9958V90M)				
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43742-0178-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
<b>Marketing Information</b>				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved homeopathic		11/07/2012		

**Labeler** - Deseret Biologicals, Inc. (940741853)

**Registrant** - Apotheca Company (844330915)

### **Establishment**

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0178) , api manufacture(43742-0178) , label(43742-0178) , pack(43742-0178)

Revised: 9/2015

Deseret Biologicals, Inc.