

**AMOLG A- betamethasone valerate, gentamicin sulfate cream**  
**OASIS TRADING**

*Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, [click here](#).*

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Betamethasone Valerate 0.61mg/g  
Gentamicin Sulfate 1mg/g

temporary relief of itching associated with minor skin irritations and rashes due to eczema, insect bites, poison ivy, poison oak, or poison sumac soaps detergents, cosmetics, jewelry, seborrheic dermatitis, psoriasis, external genital and anal itching

Keep out of reach of children

Adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily

Do not use in the eyes by putting this product into the rectum by using fingers or any mechanical device or applicator

Ask a doctor before use if you have a vaginal discharge rectal bleeding  
diaper rash

When using this product consult a doctor before exceeding recommended dosage

Stop use and ask a doctor if condition gets worse condition persists for more than 7 days  
condition clears up and occurs again within a few days. Do not begin to use any other hydrocortisone product unless you have consulted a doctor.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

cetostearyl alcohol, propylene glycol, purified water, sodium lauryl sulfate, white petrolatum

For external use only

# AMOLG A CREAM

<b>Drug Facts</b>	
<b>Active Ingredients (in 1g)</b>	<b>Purpose</b>
Betamethasone Valerate 0.61mg -----	Anti-itch
Gentamicin Sulfate 1mg -----	Anti-itch
<b>Uses</b>	
<p>■ temporary relief of itching associated with minor skin irritations and rashes due to eczema, insect bites, poison ivy, poison oak, or poison sumac soaps detergents, cosmetics, jewelry, seborrheic dermatitis, psoriasis, external genital and anal itching</p>	
<b>Warnings</b>	
<p>Do not use in the eyes by putting this product into the rectum by using fingers or any mechanical device or applicator</p> <p><b>Ask a doctor before use if you have</b> a vaginal discharge, rectal bleeding, diaper rash</p> <p><b>When using this product</b> consult a doctor before exceeding recommended dosage</p> <p><b>Stop use and ask a doctor if</b> condition gets worse condition persists for more than 7 days</p> <p>condition clears up and occurs again within a few days. Do not begin to use any other hydrocortisone product unless you have consulted a doctor.</p> <p><b>Keep out of reach of children.</b> If swallowed, get medical help or contact a Poison Control Center right away.</p>	
<b>Directions</b>	
<p>■ Adults and children 2 years of age and older: apply to the affected area no more than 3 to 4 times daily</p>	
<b>Other Information</b>	
<p>■ Store at room temperature</p>	
<b>Inactive Ingredient</b>	
<p>cetostearyl alcohol, propylene glycol, purified water, sodium lauryl sulfate, white petrolatum</p>	
<b>Questions or comments ?</b>	
<p>Call weekdays from 9 a.m to 5 p.m EST at (201) 669-8405</p> <p><b>Distributed By: P&amp;K FRONTIER MARKETING CORP.</b> 329 BROAD AVENUE # 2F, PALISADES PARK, NJ 07650, USA</p>	
<b>Made in South Korea</b>	

## AMOLG A

betamethasone valerate, gentamicin sulfate cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:72689-0033
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BETAMETHASONE VALERATE (UNII: 9IFA5XM7R2) (BETAMETHASONE - UNII:9842X06Q6M)	BETAMETHASONE VALERATE	0.61 mg in 1 g
GENTAMICIN SULFATE (UNII: 8X7386QRLV) (GENTAMICIN - UNII:T6Z9V48IKG)	GENTAMICIN	1 mg in 1 g

### Inactive Ingredients

Ingredient Name			Strength	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)				
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72689-0033-1	35 g in 1 TUBE; Type 0: Not a Combination Product	11/16/2018	
<b>Marketing Information</b>				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		11/15/2018		

**Labeler** - OASIS TRADING (689991468)

**Registrant** - OASIS TRADING (689991468)

Establishment			
Name	Address	ID/FEI	Business Operations
OASIS TRADING		689991468	manufacture(72689-0033)