## KEEPER MEDAL- mineral liquid AML Bio Co., Ltd.

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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## **Drug Facts**

mineral complex

antiseptic

KEEP OUT OF REACH OF THE CHILDREN

apply proper amount to the subject area

For external use only.

- Do not use in eyes.
- If swallowed, get medical help promptly.
- Stop use, ask doctor If irritation occurs.
- Keep out of reach of children.

for external use only

Water

Active Ingredients Mineral complex	Purpose antibacterial
Uses	
antibacterial	
Warning	
For external use only	
When using this product if following abnormal symptoms persi	st discontinue use
Irritation around the eyes, ears, mucous	
the skin irritation and rashes	
Stop immediately and consult a doctor	
<ol> <li>Hypersensitivity symptoms such as en 2) Skin Irritation</li> </ol>	thema, itching and dermatitis.
<ol> <li>Following Instructions when using me</li> </ol>	dication
(1) For external use only (Do not use inte	
(2) Avoid getting into the eyes (if contac	
Do not use the product for a lon inflammation or sickness may occur due	
It is not recommended to use this one	
with a cast or bandage.	
Directions	
apply proper amount where needed	
Other Information	
<ul> <li>read the directions and warnings before avoid fracting and expensive heat above</li> </ul>	
- avoid freezing and excessive heat abov	e 40 degree C (104 degree F)
Inactive Ingredients	
water	

KEEPER MEDAL mineral liquid							
Product Information							
Product Type	HUMAN OTC DRUG	Item Code	e (Source)	NDC:81016-0001			
Route of Administration	TOPICAL						
Active Ingredient/Active Moiety							
Ingredi	<b>Basis of Streng</b>	th Strength					
MINERAL OIL (UNII: T5L8T28FGP) (MINERAL OIL - UNII:T5L8T28FGP)			MINERAL OIL	100 g in 100 mL			

In	active Ingre	edients				
Ingredient Name					Strength	
w,	<b>ATER</b> (UNII: 0590	QF0KO0R)				
Packaging						
#	ltem Code	Package Description	Marketin Da	-	Marketing End Date	
	NDC:81016- 0001-1	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/09/2020			
Marketing Information						
	-		Markati	na Start	Markating End	
	Marketing Category	Application Number or Monograph Citation		ng Start Ite	Marketing End Date	
	approved drug Ier		11/09/2020			

Labeler - AML Bio Co., Ltd. (695067521)

Registrant - AML Bio Co., Ltd. (695067521)

## Establishment

Name	Address	ID/FEI	<b>Business Operations</b>
AML Bio Co., Ltd.		695067521	manufacture(81016-0001)

Revised: 1/2024

AML Bio Co., Ltd.