## AUVON 4% MENTHOL PAIN RELIEF GEL- menthol gel SHENZHEN YUWEN E-COMMERCE CO., LTD.

-----

## 83391-007

Auvon 4% Menthol Pain Relief Gel

Menthol 4%

**Topical Analgestic** 

Temporary relief from minor aches and paons of sore muscles and joints

For external use only

If you are under 12 years of age

If you are allergic to the listed ingredients

If you are pregnent or breast feeding

On wounds, cuts, damaged/broken/irritated skin

- Use only as directed
- Read and follow all directions and warnings on this label

You are experiencing pain, swelling or blistering

Redness is persent or irritation develops

Symptoms persist for more than 7 days or clear up and occur again within a few days If swallowed, get medical help or contact a Poison Control Center right away.

Clean and dry the patch application area (no wound or hair), pull to separate the film, peel off one side of the film, apply the exposed patch to the skin, peel off the other side of the film, and press the patch firmly 

Do not repeatedly use or repeatedly reapply the patch as far as possible

Use in te affected area no more than 4 times daily

Wash hands with cool water after use

Store it in a dray place at a room temperature between 20-25°c and away from direct sunlight

Arctium Lappa (Burdock) Extract

Arnica Montana Flower Extract

Aloe Barbadensis Leaf Juice

**Aminomethyl Propanol** 

Boswellia Serrata Extract

Camellia Sinensis (Green Tea) Leaf Extract

Camphor

Carbomer

Calendula Officinalis Flower Extract

FD&C Blue NO.1

FD&C Yellow No.5

Isopropyl Alcohol

Isopropyl Myristate,

Glycerin

Llex Paraguariensis Leaf Extract

Melissa Officinalis Leaf Extract

Methyl Methacrylate Crosspolymer

**Tocopheryl Acetate** 

Water



## **AUVON 4% MENTHOL PAIN RELIEF GEL**

menthol gel

<b>Product Information</b>
----------------------------

Product Type HUMAN OTC DRUG Item Code (Source) NDC:83391-007

**Route of Administration** TOPICAL

Active Ingredient/Active Moiety					
Ingredient Name	<b>Basis of Strength</b>	Strength			
LEVOMENTHOL (UNII: BZ1R15MTK7) (LEVOMENTHOL - UNII:BZ1R15MTK7)	LEVOMENTHOL	0.04 mg in 1 mL			

Inactive Ingredients	
Ingredient Name	Strength
ARNICA MONTANA FLOWER WATER (UNII: U7L2JP51PR)	
ALOE BARBADENSIS LEAF JUICE (UNII: ZY81Z83H0X)	
CARBOMER (UNII: 0A5MM307FC)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
AMINOMETHYL PROPANOL (UNII: LU49E6626Q)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
WATER (UNII: 059QF0KO0R)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
BOSWELLIA SERRATA OIL (UNII: 5T1XCE6K8K)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
ARCTIUM LAPPA WHOLE (UNII: 73070DU1LA)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
CAMELLIA SINENSIS LEAF OIL (UNII: VC855RRT77)	
METHYL METHACRYLATE/GLYCOL DIMETHACRYLATE CROSSPOLYMER (UNII: EG97988M5Q)	
GLYCERIN (UNII: PDC6A3C0OX)	
MELISSA OFFICINALIS LEAF OIL (UNII: PTP6R7263M)	
CAMPHOR (NATURAL) (UNII: N20HL7Q941)	

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:83391-007- 01	89 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/08/2025		
2	NDC:83391-007- 02	2 in 1 PACKAGE	02/08/2025		
2	NDC:83391-007- 01	89 mL in 1 BOTTLE; Type 0: Not a Combination Product			
3	NDC:83391-007- 03	3 in 1 PACKAGE	02/08/2025		
_	NDC:83391-007-	89 mL in 1 BOTTLE; Type 0: Not a Combination			

01	Product				
Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M017	02/08/2025			

## Labeler - SHENZHEN YUWEN E-COMMERCE CO., LTD. (544559614)

Revised: 2/2025 SHENZHEN YUWEN E-COMMERCE CO., LTD.