AVCOO MENTHOL PAIN RELIEF PATCH- menthol patch ADP Health Limited

84992-004

AVCOO Menthol Pain Relief Patch

Menthol 5%

Topical anesthetic

for the temporary pain relief

For external use only

If you are allergic to the listed ingredients

If you are pregnent or breast feeding

If you are under 12 years of age

On wounds, cuts, damaged/broken/irritated skin

On eyes or mucous membranes

With heating pads/devices or wrap with a bandage

With other topical analgesics

If the package arrives damaged or opened.

Use only as directed

Read and follow all directions and warnings on this label

avoid contact with the eyes and mucous membranes

rare cases of serious burns have been reported with products of this type

a transient burning sensation may occur upon application but generally disappears in several days

dispose of used patch in manner that always keeps product away from children and pets. Used patches still contain the drug product that can produce serious adverse effects if a child or pet chews or ingests this patch.

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 ingested, 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 Do not repeatedly use or repeatedly reapply the patch as far as possible
- Cut first if if used on joints
- Use in te affected area no more than 4 times daily

■ Wash hands with cool water after use

Avoid storing patches in direct sunlight

Protect patches from excessive moisture

P lease store it at room temperature.

Aloe Barbadensis Leaf Extract

Arnica Montana Flower Extract

Boswellia Carterii Resin Extract

Camellia Sinensis Leaf Extract

Carboxymethylcellulose Sodium

Dihydroxyaluminum Aminoacetate

Edetate Disodium

Glycerin

Hydroxyacetophenone

Kaolin

L-Tartaric Acid

Mineral Oil

Petrolatum

Polyacrylic Acid

Polysorbate 80

Propylene Glycol

PVP

Sodium Polyacrylate

Titanium Dioxide

Water



AVCOO MENTHOL PAIN RELIEF PATCH

menthol patch

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:84992-004

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LEVOMENTHOL (LINII) R71D15MTV7) (LEVOMENTHOL LINII) R71D15MTV7)	LEVOMENTHOL	500 mg

Inactive Ingredients Ingredient Name PROPYLENE GLYCOL (UNII: 6DC9Q167V3) Strength

KAOLIN (UNII: 24H4NWX5CO)

ALOE BARBADENSIS LEAF POWDER (UNII: ZY81Z83H0X)

ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)

MINERAL OIL (UNII: T5L8T28FGP)

CARBOXYMETHYLCELLULOSE SODIUM (0.7 CARBOXYMETHYL SUBSTITUTION PER SACCHARIDE; 100-200 MPA.S AT 1%) (UNII: 99H65D77XY)

HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)

WATER (UNII: 059QF0K00R)

DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6)

PVP (UNII: FZ989GH94E)

POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)

PETROLATUM (UNII: 4T6H12BN9U)

POLYSORBATE 80 (UNII: 6OZP39ZG8H)

GLYCERIN (UNII: PDC6A3C0OX)

EDETATE DISODIUM (UNII: 7FLD91C86K)

SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JNI2J)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

TARTARIC ACID (UNII: W4888I119H)

FRANKINCENSE (UNII: R9XLF1R1WM)

Packaging								
#	Item Code	Package Description	Marketing Start Date	Marketing End Date				
1	NDC:84992-004- 01	1 in 1 BAG; Type 0: Not a Combination Product	03/13/2025					
2	NDC:84992-004- 02	10 in 1 BOX; Type 0: Not a Combination Product	03/13/2025					
3	NDC:84992-004- 03	18 in 1 BOX; Type 0: Not a Combination Product	03/13/2025					

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC Monograph Drug	M017	03/13/2025				

Labeler - ADP Health Limited (101747972)

Revised: 3/2025 ADP Health Limited