KOCOWO LIDOCAINE PAIN RELIEF GEL-PATCH- lidocaine pain relief gel-patch patch
Henan Enokon Medical Instrument Co., Ltd.

83559-010

Active Ingredient

Lidocaine 4%

Purpose

Topical anesthetic

Use

Temporarily relieves muscle soreness and minor joint pains in the wrist, knees, back, neck, hips, shoulders, elbows.

Warnings

For external use only

Do not use

on wounds or damaged skin
with a heating pad or device
with other ointments, creams, sprays, or liniments
if you are allergic to any ingredients of this product

When Using

use only as directed
avoid contact with the eyes, mucous membranes or rashes
do not bandage tightly

Stop Use

Stop use and ask a doctor if
skin reactions such as redness, swelling, blistering or other discomfort occur.
symptoms persist for more than 7 days

Ask Doctor

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

Children under age of 12: Consult a doctor

Keep Out Of Reach Of Children

If swallowed accidentally, get medical help or contact with Poison Control Center immediately.

Directions

Adults and Children above 12 years old:
Clean and dry the affected area.
Tear off the protective film and apply the exposed part of the patch to the the affected area.
Carefully remove remaining film while pressing the patch firmly on the skin.
Remove patch from the skin after at most 8-hour application.

Inactive ingredients

Glycerin,Sodium polyacrylate,Polyvinyl pyrrolidone,Phenoxyethanol,Tartaric Acid,Ethyl ethyl glycerol,Sodium carboxymethyl cellulose,Aluminum Glycinate,Styrene acrylic copolymer emulsion,EDTA,Titanium Dioxide,Polysorbate 80,Sorbitol,Purified Water

PRINCIPAL DISPLAY PANEL

KOCOWO LIDOCAINE PAIN RELIEF GEL-PATCH

lidocaine pain relief gel-patch patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83559-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	4 g in 100

Inactive Ingredients	
Ingredient Name	**Strength**
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	

GLYCERIN (UNII: PDC6A3C0OX)	
ALUMINUM GLYCINATE (UNII: 1K713C615K)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
EDTA (UNII: 9G34HU7RV0)	
WATER (UNII: 059QF0KO0R)	
SODIUM ACRYLATE (UNII: 7C98FKB43H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
STYRENE/ACRYLAMIDE COPOLYMER (MW 500000) (UNII: 5Z4DPO246A)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SORBITOL (UNII: 506T60A25R)	
TARTARIC ACID (UNII: W4888I119H)	
POVIDONE (UNII: FZ989GH94E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83559-010-01	32 in 1 BOX; Type 0: Not a Combination Product	08/04/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/04/2025	

Labeler - Henan Enokon Medical Instrument Co., Ltd. (701730676)

Establishment

Name	Address	ID/FEI	Business Operations
Henan Enokon Medical Instrument Co., Ltd.		701730676	manufacture(83559-010)

Revised: 8/2025 Henan Enokon Medical Instrument Co., Ltd.