LEGERE LAVENDAR FOOT PATCH- topical starch patch Myriad 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.

Légère lavender

PURPOSE

Skin Protectant

Uses

May help relieve minor aches and pain

Active Ingredient

Warnings

For external use only / Don't eat or swallow it

Warnings

Keep out of the reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

When using this product

- Do not use otherwise than directed.
- Do not stretch out the adhesive tape.
- Do not take out the contents from the patch.
- If the patch has been ripped, discard it immediately.

Stop use and ask a doctor if

- If allergic reaction or irritation occurs
- If you are pregnant, nursing or under medial treatment

Warnings

■ On wounds, damaged skin, or face.

- If you are allergic to any ingredients of this product.
- With, or at the same time as, other external products.

Inactive Ingredients

Bamboo vinegar, Tourmaline, Eucalyptus oil, Mugwort extract, Loquat extract, Vitamin C, Highly purified silica, Perlite, Polyhydric alcohol

directions

Adults and children 4 years of age and over:

Step 1: Clean and dry area for the patch to be attached.

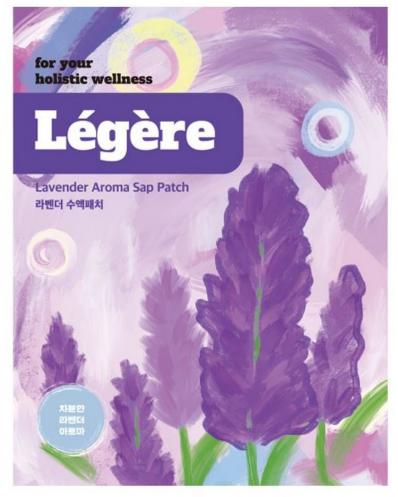
Step 2: Remove the protective cover sheet and gently apply.

Step 3: Leave it on at least 4hours before remove.

Children under 4 years of age: Consult with a doctor.

Légère lavender Foot Patch

82504-104-01



Drug Facts		
Active Ingredient		Purpose
Topical Starch	31.92%	Skin Protectant
Uses		
■ May help relieve minor a	aches and pain	
Warnings		
For external use only / Do	n't eat or swallow it	
Do Not Use		
■ On wounds, damaged sk	kin, or face.	
■ If you are allergic to any	ingredients of this produc	rt.
■ With, or at the same tim	e as, other external produ	cts.
When using this product		
■ Do not use otherwise th	an directed.	
■ Do not stretch out the a	dhesive tape.	
■ Do not take out the con	tents from the patch.	
■ If the patch has been rip	pped, discard it immediatel	y.
Stop use and ask a doctor	if	
■ If allergic reaction or irrit	tation occurs	
If you are pregnant, nurs	sing or under medial treatr	ment
Keep out of the reach o	f children. If swallowed,	get medical help of
contact a Poison Control Ce	enter right away.	
Directions		
Adults and children 4 year	s of age and over:	
Step 1: Clean and dry area f	for the patch to be attache	ed.
Step 2: Remove the protecti	ive cover sheet and gently	apply.
Step 3: Leave it on at least	4hours before remove.	
Children under 4years of a	ge: Consult with a doctor.	
Storage		
■ Avoid storing product in	the direct sunlight.	
■ Keep it in a dry and coo	l place.	
Inactive Ingredients		
Lavender oil, Tourmaline, V	Wood vinegar, Eucalyptus	oil, Mugwort extrac

Loquat extract, Vitamin C, Highly purified silica, Perlite, Polyhydric alcohol

LEGERE LAVENDAR FOOT PATCH

topical starch patch

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82504-104
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
STARCH, CORN (UNII: 08232NY3SJ) (STARCH, CORN - UNII:08232NY3SJ)	STARCH, CORN	31.92 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
LAVENDER OIL (UNII: ZBP1YXW0H8)		
PYROLIGNEOUS ACID (UNII: N4G9GAT76C)		
ERIOBOTRYA JAPONICA LEAF (UNII: Z02066SV11)		
PERLITE (UNII: 0SG101ZGK9)		
ASCORBIC ACID (UNII: PQ6CK8PD0R)		
EUCALYPTUS OIL (UNII: 2R040NI662)		
POLIGLUSAM (UNII: 82LKS4QV2Y)		
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)		
ARTEMISIA PRINCEPS LEAF (UNII: SY077EW02G)		
SILICON DIOXIDE (UNII: ETJ7Z 6XBU4)		
GLYCERYL CAPRYLATE (UNII: TM2TZ D4G4A)		
SCHORL TOURMALINE (UNII: 17308XLY6T)		
SORBITOL (UNII: 506T60A25R)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82504-104- 01	40 g in 1 BOX; Type 0: Not a Combination Product	01/10/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/10/2022	

Labeler - Myriad Co., Ltd. (695004202)

Registrant - Myriad Co., Ltd. (695004202)

Establishment				
Name	Address	ID/FEI	Business Operations	
Myriad Co., Ltd.		695004202	manufacture(82504-104)	

Revised: 12/2022 Myriad Co., Ltd.