

LEGERE BAMBOO 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 bamboo

PURPOSE

Skin Protectant

Uses

May help relieve minor aches and pain

Active Ingredient

Topical Starch 32.33%

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 medical 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 4years of age: Consult with a doctor.

Légère Bamboo Foot Patch

82504-102-01



Drug Facts	
Active Ingredient	Purpose
Topical Starch	32.33% Skin Protectant
Uses	
<ul style="list-style-type: none"> May help relieve minor aches and pain 	
Warnings	
For external use only / Don't eat or swallow it	
Do Not Use	
<ul style="list-style-type: none"> 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. 	
When using this product	
<ul style="list-style-type: none"> 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	
<ul style="list-style-type: none"> If allergic reaction or irritation occurs If you are pregnant, nursing or under medical treatment 	
Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
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 4years of age: Consult with a doctor.	
Storage	
<ul style="list-style-type: none"> Avoid storing product in the direct sunlight. Keep it in a dry and cool place. 	
Inactive Ingredients	
Bamboo vinegar, Tourmaline, Eucalyptus oil, Mugwort extract, Loquat extract, Vitamin C, Highly purified silica, Perlite, Polyhydric alcohol	

LEGERE BAMBOO FOOT PATCH

topical starch patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82504-102
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
STARCH, CORN (UNII: O8232NY3SJ) (STARCH, CORN - UNII:O8232NY3SJ)	STARCH, CORN	32.33 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
PHYLLOSTACHYS EDULIS VINEGAR (UNII: MR94DK8ZSM)	
PERLITE (UNII: 0SG101ZGK9)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
POLIGLUSAM (UNII: 82LKS4QV2Y)	

ARTEMISIA PRINCEPS LEAF (UNII: SY077EW02G)	
ERIOBOTRYA JAPONICA LEAF (UNII: Z02066SV11)	
GLYCERYL CAPRYLATE (UNII: TM2TZD4G4A)	
SCHORL TOURMALINE (UNII: 17308XLY6T)	
SORBITOL (UNII: 506T60A25R)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82504-102-01	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(82504-102)

Revised: 12/2022

Myriad Co., Ltd.