HARUTO HANGOVER DEFENCEPATCH- glutathione patch
YOUNGWOO 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.

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glutathione

N-acethylcysteine
vitamin b1 (Thiamine HCl)
vitamin b5 (D-panthenol)
vitamin b6 (Pyridoxine HCl)
vitamin b9 (folic acid)
vitamin b12 0.1% (cobalamin)
Milk Thistle extract

Hangover relief

Keep out of reach of the children

Stick the patch to target areas of concern and gently press down on the patch

1. Do not use in the following cases(Eczema and scalp wounds)

2.Side Effects

1)Due to the use of this druf if rash, irritation, itching and symptopms of hypersnesitivity occur dicontinue use and consult your phamacisr or doctor

3.General Precautions

1)If in contact with the eyes, wash out thoroughty with water If the symptoms are servere, seek medical advice immediately

2)This product is for exeternal use only. Do not use for internal use

4.Storage and handling precautions

1)If possible, avoid direct sunlight and store in cool and area of low humidity

2)In order to maintain the quality of the product and avoid misuse

3)Avoid placing the product near fire and store out in reach of children

for external use only

HARUTO HANGOVER DEFENCE PATCH

	Contents (%)
SIS Rubber	98.5
Glutathione	1.422
N-acethylcysteine	0.015
vitamin b1 (Thiamine HCl)	0.015
vitamin b5 (D-panthenol)	0.015
vitamin b6 (Pyridoxine HCl)	0.015
vitamin b9 (folic acid)	0.0015
vitamin b12 0.1% (cobalamin)	0.0015
Milk Thistle extract	0.015

HARUTO HANGOVER DEFENCEPATCH

glutathione patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80094-0001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GLUTATHIONE (UNII: GAN16C9B8O) (GLUTATHIONE - UNII:GAN16C9B8O)	GLUTATHIONE	1.422 mg in 100 mg

Inactive Ingredients	
Ingredient Name	**Strength**
THIAMINE HYDROCHLORIDE (UNII: M572600E5P)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:80094-0001-1	3 mg in 1 PATCH; Type 0: Not a Combination Product	10/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/01/2020	

Labeler - YOUNGWOO CO.,LTD (689507973)

Registrant - YOUNGWOO CO.,LTD (689507973)

Establishment

Name	Address	ID/FEI	Business Operations
YOUNGWOO CO.,LTD		689507973	manufacture(80094-0001)

Revised: 1/2022 YOUNGWOO CO.,LTD