## NEURORESCUEX HH333- scutellaria baicalensis root, coptis japonica root, phellodendron chinensis bark, gardenia jasminoides fruit, rheum palmatum root capsule

Panacura Inc.

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.

### 84074-001 NeuroRescueX HH333

■ For adults, take 2 capsules, once a day, ■ before or between meals

Scutellaria Baicalensis Root (240 mg in 400 mg)

Coptis Japonica Root (240 mg in 400 mg)

Phellodendron Chinensis Bark (240 mg in 400 mg)

Gardenia Jasminoides Fruit (240 mg in 400 mg)

Rheum Palmatum Root (60 mg in 400 mg)

prevention of cerebrovascular diseases, cardiovascular disease

Do not use if you have

- galactose intolerance
- lapp lactase deficiency
- glucose-galactose malabsorption (this medicine contains lactose)

Ask a doctor before use if you are

a patient with liver dysfunction (liver enzyme value (AST, ALT) levels more than 3 times normal)

a patient who is being treated by a doctor or taking other medications.

an elderly person (in general, elderly people should take with care, such as reducing their dosage, as their physiological function has deteriorated.)

a child (the safety of children's use is not established. (Less experience in use))

- prevention of cerebrovascular diseases, cardiovascular disease
- improvement of vascular endothelial cell function
- adjunctive use to control hypertension, hyperlipidemia
- weight loss support

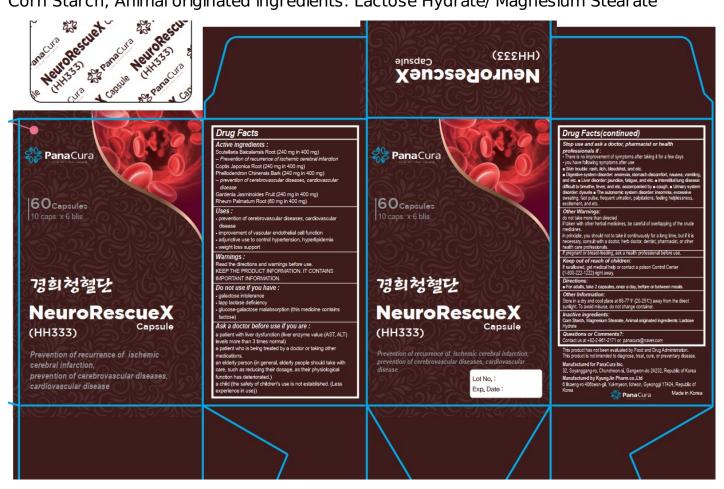
Read the directions and warnings before use.

REEP THE PRODUCT INFORMATION. IT CONTAINS IMPORTANT INFORMATION.
Stop use and ask a doctor, pharmacist or health professionals if  There is no improvement of symptoms after taking it for a few days
☐ you have following symptoms after use
■ Skin trouble: ■ rash, ■ itch, ■ bloodshot, and etc.
■ Digestive system disorder: ■ anorexia, ■stomach discomfort, ■nausea, ■vomiting
and etc.
■ Liver disorder: ■ jaundice, ■ fatigue, and etc.
■ Interstitial lung disease: ■ difficult to breathe, ■ fever, and etc. accompanied by ■

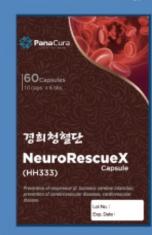
- cough. ■ Urinary system disorder: ■ dysuria
- $\blacksquare$  The autonomic system disorder:  $\blacksquare$  insomnia,  $\blacksquare$  excessive sweating,  $\blacksquare$  fast pulse,

 $\blacksquare$  frequent urination,  $\blacksquare$  palpitations,  $\blacksquare$  feeling helplessness,  $\blacksquare$  excitement, and etc. Keep out of reach of the children.

Corn Starch, Animal originated ingredients: Lactose Hydrate/ Magnesium Stearate









# NEURORESCUE X

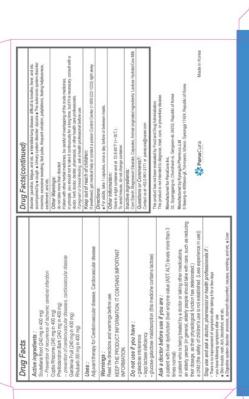
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Drug Facts (continued)

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# **NEURORESCUE X**





**NEURORESCUE X** 

Lot No.: Exp. Date:

## **NEURORESCUEX HH333**

scutellaria baicalensis root, coptis japonica root, phellodendron chinensis bark, gardenia jasminoides fruit, rheum palmatum root capsule

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84074-001	
Route of Administration	ORAL			

Active Ingredient/Active Moiety				
Ingredient Name	<b>Basis of Strength</b>	Strength		
SCUTELLARIA BAICALENSIS ROOT (UNII: 7J95K7ID2S) (SCUTELLARIA BAICALENSIS ROOT - UNII:7J95K7ID2S)	SCUTELLARIA BAICALENSIS ROOT	240 mg		
COPTIS JAPONICA ROOT (UNII: JH0SWT16D1) (COPTIS JAPONICA ROOT - UNII:JH0SWT16D1)	COPTIS JAPONICA ROOT	240 mg		
PHELLODENDRON CHINENSIS BARK (UNII: 2866QMZ434) (PHELLODENDRON CHINENSIS BARK - UNII: 2866QMZ434)	PHELLODENDRON CHINENSIS BARK	240 mg		
<b>GARDENIA JASMINOIDES FRUIT</b> (UNII: 7CTH8MD549) (GARDENIA JASMINOIDES FRUIT - UNII:7CTH8MD549)	GARDENIA JASMINOIDES FRUIT	240 mg		
RHEUM PALMATUM ROOT (UNII: G025DAL7CE) (RHEUM PALMATUM ROOT - UNII:G025DAL7CE)	RHEUM PALMATUM ROOT	60 mg		

Inactive Ingredients				
Ingredient Name	Strength			
STARCH, CORN (UNII: 08232NY3SJ)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)				

Product Characteristics			
Color	yellow	Score	no score
Shape	CAPSULE	Size	21mm
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84074- 001-03	2 in 1 CARTON	12/08/2025	
1	NDC:84074- 001-02	6 in 1 CARTON		
1	NDC:84074- 001-01	10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:84074- 001-05	3 in 1 CARTON	12/08/2025	
2	NDC:84074- 001-04	10 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/05/2024	

# Labeler - Panacura Inc. (695390655)

## Registrant - Panacura Inc. (695390655)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
KyungJin Pharm.co.,Ltd		695542024	manufacture(84074-001)	

Revised: 12/2025 Panacura Inc.