LAXATIVES RUBILAX- psyllium huskpowder granule Lydia 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.

Drug Facts

psyllium husk powder, sennae fructus powder

lactose monohydrate, acacia, corn starch, carboxymethylcellulose calcium, l-menthol

constipation

relieving the following symptoms of constipation: loss of appetite, abdominal distension, intestinal fermentation, hemorrhoids

keep out of reach of the children

adults take intake does of 3.5g twice a day morning and evening on an empty stomach, however, taking the minimum amount of the first time and watching the shape and condition of the stomach gradually increases or decreases

do not administer to the following patients,

the product contains lactose, patients with rare hereditary problems of galactose intolerance, the lapp deficiency or glucose-galactose malabsorption should not take this medicine

for oral use only

 [Active ingredient] Per intake(3.5g) Psyllium Husk Powder(BP)	 [Warnings] 1. Do not administer to the following patients, The product contains lactose, Patients with rare hereditary problems of galactose intolerance, the Lapp lactase deficiency or glucose–galactose malabsorption should not take this medicine 2. Do not take this medicine if you have following symptoms Patients with acute abdominal disease (appendicitis, intestinal bleeding, ulcerative colitis, etc.) patients with intestinal obstruction Young children and infants under the age of 3 Other warnings, please refer to attached document, [Storage] Store in an airtight container in room temperature(1~30°C)
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LAXATIVES RUBILAX psyllium huskpowder granule				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72988-0010	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				

		Ingredient Name		Basis of Strength	Strength	
PSYLLIUM HUSK (UNII: 0SH053407G) (PSYLLIUM HUSK - UNII:0SH053407G) PSYLLIUM HUSK					2168 mg in 3.5 g	
SENNA PODS (UNII: S8SJ19N2NX) (SENNA PODS - UNII:S8SJ19N2NX) SENNA PODS				SENNA PODS	496 mg in 3.5 g	
In	active Ingre	dients				
		Ingredient Name			Strength	
LE	VOMENTHOL (U	NII: BZ1R15MTK7)				
LA	СТОЅЕ МОНОН	YDRATE (UNII: EWQ57Q8I5X)				
Packaging						
#	ltem Code	Package Description	Mar	keting Start M Date	arketing End Date	
1	NDC:72988- 0010-2	30 in 1 POUCH	04/22/2	2019		
1	NDC:72988- 0010-1	3.5 g in 1 POUCH; Type 0: Not a Combination Product				
Marketing Information						
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	Marketing Category	Application Number or Monograph Citation	Ma	arketing Start N Date	Aarketing End Date	
	approved drug ner		04/22	2/2019		

Labeler - Lydia Co., Ltd. (695735569)

Registrant - Lydia Co., Ltd. (695735569)

Establishment

Name	Address	ID/FEI	Business Operations
I World Pharmaceutical Co., Ltd.		688222857	manufacture(72988-0010)

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Lydia Co., Ltd.		695735569	label(72988-0010)

Revised: 12/2023

Lydia Co., Ltd.