FESTAL PLUS- cellulase ap3 , pancreatin, simethicone tablet 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.

Cellulase AP3

Pancreatin

Simethicone

digestion

Keep out of reach of children

- children under 7 years of age: consult a doctor
- take 1-2 tablet as symptoms accur, as directed by a doctor

Ask a doctor before use if you have taking a prescription drug. Antacids may interact with certain prescription drugs.

When using this product

- Do not take more than 4 tablet in 24 hours
- Do not use the maximum dosage for more than 2 weeks

Keep out of reach of children

Dextrin, stearic acid, talc, polyethylene glycol 6000

for oral use

FESTAL PLUS

Drug FactsActive Ingredients (in one tablet)PurposeCellulase AP3 10mgDigestionPancreatin 315mgDigestionSimethicone 30mgDigestionUses

digestion

Warnings

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Directions

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Other Information

■ Store at room temperature

Inactive Ingredient

Dextrin, stearic acid, talc, polyethylene glycol 6000

Distributed By: Bio Mission Group,Inc.

9925 Painter Ave #O, Whittier, CA 90605 USA

Made in South Korea

cellulase ap3, pancreatin, simethicone tablet

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Product Type HUMA	N OTC DRUG Item Cod	e (Source) ND	C:72988-0038
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Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PANCRELIPASE (UNII: FQ3DRG0N5K) (PANCRELIPASE LIPASE - UNII:8MYC339320)	PANCRELIPASE LIPASE	315 mg
DIMETHICONE (UNII: 92RU3N3Y10) (DIMETHICONE - UNII:92RU3N3Y10)	DIMETHICONE	30 ma

Inactive Ingredients

Ingredient Name	Strength

STEARIC ACID (UNII: 4ELV7Z65AP)

Product Characteristics

Color	white	Score	no score
Shape	OVAL	Size	14mm
Flavor		Imprint Code	PLUS
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72988- 0038-1	10 in 1 PACKAGE; Type 0: Not a Combination Product	09/01/2024	

Marketing Information

Marketing Application Number or Monogo Category Citation		Marketing Start Date	Marketing End Date
unapproved drug other		09/01/2024	

Labeler - Lydia Co., Ltd. (695735569)

Registrant - Lydia Co., Ltd. (695735569)

Establishment

Name	Address	ID/FEI	Business Operations	
Lydia Co., Ltd.		695735569	manufacture(72988-0038)	

Revised: 9/2024 Lydia Co., Ltd.