ANTI-H.PYLORI FORMULA- reishi yeast fu ling ophiopogon japonicus whole capsule
Bispit Canada 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.

85631-016-01ANTI-H.PYLORI FORMULA

Active ingredients: Reishi, Yeast, Fu lingOphiopogon japonicus whole.

Inactive ingredient: Maltodextrin, Heavy calciumcarbonate, Povidone K30.

Storage: Store in a tightly sealed container in acool, dry place,Keep out of children

INHIBITING BACTERIAL GROWTH
AND PROMOTING METABOLIC EXCRETION

1. This product can be taken for an extended periodof time, but it is not intended to substitutemedication. Seek medical advice promptly for severecases.
2. Avoid spicy and irritating foods for the first 3 daysof administering this product.3. Pregnant women and individuals with allergiesshould use this product only under the guidance of aphysician.

Dosage: 3 capsules each time, twice a day(morning and evening)

Avoid spicy and irritating foods for the first 3 daysf administering this product.

ANTI-H.PYLORI FORMULA

reishi yeast fu ling ophiopogon japonicus whole capsule

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:85631-016
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OPHIOPOGON JAPONICUS WHOLE (UNII: 2983RA66TS) (OPHIOPOGON JAPONICUS WHOLE - UNII:2983RA66TS)	OPHIOPOGON JAPONICUS WHOLE	3.5 g in 100 g
YEAST (UNII: 3NY3SM6B8U) (YEAST - UNII:3NY3SM6B8U)	YEAST	6 g in 100 g
FU LING (UNII: XH37TWY5O4) (FU LING - UNII:XH37TWY5O4)	FU LING	3 g in 100 g

REISHI (UNII: TKD8LH0X2Z) (REISHI - UNII:TKD8LH0X2Z)	REISHI	50 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
POVIDONE K30 (UNII: U725QWY32X)	

Product Characteristics

Color	blue	**Score**	3 pieces
Shape	DOUBLE CIRCLE	**Size**	12mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85631-016-01	90 in 1 BOTTLE	12/22/2025	
1		0.8 g in 1 CAPSULE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		12/22/2025	

Labeler - Bispit Canada Ltd. (243332192)

Registrant - Bispit Canada Ltd. (243332192)

Establishment

Name	Address	ID/FEI	Business Operations
Bispit Canada Ltd.		243332192	manufacture(85631-016)

Revised: 12/2025 Bispit Canada Ltd.