GOUT RELIEF 3310-S- medicago sativa whole chicory root capsule
Bispit Canada Ltd.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

85631-006-01-Gout Relief 3310-S-NEW

MEDICAGO SATIVA WHOLE 20.00%
CHICORY ROOT 23.00%
GLUCOSAMINE HYDROCHLORIDE 10.00%
BUCKWHEAT 7.00%
TEA LEAF 5.00%
MORUS ALBA LEAF 13.00%
LICORICE 0.20%
FU LING 2.00%
COIX LACRYMA-JOBI SEED 8.00%

This product has a significanteffect on gout, crystal deposits,and hyperuricemia; elt has an auxiliary effectof mitigating kidney stonesand bladder stones

Keep out of reach of children

The product has great somatosensation after taking it atleast 21 days in the earty stage;3. Elevated uric acid or gout recurrence and site changes arenoral in the earty stage of taking this product;

Dissolved CrystallizationKidney StoneGallstone< Urinary Calculus

Do not use if seal is broken.

Do not eat freshness packet enclosed.

Keep out of direct sunlight, high temperature, and humidity.

Store in a cool, dry place.

For your health, please read the followingcarefully
1. The use of three vials can effectivety dissolve crystallization (ittakes time to dissolve crystallization);2. The product has great somatosensation after taking it atleast 21 days in the early stage;3. Elevated uric acid or gout recurrence and site changes arenormal in the early stage of taking this product;4. Try not to drink alcohol or drink less 15 days before takingthis product;
5. This product certainly has significant effect on gout,crystallization and hyperuricemia;

This product has a significanteffect on gout, crystal deposits,and hyperuricemia; elt has an auxiliary effectof mitigating kidney stonesand bladder stones

GOUT RELIEF 3310-S

medicago sativa whole chicory root capsule

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:85631-006
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
COIX LACRYMA-JOBI SEED (UNII: 4Q4V6NTZ1F) (COIX LACRYMA-JOBI SEED - UNII:4Q4V6NTZ1F)	COIX LACRYMA-JOBI SEED	104 mg in 1.3 g
MEDICAGO SATIVA WHOLE (UNII: DJO934BRBD) (MEDICAGO SATIVA WHOLE - UNII:DJO934BRBD)	MEDICAGO SATIVA WHOLE	260 mg in 1.3 g
BUCKWHEAT (UNII: N0Y68724P3) (BUCKWHEAT - UNII:N0Y68724P3)	BUCKWHEAT	91 mg

BUCKWHEAT (UNII: N0T68724R3) (BUCKWHEAT - UNII:N0T68724R3)	BUCKWHEAT	in 1.3 g
TEA LEAF (UNII: GH42T47V24) (TEA LEAF - UNII:GH42T47V24)	TEA LEAF	65 mg in 1.3 g
MORUS ALBA LEAF (UNII: M8YIA49Q2P) (MORUS ALBA LEAF - UNII:M8YIA49Q2P)	MORUS ALBA LEAF	169 mg in 1.3 g
CHICORY ROOT (UNII: 090CTY533N) (CHICORY ROOT - UNII:090CTY533N)	CHICORY ROOT	299 mg in 1.3 g
FU LING (UNII: XH37TWY5O4) (FU LING - UNII:XH37TWY5O4)	FU LING	26 mg in 1.3 g
GLUCOSAMINE HYDROCHLORIDE (UNII: 750W5330FY) (GLUCOSAMINE - UNII:N08U5BOQ1K)	GLUCOSAMINE HYDROCHLORIDE	130 mg in 1.3 g
LICORICE (UNII: 61ZBX54883) (LICORICE - UNII:61ZBX54883)	LICORICE	2.6 mg in 1.3 g

Inactive Ingredients

Ingredient Name	Strength
MALTOSYL-ISOMALTOTETRAOSE (UNII: B9HR7A9UFR)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	

Product Characteristics

Color	white	**Score**	score with uneven pieces
Shape	CAPSULE	**Size**	20mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85631-006-01	60 in 1 BOTTLE	09/01/2025	
1		1.3 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 homeopathic		09/01/2025	

Labeler - Bispit Canada Ltd. (243332192)

Registrant - Bispit Canada Ltd. (243332192)