

GOUT RELIEF 3310-S- celery seed, medicago sativa whole, chicory root capsule
Bispi 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.

CELERY SEED 1G1EAA320L 20.00%
MEDICAGO SATIVA WHOLE DJO934BRBD 20.00%
CHICORY ROOT 090CTY533N 18.00%
GLUCOSAMINE HYDROCHLORIDE 750W5330FY 10.00%
BUCKWHEAT N0Y68724R3 7.00%
TEA LEAF GH42T47V24 5.00%
PUERARIN Z9W8997416 5.00%
MORUS ALBA LEAF M8YIA49Q2P 3.00%
LICORICE 61ZBX54883 0.20%
SODIUM CHLORIDE 451W47IQ8X 2.00%

This product has a significant effect on gout, crystal deposits, and hyperuricemia; it has an auxiliary effect of mitigating kidney stones and bladder stones

Keep out of reach of children

The product has great somatosensation after taking it at least 21 days in the early stage; 3. Elevated uric acid or gout recurrence and site changes are normal in the early stage of taking this product;

Dissolved Crystallization Kidney Stone Gallstone < Urinary Calculus

For your health, please read the following carefully

1. The use of three vials can effectively dissolve crystallization (it takes time to dissolve crystallization);
2. The product has great somatosensation after taking it at least 21 days in the early stage;
3. Elevated uric acid or gout recurrence and site changes are normal in the early stage of taking this product;
4. Try not to drink alcohol or drink less 15 days before taking this product;
5. This product certainly has significant effect on gout, crystallization and hyperuricemia;

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.

MICROCRYSTALLINE CELLULOSE
MALTOSYL-ISOMALTOTETRAOSE
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED
MAGNESIUM STEARATE



GOUT RELIEF 3310-S

celery seed, medicago sativa whole, chicory root capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85631-003
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GLUCOSAMINE HYDROCHLORIDE (UNII: 750W5330FY) (GLUCOSAMINE - UNII:N08U5BOQ1K)	GLUCOSAMINE HYDROCHLORIDE	130 mg in 1.3 g
MORUS ALBA LEAF (UNII: M8YIA49Q2P) (MORUS ALBA LEAF - UNII:M8YIA49Q2P)	MORUS ALBA LEAF	39 mg in 1.3 g
SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION -	SODIUM CHLORIDE	26 mg

UNII:LYR4M0NH37)	SODIUM CHLORIDE	in 1.3 g
MEDICAGO SATIVA WHOLE (UNII: DJO934BRBD) (MEDICAGO SATIVA WHOLE - UNII:DJO934BRBD)	MEDICAGO SATIVA WHOLE	260 mg in 1.3 g
TEA LEAF (UNII: GH42T47V24) (TEA LEAF - UNII:GH42T47V24)	TEA LEAF	65 mg in 1.3 g
CELERY SEED (UNII: 1G1EAA320L) (CELERY SEED - UNII:1G1EAA320L)	CELERY SEED	260 mg in 1.3 g
CHICORY ROOT (UNII: 090CTY533N) (CHICORY ROOT - UNII:090CTY533N)	CHICORY ROOT	234 mg in 1.3 g
PUERARIN (UNII: Z9W8997416) (PUERARIN - UNII:Z9W8997416)	PUERARIN	65 mg in 1.3 g
BUCKWHEAT (UNII: N0Y68724R3) (BUCKWHEAT - UNII:N0Y68724R3)	BUCKWHEAT	91 mg in 1.3 g
LICORICE (UNII: 61ZBX54883) (LICORICE - UNII:61ZBX54883)	LICORICE	2.6 mg in 1.3 g

Inactive Ingredients

Ingredient Name	Strength
CELLULOSE GUM (UNII: K679OBS311)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
MALTOSYL-ISOMALTOTETRAOSE (UNII: B9HR7A9UFR)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

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:85631-003-01	60 in 1 BOX	07/28/2025	
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		07/28/2025	

Labeler - Bispit Canada Ltd. (243332192)

Registrant - Bispit Canada Ltd. (243332192)

