GOUT RELIEF 3310A PLUS- gout relief 3310a plus tablet
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.

MEDICAGO SATIVA WHOLE
CHICORY ROOT
GLUCOSAMINE HYDROCHLORIDE
BUCKWHEAT
TEA LEAF
MORUS ALBA LEAF
LICORICE
FU LING
COIX LACRYMA-JOBI SEED
ANSERINE

CALCIUM CARBONATE
SODIUM CHLORIDE

Ulsing three bottles is generally effective fordissolving crystals (as crystal dissoiution takestime);
You should experience noticeable effects afterat least 21 days of conbnlous use,It is noral to experence increose in uric acidlevels or gout recurrencelchanges imn affectedareas during the early days of use;Refrain from olcohol consumption, or limit itsionificantly, for at least 15 days before takingthis product;
lt has an auxiliary effect on improving libidoand supporting prostate health,
Do not use if you have a seafood allergy.
Store properly and out of reach of children.

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This product is a pure plant extract; it canreduce the frequency of goutattacks; dissolve the crystals in thejoints; promote the excretion of blood uricacids,andimprove the purine metabolismand excretory system.

2 tablets in the momingand 2 toblets in the evening.

Applicable Population:patient with gout,high uric acid, depositionof urate crystal, kidney stone and bladder
stone.

GOUT RELIEF 3310A PLUS

gout relief 3310a plus tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:85631-009
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LICORICE (UNII: 61ZBX54883) (LICORICE - UNII:61ZBX54883)	LICORICE	1 mg in 1 g
MEDICAGO SATIVA WHOLE (UNII: DJO934BRBD) (MEDICAGO SATIVA WHOLE - UNII:DJO934BRBD)	MEDICAGO SATIVA WHOLE	150 mg in 1 g
BUCKWHEAT (UNII: N0Y68724R3) (BUCKWHEAT - UNII:N0Y68724R3)	BUCKWHEAT	40 mg in 1 g
FU LING (UNII: YH37TWX5O4) (FU LING - UNII:YH37TWX5O4)	FU LING	10 mg

FU LING (UNII: XH371WT3O4) (FU LING - UNII:XH371WT3O4)	FU LING	in 1 g
CHICORY ROOT (UNII: 090CTY533N) (CHICORY ROOT - UNII:090CTY533N)	CHICORY ROOT	180 mg in 1 g
MORUS ALBA LEAF (UNII: M8YIA49Q2P) (MORUS ALBA LEAF - UNII:M8YIA49Q2P)	MORUS ALBA LEAF	75 mg in 1 g
ANSERINE (UNII: HDQ4N37UGV) (ANSERINE - UNII:HDQ4N37UGV)	ANSERINE	350 mg in 1 g
TEA LEAF (UNII: GH42T47V24) (TEA LEAF - UNII:GH42T47V24)	TEA LEAF	34 mg in 1 g
COIX LACRYMA-JOBI SEED (UNII: 4Q4V6NTZ1F) (COIX LACRYMA-JOBI SEED - UNII:4Q4V6NTZ1F)	COIX LACRYMA-JOBI SEED	40 mg in 1 g
GLUCOSAMINE HYDROCHLORIDE (UNII: 750W5330FY) (GLUCOSAMINE - UNII:N08U5BOQ1K)	GLUCOSAMINE HYDROCHLORIDE	70 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Product Characteristics

Color	brown	**Score**	score with uneven pieces
Shape	ROUND	**Size**	14mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85631-009-01	60 in 1 BOX	09/05/2025	
1		1 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		09/05/2025	

Labeler - Bispit Canada Ltd. (243332192)

Registrant - Bispit Canada Ltd. (243332192)

Revised: 9/2025 Bispit Canada Ltd.