Quercetin SAP

Science-based bioflavonoid for allergy and inflammation management

Quercetin is a bioflavonoid found in onions, apples, black tea, and grapefruit. Quercetin has antiallergic, antioxidant, and anti-inflammatory activity. Mast cells contribute to the inflammatory process in allergic reactions, where immunologic stimulation leads to degranulation of the mast cell as well as the generation of numerous cytokines and inflammatory mediators. Quercetin is an effective treatment option for allergies, as it has been shown to inhibit mast-cell secretions, and it has the ability to downregulate histamine production via inhibition of the enzyme histidine decarboxylase (HDC) mRNA from human mast cells. Quercetin also causes a decrease in the release of IL-6 and tryptase, and is known to inhibit phospholipase A_2 and 5-lipoxygenase, thereby reducing synthesis of leukotrienes and the series-2 prostaglandins via the arachidonic-acid pathway. The anti-inflammatory properties of quercetin were explored in a double-blind, placebo-controlled study on men with chronic prostatitis syndrome. The study found that after one month of treatment, 67% of men in the treatment group reported at least a 25% improvement in their pain levels.

Bromelain is a proteolytic enzyme derived from the stem of the pineapple plant. Bromelain has been found to be a mucolytic agent effective in respiratory disorders, and functions as an anti-inflammatory agent. Bromelain counteracts the fibrin and kinin pathways, which stimulate plasmin. Plasmin blocks endogenous arachidonic-acid mobilization and reduces prostaglandin synthesis, which reduces localized inflammation and edema. This is thought to be the mechanism by which bromelain helps reduce symptoms of allergic rhinitis. Bromelain may also enhance the effectiveness of quercetin.

ACTIVE INGREDIENTS

Each non-GMO vegetable capsule contains:

 Quercetin.
 500 mg

 Bromelain (Ananas comosus), 2400 GDU/g
 [3,600,000 FCC PU].
 100 mg

This product is non-GMO.

Contains no: Gluten, soy, wheat, corn, eggs, dairy, yeast, citrus, preservatives, artificial flavour or colour, starch, or sugar.

WARNING: Take caution for use in patients with pineapple allergies.

DIRECTIONS FOR USE

Adults: Take 1 capsule twice daily with food in divided doses or as directed by your healthcare practitioner. Consult a healthcare practitioner for use beyond 4 weeks.

INDICATIONS

Quercetin SAP:

- Is an effective treatment for mild/moderate allergic reactions. It has been shown to prevent histamine release and reduce systemic inflammation, and therefore is a good proactive therapy for allergy treatment and prevention.
- Can be used to treat chronic asthma symptoms.
- Has anti-inflammatory properties, and can be used to treat men with chronic pelvic pain syndrome/chronic prostatitis.

PURITY, CLEANLINESS, AND STABILITY

All ingredients listed for all **Quercetin SAP** lot numbers have been tested by a third-party laboratory for identity, potency, and purity.



Bioflavonoid / Bioflavonoide For allergy and inflammation management Pour le control des allergies et de l'inflammation All Ingredients have been tested by a third-arty laboratory Tous tes ingrédients ont été testés par un laboratorie externe pour l'indriti, la puissance et la purété

> NPN 80039352 60 capsules

Scientific Advisory Panel (SAP): adding nutraceutical research to achieve optimum health



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ALLERGIES AND MAST CELLS

The IgE receptor plays a key role in inducing an allergic reaction. When a mast cell is exposed to an antigen, it binds to the IgE receptor and causes degranulation, leukotriene and prostaglandin secretion, and cytokine production.^[1] While mast cells are found throughout the body, they are found in high concentrations in connective tissue of the respiratory tract and nasal mucosa, and in blood vessels.^[2] Degranulation of mast cells accounts for almost 50% of symptoms of nasal rhinitis through the release of histamine.^[2] This degranulation also leads to symptoms that can include nasal congestion, sneezing, itching, inflammation, conjunctivitis, hives, throat irritation, vasodilation, increased capillary permeability, and smooth-muscle contraction. Allergy symptoms can be caused by a variety of sources such as environmental sources, foods, medications, and metabolic conditions. Studies on allergy sufferers demonstrate that during symptoms, there is a decline in the ability to perform daily activities and a measurable decline in physical and mental health status.^[2] Typical treatments of allergies include antihistamines, decongestants, anticholinergic agents, and corticosteroid drug therapy, alone or in combination; these can lead to adverse side effects including sedation, impaired learning/ memory, and cardiac arrhythmias.^[2]

QUERCETIN FOR ALLERGIES

Quercetin is an effective treatment option for allergies, as it has been shown to inhibit mast-cell secretion and has the ability to downregulate histidine decarboxvlase (HDC) mRNA from human mast cells.^[3] Quercetin inhibits the inflammatory process that is regulated by an increased release in neutrophils. The release of neutrophils leads to the destabilization of mast cells and their subsequent release of histamine and leukotrienes.^[2] By preventing the release of neutrophils, quercetin is therefore able to add stability to the mast-cell membrane, to help prevent degranulation and subsequent allergy symptoms. A placebo-controlled study performed by Hirano et al. (2009), in which subjects received 100 mg/d quercetin, demonstrated a reduction in allergic symptoms including congestion, lacrimation and itching, and a reduction in use of other allergy medications when compared to controls.[4]

QUERCETIN FOR ASTHMA

In a mouse model, quercetin was dosed between 3 and 10 mg/ kg and worked in a dose-dependent manner to inhibit eosinophil recruitment to the bronchoalveolar lavage fluid, and also significantly reduced both IL5 and IL4 levels. Quercetin also reduced P-selectin expression and the mucus production in the lung.^[5] These findings suggest the potential of therapeutic treatment of inflammatory airway diseases using quercetin.^[5]

OUERCETIN FOR CHRONIC PROSTATITIS

In a double-blind, placebo-controlled study by Shoskes et al. (1999), male subjects with chronic pelvic pain were randomly assigned to either placebo group or to receive 500 mg of quercetin twice a day.^[6] Pain was then evaluated before the study commenced and after 1 month, using the NIH chronic prostatitis symptom score. 67% of patients taking quercetin had a statistically significant improvement in symptoms of at least 25%.^[6] An unblinded followup arm of this study was also performed where participants received guercetin with bromelain and papain for one month, and 82% of these patients demonstrated an improvement in symptoms of at least 25%.[6]

Research Monograph

A separate randomized, double-blind, placebo-controlled trial was also performed to test the efficacy of quercetin in men with chronic prostatitis (CP). This study assessed 30 men with CP to receive quercetin, 500 mg twice daily, or the placebo pill for 1 month.^[7] Significant change in the NIH-CPSI score was observed in the quercetin group (p = 0.003) versus the placebo group, who had an insignificant mean improvement in the NIH-CPSI score.[7] These 2 trials indicate that quercetin is a viable treatment option for men suffering from chronic prostatitis.

BROMELAIN

Bromelain has been found to be a mucolytic agent in respiratory disorders, and works as an anti-inflammatory. Bromelain counteracts the fibrin and kinin pathways, which stimulates plasmin. Plasmin blocks endogenous arachidonic acid by prostaglandins, which reduces localized edema.^[2] Bromelain also has a long-term effect of inhibiting the firm adhesion of leukocytes to blood vessels at the site of inflammation, which also reduces edema.^[9] A study performed by Secor et al. (2005) dosed mice with 2 g/kg or 6 g/kg of bromelain twice a day for 4 d and resulted in attenuated development of allergic airway disease while altering CD4⁺ to CD8⁺ T-lymphocyte populations.^[8] The reduction in allergic airway disease outcomes suggests that bromelain may have similar effects in the treatment of human asthma and hypersensitivity disorders.^[9] Bromelain has also shown the ability to enhance the absorption and therefore the effectiveness of quercetin.^[2, 6]

SAFETY

Both bromelain and quercetin have an excellent safety profile, with minimal toxicity pictures. Bromelain demonstrates very low toxicity with an LD₅₀ greater than 10 g/kg.^[10] Due to the efficacy of bromelain after oral administration, its safety and lack of undesired side effects, it has high compliance among patients.^[11] Since bromelain is derived from the stem of pineapple, caution must be taken in patients with pineapple allergy. Caution should also be taken in patients using heparin, because in the study by Grabovac et al. (2006), evidence was found that heparin and bromelain form stable complexes leading to a significantly improved uptake of heparin, which could have an impact on blood-thinning potential.[12] Quercetin has an LD_{50} of 159 mg/kg in a mice model orally before reaching toxicity.^[13]

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Quercetin SAP Sciènce-based bioflavonoid for allergy and inflammation management



INDICATION SPECIFIC DOSAGE SUMMARY BASED ON HUMAN CLINICAL RESEARCH*

*Please note these suggestions are guidelines based on the clinical studies. Evidence for efficacy and safety have been qualitatively (study quality in terms of study design, sample size, appropriate methods of analysis, use of appropriate placebo/control, bias) assessed and have been rated using a 5 star \star rating classification.

Indication	Suggested Dosage	Supporting evidence and study outcomes	Study design	Outcomes measures/ selection criteria for studies	Safety	Evidence quality rating
Pulmonary Diseases						
SARS-Cov-2 infection ¹	1 capsule/day	Reduced the hospitalization period	Randomized open label control study, n=60, dose = 1000 mg/day for 1 week. Patients in the intervention group received quercetin in addition to remdesivir or favipiravir	Peripheral blood samples, serum levels of interleukin 1 beta, tumor necrosis factor alpha (TNF α), and interleukin-6 cytokines	No adverse events were reported	***
COPD and URTI ^{2,3}	2 capsules/ day	Lowered severity of URTI	2 Randomised, double- blind, placebo-controlled, single-centre dose- escalation studies, n=1012, avg dose = 1250 mg/day for 1-12 weeks	Wisconsin upper respiratory symptom survey, total plasma quercetin and its primary conjugates, COPD assessment test	Adverse events were gastro- esophageal reflux disease, stomach upset, breathlessness, chest congestion, headache, and nausea	****
Oxidative Stress						
Inflammation in sarcoidosis ⁴	2 capsules/ day	Reduced markers of oxidative stress and inflammation	Double-blind intervention study, n=18, dose = 2000 mg/day	Forced expiratory volume, forced vital capacity and diffuse capacity for carbon monoxide, plasma antioxidant status, vitamin C, uric acid, total glutathione, TNF- α , interleukin-8, and interleukin-6.	No adverse events were reported	**
Exercise in humans⁵	2 capsules/ day	Improved oxidative stress	Randomized, crossover, intervention study, n=14, dose = 1000 mg/day for 2 weeks.	Hemolysis assay, glutathione homeostasis, thiobarbituric acid reactive substances, intracellular superoxide dismutase, catalase and glutathione peroxidase activities	No adverse events were reported	**
Muscular Function						
Muscle Damage ^{6,7}	2 capsules/ day	Improved muscle weakness and enhanced recovery	2 Randomized, double- blind, crossover studies, n=24, avg dose = 1000 mg/ day for 2 weeks.	Neuromuscular tests, maximal voluntary isometric contractions, force-velocity relationship task, creatine kinase, interleukin-6, insulin-like growth factor (IGF-1 & IGF-II) levels	No adverse events were reported	***
Neuromuscular function ⁸	2 capsules/ day	Accelerated the recovery of neuromuscular function	Double-blind, placebo- controlled, crossover study, n=16, dose = 1000 mg/day for 2 weeks	Neuromuscular test, maximal voluntary isometric contraction, and force-velocity relationship	No adverse events were reported	***
Cardiovascular Health						
ΜI°	1 capsule/day	Enhanced total antioxidant capacity and improved the insecurity dimension of quality of life	Randomized double-blind, placebo-controlled trial, n=88, dose = 500 mg/day for 8 weeks	Body weight, BMI, body fat percentage, muscle mass percentage, visceral fat and blood pressure, 2-day food recall questionnaire, quality of life assessment, myocardial infarction and dimensional assessment scale	No adverse events were reported	***
Hypertension ¹⁰	1 capsule/day	Reduced systolic blood pressure and plasma, oxidized low- density lipoprotein concentrations	Randomized, double- blind, placebo-controlled, crossover study, n=41, dose = 730 mg/day for 4 weeks	Blood pressure measurement, blood lipid, glucose, urine collection, indices of oxidative stress, urinary isoprostane measurement, dietary analysis	No adverse events were reported	***

Women's Health



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★★ (For the

open-label

follow up

study)

Metabolic and 2 capsules/ Significant changes 2 Randomized, double-No adverse Glucose homeostasis, circulatory Hormonal day in lowering major blind, placebo-controlled testosterone, luteinizing hormone, effects were contributors of PCOS sex hormone-binding globulin, and reported Parameters trials, n = 79, avg dose = (PCOS)11, 12 1000 mg/day for 12 weeks anthropometric measurements, fasting serum levels of total adiponectin, highmolecular-weight (HMW) adiponectin Rheumatoid 1 capsule/day Reduced clinical Double-blind, placebo-Physician global assessment, disease No adverse Arthritis¹³ symptoms, and pain controlled randomized activity score-28, validated HAQ, plasma events were clinical trial, n=50, dose = hs-TNFα reported 500 mg/day for 8 weeks. Men's Health No adverse Pre-1 capsule/day Lowered plasma uric Randomized, double-Plasma glucose level, uric acid in hyperuricaemic acid levels blinded, placebo-controlled, plasma and urine samples events were males¹⁴ cross-over trial, n=22, dose = reported 500 mg/day for 4 weeks. Prostatitis¹⁵ 1 capsule/day Significant symptomatic Randomized double-Chronic prostatitis symptom score No adverse reductions blinded, placebo-controlled events were trial. n=30. dose = 500 mg/ reported In the open-label study day for 4 weeks with 17 patients who received a supplement In a follow-up unblinded, containing quercetin, open-label study, 17 bromelain, and papain, additional men received 82% had at least a a supplement containing 25% improvement in quercetin, bromelain, and symptom score papain (amounts unknown) to enhance bioflavonoid absorption for 4 weeks Thalassemia Iron 1 capsule/day Removed excess Randomized placebo-TNF-α hemoglobin, serum iron, ferritin, Some gastroiron, ferritin, hs-CRP, controlled trial, n=84, dose = transferrin, total iron binding overload and intestinal events inflammation¹⁶ capacity, transferrin saturation, and transferrin saturation, 500 mg/day for 12 weeks were reported and increased high sensitivity C-reactive protein transferrin Antiviral Chronic 1 capsule/day Decrease in hepatitis Phase 1 dose escalation Aspartate aminotransferase (AST) and No adverse Hepatitis C¹⁷ study, n=30, dose = 250-5000 alanine aminotransferase (ALT), Alkaline C virus events were mg/day for 4 weeks phosphate, total bilirubin, direct reported bilirubin, prothrombin time, partial thromboplastin time, international

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normalized ratio, hepatitis C viral load,

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