

Glucose Control Support



Nutritional Support for Healthy Glucose Metabolism

DESCRIPTION

Glucose Control Support contains a complementary blend of thiamin, biotin, chromium, alpha-lipoic acid, N-acetyl-L-cysteine, and standardized extract of *Gymnema sylvestris* leaf for targeted nutritional support of glucose metabolism.

FUNCTIONS

Glycation is the non-enzymatic attachment of sugars to major molecules in the body, including proteins, lipids, and nucleic acids. Glycation reactions generate advanced glycation end-products (AGEs) and glycotoxin intermediates. AGEs can cause abnormal and destructive functioning of body proteins, lipids, and nucleic acids. AGE-associated damage is suspected in the pathogenesis of many diseases and age-related deteriorations.

AGEs can be created endogenously, often as a consequence of excessive blood glucose levels. AGEs are also present in browned foods and cured tobacco. As uncontrolled blood glucose levels give rise to the formation of dangerous AGEs, it is crucial to maintain blood glucose levels within normal, healthy limits to avoid excessive AGE induced damage.

Maintenance of normal blood glucose levels is fundamental to preventing AGE formation. Chromium is an integral component of the glucose tolerance factor (GTF), a naturally occurring compound of chromium, nicotinic acid, and amino acids that is essential for proper glucose metabolism. Adequate chromium nutrition is essential for the formation of GTF and subsequent healthy metabolism of normal blood glucose levels.

Biotin serves as a cofactor of glucose metabolism and induces glucokinase, an enzyme that encourages cells to retain glucose for energy production rather than release it into the blood stream. Alpha-lipoic acid appears to enhance glucose use by muscles by augmenting muscle protein content. *Gymnema sylvestris* is an Ayurvedic botanical that may assist in the normal regeneration and repair of healthy pancreatic beta cells. *Gymnema* may also support healthy intestinal glucose absorption.

Select nutrients help to directly arrest glycation reactions. A derivative of thiamin (vitamin B1), called thiamin pyrophosphate, and alpha-lipoic acid have both been shown to prevent in vitro AGE production.

Also crucial to controlling AGE formation and subsequent damage is antioxidant protection. Oxidative stress is closely related to AGE production and is a known contributing factor to many of the same health concerns thought to be associated with AGE damage. In vitro studies of N-acetyl-L-cysteine have documented inhibition of glycation induced damage to pancreatic cells. N-acetyl-L-cysteine is a potent antioxidant, serving as an intracellular precursor of glutathione.

INDICATIONS

Glucose Control Support may be a useful dietary supplement for those who wish to support nutritional control of healthy blood sugar metabolism.

FORMULA (WW #10075)

2 Capsules Contain:

Thiamine (pyrophosphate)	50 mg
Biotin	2,500 mcg
Chromium (ChromeMate® polynicotinate*).....	800 mcg
Alpha-Lipoic Acid**	200 mg
N-Acetyl-L-Cysteine	500 mg
<i>Gymnema sylvestris</i> leaf,	400 mg
dried extract, 25% gymnemic acids	

* ChromeMate® brand. U.S. Patent Number 4,923,855
InterHealth Co.

** Lipoic Acid used in this product is of the highest quality available and of Italian origin.

SUGGESTED USE

Two capsules daily, or as directed by a healthcare professional.

SIDE EFFECTS

No adverse effects have been reported.

STORAGE

Store in a cool, dry place, away from direct light. Keep out of reach of children.

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REFERENCES

- Anderson RA. Nutritional factors influencing the glucose/insulin system: chromium. *J Am Coll Nutr* 1997;16:404-10.
- Anderson, RA. Chromium, glucose intolerance, and diabetes. *J Am Coll Nutr* 1998;17:548-55.
- Baskaran K, Kizar Ahamath B, Radha Shanmugasundaram K, Shanmugasundaram ER. *J Ethnopharmacol* 1990;30:295-300.
- Bierhaus A, Chevion S, Chevion M, Hofmann M, et al. Advanced glycation end product-induced activation of NF-kappaB is suppressed by alpha-lipoic acid in cultured endothelial cells. *Diabetes* 1997;46:1481-90.
- Booth AA, Khalifah RG, Hudson BG. Thiamine pyrophosphate and pyridoxine inhibit the formation of antigenic advanced glycation end-products: comparison with aminoguanidine. *Biochem Biophys Res Commun* 1996;220:113-9.
- Chauhan J, Dakshinamurti K. Transcriptional regulation of the glucokinase gene by biotin in starved rats. *J Biol Chem* 1991;266:10035-8.
- Cunningham JJ. Micronutrients as nutraceutical interventions in diabetes mellitus. *J Am Coll Nutr* 1998;17:7-10.
- De Mattia G, Bravi MC, Laurenti O, Cassone-Faldetta M, et al. Reduction of oxidative stress by oral N-acetyl-L-cysteine treatment decreases plasma soluble vascular cell adhesion molecule-1 concentrations in non-obese, non-dyslipidaemic, normotensive, patients with non-insulin dependent diabetes. *Diabetologia* 1998;41:1392-6.
- Frye EB, Degenhardt TP, Thorpe SR, Baynes JW. Role of the Maillard reaction in aging of tissue proteins. Advanced glycation end product-dependent increase in imidazolium cross-links in human lens proteins. *J Biol Chem* 1998;273:18714-9.
- Kaneto H, Fujii J, Myint T, et al. Reducing sugars trigger oxidative modification and apoptosis in pancreatic beta-cells by provoking oxidative stress through the glycation reaction. *Biochem J* 1996;320:855-63.
- Koutisikos D, Agroyannis B, Tzanatos-Exarchou H. Biotin for diabetic peripheral neuropathy. *Biomed Pharmacother* 1990;44:511-4.
- Levi B, Werman MJ. Long-term fructose consumption accelerates glycation and several age-related variables in male rats. *J Nutr* 1998;128:1442-9.
- Low PA, Nickander KK, Tritschler HJ. The roles of oxidative stress and antioxidant treatment in experimental diabetic neuropathy. *Diabetes* 1997;46 Suppl 2:S38-42.
- Mertz W. Chromium in human nutrition: a review. *J Nutr* 1993;123:626-33.
- Obrosova I, Cao X, Greene DA, Stevens MJ. Diabetes-induced changes in lens antioxidant status, glucose utilization and energy metabolism: effect of DL-alpha-lipoic acid. *Diabetologia* 1998;41:1442-50.
- Okabayashi Y, Tani S, Fujisawa T, et al. Effect of *Gymnema sylvestre*, R.Br. on glucose homeostasis in rats. *Diabetes Res Clin Pract* 1990;9:143-8.
- Preuss HG. Effects of glucose/insulin perturbations on aging and chronic disorders of aging: the evidence. *J Am Coll Nutr* 1997;16:397-403.
- Shanmugasundaram ER, Rajeswari G, Baskaran K, et al. Use of *Gymnema sylvestre* leaf extract in the control of blood glucose in insulin-dependent diabetes mellitus. *J Ethnopharmacol* 1990;30:281-94.
- Shanmugasundaram KR, Panneerselvam C, Samundram P, Shanmugasundaram ER. Enzyme changes and glucose utilization in diabetic rabbits: the effect of *Gymnema sylvestre*, R. Br. *J Ethnopharmacol* 1983;7:205-34.
- Thorpe SR, Baynes JW. Role of the Maillard reaction in diabetes mellitus and diseases of aging. *Drugs Aging* 1996;9:69-77.
- Yang CW, Vlassara H, Peten EP, et al. Advanced glycation end products up-regulate gene expression found in diabetic glomerular disease. *Proc Natl Acad Sci USA* 1994;91:9436-40.
- Ziegler D, Hanefeld M, Ruhnau KJ, et al. Treatment of symptomatic diabetic peripheral neuropathy with the antioxidant alpha-lipoic acid. A 3-week multicentre randomised controlled trial (ALADIN Study). *Diabetologia* 1995;38:1425-33.

These statements have not been evaluated by the Food and Drug Administration. This product is not intended to diagnose, treat, cure, or prevent any disease.

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