Glycomacropeptide (GMP)-Based Food for the Treatment of PKU and Other Metabolic Disorders

INVENTORS • Denise Ney, Mark Etzel

WARF: PO9323US02
View U.S. Patent No. 8,604,168 in PDF format.

The Wisconsin Alumni Research Foundation (WARF) is seeking commercial partners interested in developing improved medical food for the nutritional management of phenylketonuria (PKU).

OVERVIEW

Phenylketonuria (PKU) is a genetic disorder in which an individual lacks the enzyme phenylalanine hydroxylase (PAH) that converts the amino acid phenylalanine into tyrosine. If left untreated, the buildup of phenylalanine in the blood can lead to mental retardation and central nervous system disorders.

The treatment for PKU is a lifelong reduced protein diet. Because most proteins contain significant amounts of phenylalanine, the specialized diet usually is protein poor and supplemented with other amino acids. This diet is difficult to follow, restrictive and unpalatable. Non-compliance is a common problem and can cause severe neuropsychological impairment.

THE INVENTION

UW-Madison researchers have developed an improved medical food for treating PKU. This food is made with highly purified glycomacropeptide (GMP) as its primary protein source and supplemented with other amino acids, including arginine, histidine, leucine, tyrosine and tryptophan. It provides a complete, low-phenylalanine source of protein and is more palatable than the standard specialized diet.

GMP is a naturally occurring protein that is formed during cheese making and contains no phenylalanine. The purity of GMP is one key to producing this medical food. The other is the amount and type of amino acid supplementation. For example, if too many sulfur-containing amino acids are used, the food tastes bad and patients will not eat it.
APPLICATIONS

- Nutritional management of PKU

KEY BENEFITS

- Provides the highest quality, most effective medicinal foodstuff available for treating patients with PKU
- Provides a complete source of protein while reducing the levels of phenylalanine in the blood of patients with PKU
- Tastes good enough for patients to remain compliant with the diet
- Economically feasible to manufacture
- Can be produced as a variety of food types, including beverages, bars, wafers, puddings, gelatins, crackers, fruit leathers, nut butters, sauces, flakes, crisp cereal pieces, puffs, pellets and extruded solids
- May be used to treat other metabolic disorders in addition to PKU

STAGE OF DEVELOPMENT

This food was successfully tested in human trials. It also was shown to support normal growth and reduce levels of phenylalanine in the blood and brain of PKU mice.

ADDITIONAL INFORMATION

Publications


Click here for a news release describing this technology.


Tech Fields

Food & Supplements - Functional foods
Pharmaceuticals & Vitamin D - Metabolic disorders
CONTACT INFORMATION

For current licensing status, please contact Mark Staudt at mstaudt@warf.org or 608-960-9845.