



Ergothioneine

CAS

Molecular Weight

Molecular Formula

Appearance

Structure

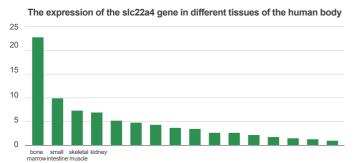
497-30-3

C₀H₁₅N₂O₂S

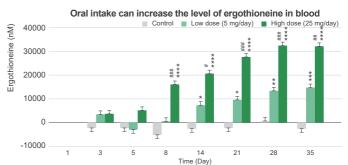
White powder

Product Information

Ergothioneine (EGT) is a natural amino acid derivative found in many plants and animals. Unlike other amino acids, ergothioneine cannot be synthesized by the human body, and thus, it can only be obtained through a balanced diet. Transporting ergothioneine into cells is made possible by a highly specific cationic transporter protein known as slc22a4 (previously known as OCTN1). With its potent antioxidant properties, ergothioneine is an effective ingredient in reducing free radicals and has a significant impact on human physiological activities.



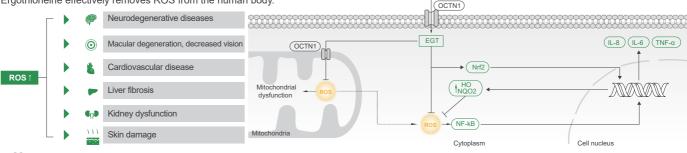
The gene slc22a4 encoding the transporter of ergothioneine is widely expressed in in various tissues and organs, suggesting the universal presence of ergothioneine in human body.



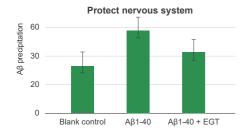
45 healthy volunteers were divided into three groups and were provided with a placebo, a low dose of ergothioneine (5 mg/day), and a high dose of ergothioneine (25 mg/day), respectively. It was found that the level of ergothioneine was significantly increased in blood after 35 days, without any adverse effects.

Mechanism

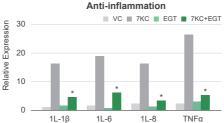
Ergothioneine effectively removes ROS from the human body



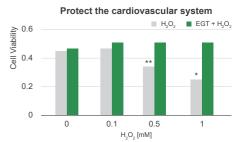
Efficacy



The control group was fed with physiological saline, while the $A\beta1-40$ group was injected with amyloid protein ($A\beta$ 1-40) (5 μ L 400 pmol/mouse) into the hippocampus and then fed with physiological saline. The A β 1-40 + EGT group was additionally fed with 0.5 mg/kg EGT. After 39 days, and it was found that EGT significantly reduced the number of $A\beta$ deposits in the hippocampus



In vitro cultured endothelial cells were divided into four groups (control group, 30 μ M 7-ketocholesterol (7KC) as the negative control, 1 mM EGT, 7KC + EGT). After 24 hours of culturing, an increase in the expression of IL-18. IL-6, IL-8 and TNF α in endothelial cells induced by 7KC was observed. Addition of EGT significantly reduced the expression of inflammatory factors



Arterial endothelial cells were cultured in vitro and treated with varying concentrations of $\mathrm{H_2O_2}$, resulting in a significant decrease in cell viability. However, upon addition of 0.5 mM EGT, a significant increase in cell viability was observed, preventing damage to the endothelial cells.

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