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(Article)

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## Abstract

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Obesity has been often associated with the occurrence of cardiovascular diseases, type 2 diabetes, and cancer. The development of obesity is also accompanied by significant differentiation of preadipocytes into adipocytes. In this study, we investigated the activity of  $\alpha$ -mangostin, a major xanthone component isolated from the stem bark of *G. malaccensis*, on glucose uptake and adipocyte differentiation of 3T3-L1 cells focusing on PPAR $\gamma$ , GLUT4, and leptin expressions.  $\alpha$ -Mangostin was found to inhibit cytoplasmic lipid accumulation and adipogenic differentiation. Cells treated with 50  $\mu$ M of  $\alpha$ -mangostin reduced intracellular fat accumulation dose-dependently up to 44.4% relative to MDI-treated cells. Analyses of 2-deoxy-D-[<sup>3</sup>H] glucose uptake activity showed that  $\alpha$ -mangostin significantly improved the glucose uptake ( $P < 0.05$ ) with highest activity found at 25  $\mu$ M. In addition,  $\alpha$ -mangostin increased the amount of free fatty acids (FFA) released. The highest glycerol release level was observed at 50  $\mu$ M of  $\alpha$ -mangostin. qRT-PCR analysis showed reduced lipid accumulation via inhibition of PPAR $\gamma$  gene expression. Induction of glucose uptake and free fatty acid release by  $\alpha$ -mangostin were accompanied by increasing mRNA expression of GLUT4 and leptin. These evidences propose that  $\alpha$ -mangostin might be possible candidate for the effective management of obesity in future.

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## Indexed keywords

EMTREE drug terms: alpha mangostin fatty acid glucose glucose transporter 4 leptin peroxisome proliferator activated receptor gamma plant medicinal product unclassified drug

EMTREE medical terms: adipocyte adipogenesis animal cell Article bark cell differentiation cell viability controlled study drug isolation *Garcinia* *Garcinia malaccensis* gene expression glucose transport lipid storage mouse MTT assay nonhuman priority journal reverse transcription polymerase chain reaction

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