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Human peroxisome proliferator-activated receptor (PPAR) γ agonists identified in selected plant extracts by bioassay-guided fractionation

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Introduction

Abdominal obesity is associated with a set of risk factors known as the metabolic syndrome. Type 2 diabetes (TZD) is one of these and is caused by a combination of insulin resistance and δ-cell failure. Lifestyle modifications are rarely sufficient for the treatment of T2D, thus thiazolidinedione (TZD) drugs are often prescribed. TZDs are agonists of PPARγ, a key regulator of adipogenesis. Activation of PPARγ causes re-distribution of fat in the body, which leads to increased insulin sensitivity. Severe side-effects are associated with the use of TZDs e.g. weight gain and edemas. These side-effects occur because TZDs are full PPARγ agonists. However, it has been suggested that partial PPARγ agonists do not have severe side-effects.

Plants have been used in the traditional treatment of diabetes but the mechanism of action of the bioactive compounds is rarely known. In this study, we focused on the identification of natural products from medicinal plants able to activate PPARγ. In a screening for partial PPARγ agonists we identified several plant species (60% hit rate) able to activate PPARγ and increase insulin-stimulated glucose uptake in adipocytes without having an adipogenic potential. Many of these were also activators of PPARα and δ [1].

Purple coneflower (Echinacea purpurea)

The n-hexane extract of the flowers of purple coneflower was one of the most promising extracts with respect to activation of PPARγ. Bioassay-guided fractionation led to the isolation and identification of α-linolenic acid, linoleic acid, palmitoleic acid, dodeca-2E,4Z,10Z-trien-8-ynoic acid isobutylamide, dodeca-2E,4E,8Z,10E/Z-tetraenoic acid isobutylamide, dodeca-2E,4E-dienic acid isobutylamide, and the new hexadeca-2E,9Z,12Z,14E-tetraenoic acid isobutylamide (1).

A luciferase-based PPARγ transactivation assay using mouse embryonic fibroblasts was used for initial assessment of bioactivity. All three fatty acids are well-known PPARγ agonists and were also found to be active in our test-system. The alkamides also activated PPARγ, of which compound 1 was the most effective of these (Fig. 1) [2].

Compound 1 had no adipogenic potential and was able to positively affect insulin-stimulated glucose uptake in a manner similar to that of Rosiglitazone, which was used as positive control (Fig. 2) [2].

Conclusions

- High hit-rate among plants for identification of PPAR modulators.
- PPAR modulators from plants are in general fatty acids together with specific metabolites such as:
  - Alkamides (preferably ≥ C16)
  - Flavonoid aglycones
- In general, flavonoid glycosides and phenolic acids do not activate PPARs.
- Potential synergistic effects exist between some of the identified PPAR modulators as overall bioactivity could not be explained completely by individual activities.

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