

Orally administered *Polypodium leucotomos* extract decreases psoralen-UVA-induced phototoxicity, pigmentation, and damage of human skin.

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Abstract

BACKGROUND:

The use of psoralen-UVA (PUVA) in patients of skin phototype I to II is limited by side effects of acute phototoxicity and possible long-term carcinogenesis.

OBJECTIVE:

We sought to assess oral *Polypodium leucotomos* (PL) extract in decreasing PUVA-induced phototoxicity of human skin on a clinical and histologic level.

METHODS:

A total of 10 healthy patients with skin phototypes II to III were exposed to PUVA alone (using 0.6 mg/kg oral 8-methoxypsoralen) and to PUVA with 7.5 mg/kg of oral PL.

RESULTS:

Clinically, phototoxicity was always lower in PL-treated skin after 48 to 72 hours ($P < .005$), and pigmentation was also reduced 4 months later. Histologically, PL-treated skin showed a significant numeric reduction of sunburn cells ($P = .05$), preservation of Langerhans cells ($P < \text{ or } = .01$), decrease of tryptase-positive mast cell infiltration ($P < .05$), and decrease of vasodilation ($P < \text{ or } = .01$). No differences were found in Ki-67+ proliferating cells.

CONCLUSIONS:

PL is an effective chemophotoprotector against PUVA-induced skin phototoxicity and leads to substantial benefits of skin protection against damaging effects of PUVA as evidenced by histology.

PMID: 14699363

DOI: [10.1016/S0190](#)