# A New Topical Agent, Zanfel®, Ameliorates Urushiol-Induced *Toxicodendron* Allergic Contact Dermatitis

ACEP's 2003 Research Forum



A Davila, MD, J Lucas, DO, M Laurora, DO, J Jacoby, MD, J Reed, PhD, and M Heller, MD, Emergency Medicine Residency, and Research Institute, St. Luke's Hospital, Bethlehem, PA

# BACKGROUND/OBJECTIVE

Toxicodendron exposure and subsequent allergic contact dermatitis (ACD) is a significant health issue and frequent cause for emergency department visits. The resulting monetary cost to states with *Toxicodendron* plants can be substantial. Poison ivy, poison oak, and poison sumac are all common causes of ACD and the vast majority of the general population is susceptible, with perhaps one million victims per year. Treatment for the condition may be difficult, however, with systemic anti-histamines and steroids as well as topical agents all being limited by concerns of efficacy or side effects. A new topical agent, Zanfel®,(Zanfel Laboratories Inc., Morton, IL) a mixture of alcohol solubles and anionic surfactants. has the potential to bind the urushiol resin post exposure, even after symptom onset. If effective, this would provide the emergency physician with a new mode of therapy for this common problem. The objective of this study was to determine if a new topical wash, Zanfel®, is an effective post exposure treatment of Toxicodendron ACD

# **METHODS**

This was a randomized. controlled, double-blind trial with 24 consenting volunteers. With each subject serving as his or her own control, both forearms received three one square inch sites of skin exposure to 1.5 micrograms of purified T. radicans urushiol resin (see Figure 1). At three subsequent times post exposure (48, 96 and 144 hours), the sites were treated





FIGURE 1

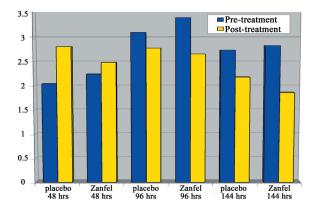
with placebo or Zanfel®. Pre-treatment and posttreatment ACD reaction scores and subjective itch scores were compiled at each time. ACD was scored using a zero to six point Likert scale based upon erythema,

induration, and vesiculation; itch was quantified using a tencentimeter visual analog scale. Comparisons were by paired ttest, with alpha set at 0.05.

At all times following urushiol exposure, Zanfel® significantly improved the Toxicodendron-induced ACD compared to placebo. The difference in ACD scores at times 48, 96 and 144 hours respectively, all favored the Zanfel® group versus placebo (-0.76 vs. -0.24; 0.32 vs. 0.74; and 0.55 vs. 0.96, all p<0.05). (See Figure 2) Itch scores improved for both groups at all three time points with a trend favoring the Zanfel® group at 96 hours (p=.058).

RESULTS

## FIGURE 2 - ACD REACTION RESULTS



Reprinted from Annals of Emergency Medicine, V42 (4 suppl 1), Davila A, et al, "A New Topical Agent, Zanfel, Ameliorates Urushiol-Induced Toxicodendron Allergic Contact Dermatitis," Abstract 364, © 2003, with permission from American College of Emergency Physicians.

Traditional treatments for urushiol-

induced ACD have utilized one of two

inflammatory (using steroids and anti-

histamines to treat the reaction, once

different strategies: prophylaxis (barrier

creams to prevent resin exposure) and anti-

established). Zanfel, however, appears to

provide an additional approach. In our

study it appeared to effectively bind the

toxin thus ameliorating the ACD symptoms

within a short time after application. The

Zanfel treated sites demonstrated less

ervthema and induration than those sites

treated with placebo. In some cases the

Zanfel appeared to prevent a reaction

entirely. (See Figure 3) The effect of

Zanfel on itch was more difficult to discern

the degree of itch shortly after washing the

sites either with Zanfel or the placebo. The

physical act of washing caused a clinically

significant decrease in itch, which may

have attenuated the effect of the active

agent.

possibly due to our protocol, which rated

# DISCUSSION

Zanfel has thus far had almost no role in emergency practice. No peer-reviewed study documenting or refuting its effectiveness has appeared. The manufacturer claims that relief from ACD symptoms occurs within seconds to minutes of a single application; subsequent or multiple applications in recalcitrant cases are recommended. Our study documented the potential of Zanfel for treatment of experimentally induced ACD. A clinical study of naturally occurring poison ivy rash is underway. Zanfel

FIGURE 3

# CONCLUSIONS

The new topical agent, Zanfel®, when applied after exposure to urushiol, ameliorates or prevents Toxicodendroninduced allergic contact dermatitis.