

**A RANDOMIZED, DOUBLE-BLIND, PLACEBO-
CONTROLLED, PARALLEL-GROUP PILOT STUDY TO
INVESTIGATE THE EFFECTS OF HUMIC ACID ON
SYMPTOMS OF INFLUENZA**

FINAL REPORT SUMMARY ABSTRACT

Prepared By:

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Study Carried Out By:

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Objectives:

The primary objective of this Study was to assess the efficacy of the Humic Acid used in Viracillin® - Activated Humic Acid™ on flu symptoms as assessed by the alleviation of symptoms after 7 and 14 days of treatment in adults with influenza A or B.

Methodology:

This was a multi-center, randomized, double-blind, placebo-controlled, parallel-group study with a single 14-day treatment period to assess the efficacy of the Humic Acid used in Viracillin® - Activated Humic Acid™ on flu symptoms in adults. This Study was conducted at two sites in the US: Advanced Research Institute, Inc. (Trinity, FL) and Vertitas Research Corp. (Miami Lakes, FL). The Study was managed by KGK Synergize, Inc.

Efficacy Results:

Subjects on the Humic Acid used in Viracillin® - Activated Humic Acid™ reported higher scores for cough, fever, runny nose, stuffy nose, aches, chills, sneezing, earaches and fatigue on Day 1. After supplementation for 7 days with the Humic Acid used in Viracillin® - Activated Humic Acid™, symptom scores improved by a greater percentage than did those for Subjects taking Placebo for the same duration for cough (61.9% vs. 36.8%), fever (91.7% vs. 81.8%), runny nose (66.7% vs. 62.5%), stuffy nose (66.7% vs. 62.5%), aches (86.4% vs. 62.5%), chills (91.7% vs. 66.7%), sneezing (70.6% vs. 61.5%) and fatigue (80.0% vs. 54.5%). Mean scores for Subjects on the Humic Acid used in Viracillin® - Activated Humic Acid™ were also lower than for those on Placebo after 7 days of treatment for cough, fever, aches, chills and fatigue in spite of being worse than Placebo on Day 1.

Conclusions:

The Humic Acid used in Viracillin® - Activated Humic Acid™ demonstrated clinically-noteworthy efficacy on influenza viral infection in the 2010 flu season by improving flu-related symptoms, cytokine response, immune-system modulators CD4+ and CD8+, and VAS scores when provided as 6 x 250-mg tablets daily for 14 days within the demographic of Subjects studied.