Please Join

American Herbal Products Association, Consumer Healthcare Products Association,
Council for Responsible Nutrition, Natural Products Association and
United Natural Products Alliance

In Cooperation With The

Congressional Dietary Supplement Caucus

For A Mid-Afternoon Snack Briefing On

Serious Adverse Event Reporting: What It Reveals About Dietary Supplement Safety

Thursday, February 27th ★ 2:30 – 3:30 PM

121 Cannon House Office Building

The Dietary Supplement and Nonprescription Drug Consumer Protection Act, signed into law in December 2006, requires manufacturers to report to FDA serious adverse events (AERs) associated with dietary supplements. In addition, supplement companies must maintain accurate records for six years for any non-serious adverse event reports they receive. Over the past 12 years, the data collected from these requirements have provided insights into the safety of dietary supplements and given FDA early warnings of potential safety concerns. AER data is already available for hemp-derived CBD products.

This briefing will examine the AER data collected and what it reveals about the safety profile of dietary supplements: Does the industry have a safety problem? What products are most often associated with adverse events? Are safety concerns already evident for CBD-containing supplements? Dr. Rick Kingston will explain how AER data has helped ensure consumer safety and explain what adverse event reporting exposes from the introduction of hemp-derived CBD products.

Speaker: Richard "Rick" Kingston, PharmD – Clinical Professor, University of Minnesota/College of Pharmacy, Co-Founder and President, Regulatory and Scientific Affairs, SafetyCall International

Interns welcome when accompanied by Permanent Staff
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