EXPERT PANEL

MEADERS: Medication Events and Adverse Drug Events Recording System

Meeting: APRIL 5-6, 2006 Rockville, MD

Expert	Area of Expertise
Alan Lembitz, MD	Vice President of COPIC and the Director of Risk Management.
	Assists physicians in risk reduction and practice improvement
	that both improves care and lowers overall malpractice risk.
Diane Cousins, RPh	Developed the US Pharmacopeia Practitioners' Reporting
	Network into four nationwide reporting programs that focus on
	the quality, efficacy, and safety of over-the-counter and
	prescription drugs used in human and veterinary medicine.
Gurdev Singh, MD	Director of Patient Safety Research Center at SUNY-Buffalo.
	Civil engineer with a background in safety in primary care. He
	has designed a web-based error reporting system.
Harold Kaplan, MD	Professor of Clinical Pathology and Director of Transfusion
	Medicine at the Columbia University Medical Center. Current
	research effort is directed at establishing the usefulness of
	standardized medical event reporting for error prevention and
	management throughout healthcare delivery systems.
Kathleen Stevens, RN,	Professor at the University of Texas Health Science Center
EdD	School of Nursing, Family Care Department and the Graduate
	School of Biomedical Sciences and a Certified Advance
	Practice Nurse. Her research includes children's health
	promotion, evidence synthesis, evidence-based practice, and
	informatics applications.
Kim Galt, PhD	Professor of Pharmacy, developed and managed pharmacists'
	primary care consultative ambulatory practice sites in the
	private and Veternas Affairs systems, supervised specialized
	drug information and clinical pharmacy services, and practiced
	general hospital, community and long-term care pharmacy.
Mark Lehto, PhD	Associate Professor and Industrial engineer whose research
	focuses on human decision-making, hazard communication,
	safety and ergonomics. His work in the area of product warning
	and instructions is nationally recognized.
Michael Cohen, RPh,	President of the Institute for Safe Medication Practices (ISMP),
MS, ScD	member of the Drug Safety and Risk Management Advisory
	Panel for the U.S. Food and Drug Administration (FDA), the
	committee on identifying and preventing medication errors for
	the Institute of Medicine, and the national Quality Forum's
	National Voluntary Consensus Standards Maintenance
	Committee on Safe Practices.

Morris Weinberger,	Professor of HealthCare quality at the University of North
PhD	Carolina, Chapel Hill. A behavioral scientist who has
	performed at number of interventional and observational studies
	involving drug therapy in PBRNs and community drugstores.
Ranjit Singh, MD,	Director of Patient Safety Research Center at the Family
MBA	Medicine Research Institute, SUNY-Buffalo. Experience in
	software development and systems engineering issues.
R. Scott Evans, PhD,	Director of Research for the Department of Clinical
MS	Epidemiology. Designs, develops and evaluates computerized
	tools for the selection and management of anti-infective agents,
	computer methods to identify and reduce adverse drug events,
	computerized methods to identify patients needing isolation,
	and computerized methods to identify and reduce hospital-
	acquired infections.