



The University of Chicago

The Division of Biological Sciences • The Pritzker School of Medicine

The University of Chicago Hospitals

Institutional Review Board

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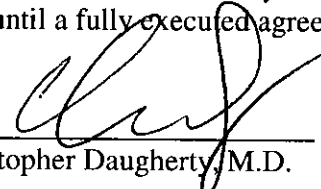
INSTITUTIONAL REVIEW BOARD CERTIFICATION

Principal Investigator:	Chris Shea
Facex:	MC 5067
IRB Protocol Number:	13844B
Type of Submission:	Revised Deferral
Meeting Date:	09/27/2005
Risk Determination:	Minimal Risk
Informed Consent:	Written
Special Population:	Minors, Economically Disadvantaged, Non-English Speaking, Illiterate
Advertisement(s) Used:	No
Protocol Title:	Selenium Dietary Supplements Against Arsenic Toxicity in Bangladesh: A Proposed, 48-week, 800-patient, Randomized, Placebo-Controlled, Double Blind Clinical Trial
Protocol Version:	20 January 2005
Consent Version:	Texas Tech IRB approved consent form (U of C IRB received 15 September 2005)
Grant:	NIH Proposal (dated 8 August 2005)

STATUS: Approved*

THE RESEARCH PROTOCOL AND/OR CONSENT FORM DESCRIBED ABOVE HAVE BEEN REVIEWED BY THE IRB WITH THE RESULTS AS INDICATED. Please note that any externally funded research, even if approved by the IRB, may not be initiated until a fully executed agreement has been approved by University Research Administration.

Date: **SEP 27 2005**

Signature of Vice-Chair: 
Christopher Daugherty, M.D.

Federal regulations require that any severe drug reaction and unexpected or adverse occurrence to subjects during the conduct of this research be reported to the IRB. Any changes to this protocol must be submitted for review to the IRB.

**Approval Period: September 27, 2005 through September 26, 2006*