INDEPENDENT
INVESTIGATIONAL
REVIEW BOARD INC.
Your Advocate for Clinical Research Participants

DATE: April 4, 2006

TO: Scott Carroll, Ph.D.
Principal Investigator

FROM: Kim Lerner, Chairman or
Anita McSharry, Vice-Chairman
Independent Investigational Review Board, Inc.

SUBJECT: Approval Template Clinical Research Protocol dated: March 30, 2006

PROTOCOL: (CL-001)

The Independent Investigational Review Board, Inc. is an institutional review committee structured in compliance with the regulations of the Food and Drug Administration contained in the Code of Federal Regulations (2ICFR 50 and 56), Environmental Protection Agency (40 CFR parts 9 and 26), and is in compliance with the International Conference of Harmonization (ICH) Good Clinical Practice (GCP) guidelines for IRB/IECs.

At the April 4, 2006 meeting, the Committee unanimously approved the Protocol as a template for future research. Specific protocol review and approval for any future research is required prior to its implementation. The template research protocol must include test substance information, specific research procedures, and an Informed Consent Form.

Thank you for your cooperation.

KL/AMS/ds/rr

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