Texas Applied Biomedical Services

September 22, 2004

Panos Pappas, PhD
11502 North Poema Place #204
Chatsworth, CA 91311

RE: Clinical Study Accession Number: TABS 2004-0041
A Pre-Clinical Study to Evaluate the PAP Ion Magnetic Inductor Device in the Management of Myalgias and Arthralgias

Dear Pappas:

I am happy to inform you that T.A.B.S. Research Review Committee (IRB) approved the Clinical Investigational Study Protocol entitled **"A Pre-Clinical Study to Evaluate the PAP Ion Magnetic Inductor Device in the Management of Myalgias and Arthralgias"** at our September 21, 2004 meeting. A copy of the Clinical Investigational Plan and Protocol, as approved is enclosed for your records. This approval expires September 21, 2005.

Each clinical study site and Principal Investigator (PI) Application must be submitted to TABS Research Review Committee for their review and approval prior to initiating the clinical study. A copy of the PI Application Packet and Principal Investigator Obligations are also enclosed for your convenience.

Each Investigational Device must be labeled in accordance with 21 CFR 812.5 with the following information:

1. Name and place of business of the device manufacturer
2. Caution Statement:
   CAUTION: Investigational device. Limited by Federal (or United States) law to investigational use.

TABS Research Review Committee follows the procedures described in the Federal Food and Drug Administration's Code of Federal Regulations, 21 CFR Part 56. Should you have any questions, please do not hesitate to contact me at 713 / 734-4433.

Best regards,

M. Joyce Heinrich
Chairperson
TABS Research Review Committee

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Let us keep TABS for you!

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