

US EPA ARCHIVE DOCUMENT

Tuesday, November 07, 2006  
MINUTES

**ATTENDANCE:**

**PRESENT**

David Wells, MD  
Anita McSharry, RN  
Shari Somerstein, RPh  
Edward Wiederhorn  
Rabbi Akiva Mann  
Kim Lerner

**ABSENT**

George Garbarino

**GUEST**

Katy Kysela

**ALSO PRESENT**

Marcos Rejtman, DO

**I. CALL TO ORDER**

The meeting was called to order at 10:32 AM, by Chairman, Kim Lerner. The meeting was held at 6738 West Sunrise Blvd., Suite 102, Plantation, FL 333313.

**II. APPROVAL OF THE 10/31/2006 MINUTES**

The minutes of the meeting held 10/31/2006 were reviewed and unanimously approved as reviewed.

**III. REVIEW PROTOCOLS**

**L (Protocol SCI-001) Test of Personal Insect Repellents**

Principal Investigator: Scott P. Carroll, PhD

- Approval Clinical Research Protocol dated: 11/2/2006
- Informed Consent Form (Ver. 11/7/2006)
- Site Questionnaire
- The Experimental Subject's Bill of Rights

Motion was made, seconded and the Committee unanimously approved the Research Protocol, the Investigator(s), Informed Consent Form, MSDS, The Experimental Subject's Bill of Rights for the above noted research study. The Site Questionnaire was reviewed and unanimously accepted.

The Informed Consent Form is unanimously approved as submitted. The approved revised Informed Consent Form is identified as Version 11/7/2006 and stamped, "Approved 11/7/2006". The Informed Consent Form contains all regulatory required consent elements. The Experimental Subject's Bill of Rights is stamped "Approved 11/7/2006".

The Committee evaluated that the risks to the subjects were minimized and that a reasonable risk/benefit ratio is established. Based on the duration of the study and the risks to the subjects, the approval is granted for a one year period, with a progress report required prior to continued approval. See Approval letter for Investigator's responsibilities and file for supporting documentation.