**CSU CRB Client Consent Form Review Checklist:**

Does the client consent form contain the following recommended elements?

\_\_\_\_\_ An explanation of the purposes of the study

\_\_\_\_\_ The expected duration of the subject's participation

\_\_\_\_\_ A description of the procedures to be followed

\_\_\_\_\_ Identification of any procedures which are experimental

\_\_\_\_\_ A description of any reasonably foreseeable risks or discomforts to the subject that are related to the study.

\_\_\_\_\_ A description of any benefits to the subject or to others which may reasonably be expected from the research

\_\_\_\_\_ A disclosure of appropriate alternative procedures or courses of treatment, if any, that might be advantageous to the subject

\_\_\_\_\_ A statement describing the extent, if any, to which confidentiality of records identifying the subject will be maintained

\_\_\_\_\_ For research involving more than minimal risk, an explanation as to whether any compensation, and an explanation as to whether any medical treatments are available, if injury occurs and, if so, what they consist of, or where further information may be obtained

\_\_\_\_\_ A statement that participation is voluntary, refusal to participate will involve no penalty or loss of benefits to which the subject is otherwise entitled, and the subject may discontinue participation at any time without penalty or loss of benefits, to which the subject is otherwise entitled.

\_\_\_\_\_ A statement appropriately explaining any Conflict of Interest

\_\_\_\_\_ Dates and signatures

**Reviewer Comments:**

Potential areas for comment on Scientific Merit and Experimental Design:

- Significance/impact (Is study clinically relevant?)

- Approach

- Is the study sufficiently powered to achieve significant results?

-Other