

**Principal Investigator:**  
**IRB #:**

**DOCUMENTATION OF CONSENT PROCESS**

**Subject Initials:** \_\_\_\_\_

Person obtaining consent initial each completed step in the process:

\_\_\_\_\_ Informed consent was discussed with subject for the above referenced study. Copy of the consent form was provided for subject and/or authorized subject representative review.

\_\_\_\_\_ Subject and/or authorized subject representative was given adequate time to read the consent form and discuss the study with study investigators and/or family members.  
subject representative upon conclusion of the consent process.

\_\_\_\_\_ During informed consent process, the following questions were asked by the subject and/or authorized representative and were answered by study personnel:

\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

\_\_\_\_\_ Consent has been signed prior to any study procedures being performed.

Consent process documented by: \_\_\_\_\_  
Print Name

\_\_\_\_\_  
Signature

\_\_\_\_\_  
Date