

# **INFORMED CONSENT FORM**

**Title of the Study:**

**Principal Investigators, Department, Institution:**

**Co-Investigator(s), Department/Hospital/Institution:**

## **1. INFORMATION**

## **2. PURPOSE OF THIS RESEARCH STUDY**

## **3. DESCRIPTION OF THE RESEARCH PROCEDURE**

## **4. ASPECTS OF THE TRIAL THAT ARE NOT STANDARD CARE FOR YOUR CONDITION OR ARE EXPERIMENTAL**

**5. WHO CAN PARTICIPATE**

**6. LENGTH OF YOUR PARTICIPATION**

**7. POTENTIAL RISKS (INCLUDING PHYSICAL, PSYCHOLOGICAL, SOCIAL HARM, DISCOMFORT, OR INCONVENIENCE)**

**8. BENEFITS OF THE RESEARCH TO SOCIETY AND/OR TO THE INDIVIDUAL**

**9. CONFIDENTIALITY AND RELEASE OF PERSONAL INFORMATION**

## **10. ALTERNATIVE PROCEDURES OR TREATMENTS**

## **11. IN CASE OF RESEARCH RELATED INJURY**

## **12. COSTS AND PAYMENT FOR PARTICIPATION**

## **13. WHOM TO CONTACT**

#### 14. CONSENT STATEMENT

I therefore certify the following:

- I have read the above information form and understand that the study involves research. I understand the purpose of the study as well as the potential benefits and risks of participating in the study.

**I agree to participate in this study and I understand that I will receive a signed copy of this form**

_____ Participant/Legal Representative's name (printed)	_____ Signature	_____ Date: (DD MMM YYYY) and Time (24 hr clock)
_____ Witness name (printed)	_____ Signature	_____ Date: (DD MMM YYYY) and Time (24 hr clock)
_____ Name of person obtaining consent (printed)	_____ Signature	_____ Date: (DD MMM YYYY) and Time (24 hr clock)
_____ Name of Investigator (printed)	_____ Signature	_____ Date: (DD MMM YYYY) and Time (24 hr clock)