Public reporting burden for this collection of information is estimated to average 3 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information.  **An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.**  Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to: NIH, Project Clearance Branch, 6705 Rockledge Drive, MSC 7974, Bethesda, MD 20892-7974, ATTN: PRA (0925-xxxx\*).  Do not return the completed form to this address.

**OMB Number: 0925-XXXX**

 **OMB Expiration Date: TBD**

**Interactive Informed Consent for Pediatric Clinical Trials**

**Clinical trial concepts Pre-test (baseline) interview:**

1. Have you (or your child) ever been in a research study before?

 Yes No

1. When researchers talk about clinical studies, they often use words to describe how the study will be done. Can you tell me what the following words mean?

Clinical trial

Randomization

Placebo

Blinding