Public reporting burden for this collection of information is estimated to average 9 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

APPENDIX B

Interactive Informed Consent for Pediatric Clinical Trials

Perceptions Questionnaire:

ceptions Questionnane.											
1.	If this had been a real study how likely would you (or your child) have been to participate?										
	0 Not like	1 ely	2	3	4	5	6	7		9 Extreme	10 ly likely
2.	Please rate the quality of the information presented regarding the study										
	0 Poor q	1 Juality	2	3	4	5	6	7		9 Excellen	10 t quality
3.	Please rate your ability to follow the information presented regarding the study								study		
	0 Imposs	1 sible to	2 follow	3	4	5	6	7 Extrem	8 nely ea	9 sy to foll	10 low
4.	How e study?		was the	e prese	ntation	of infor	mation	in helpi	ng you	unders	tand the
	\square Not effective at all \square Somewhat effective \square Extremely effective										
5.	Was th	Was the amount of information that you received about the study :									

	☐ Too little		□ Just right			□ Too much					
6.	Overal	l, how o	clearly v	vas the	informa	tion abo	out stud	y prese	nted?		
	□ not a	at all cle	ear		□ fairly	y clear		□ very	clear		
7.		rate h the stud	-	y it was	to use	the co	mputer	prograr	n to ob	otain info	ormation
Ext	0 tremely		2 t	3	4	5	6	7		9 Extreme	10 ly easy
8.	Please study	rate y	our ove	erall sat	isfactio	n with t	he com	iputer p	rogran	n descril	oing the
No	0 t at all s	1 satisfied	2	3	4	5	6	7	8 Extrer	9 nely sati	10 sfied

9. If you are asked to be in a real clinical study in the future, would you prefer the information about the study to be presented as: written information only, verbal information only, computer program only (like this one), written and verbal

information, computer program and verbal information