Project Title: **Date Submitted:**		Yes/No N/A
1	The purpose of the research is clearly stated	
2	The time it will take the subject to participate in the study	
3	Procedures to be followed and which procedures are experimental	
4	Reasonably foreseeable risks or discomforts	
5	Benefits to the subject or others	
6	Alternative procedures for treatment	
7	Confidentiality of records which identify the subject	
8	If more than minimal risk: availability of medical treatment in the case of research-related injury, including who will pay for the treatment and whether other financial compensation is available, if there is a potential for emotional/psychological risk, you must be able to provide proper counseling and indicate who will provide the counseling	
9	Whom to contact for answers to pertinent questions about the research and the research subject's rights, and whom to contact in the event of a research-related injury	
10	Participation is voluntary; refusal will involve no penalty or loss of benefits to which the subject is otherwise entitled	
11	Risks to the subject (or the fetus or embryo if the subject is or may become pregnant)	
12	Are children, minors, pregnant women, fetuses, human in vitro fertilization, prisoners, employees, military persons and students in hierarchical organizations, terminally ill, comatose, physically and intellectually challenged individuals, institutionalized or elderly individuals, visually or hearing impaired, ethnic minorities, refugees, and economically and educationally disabled individuals involved in this study?	
13	Participation may be terminated by the investigator without regard to the subject's consent; procedure(s) for termination	
14	Cost of participation in the subject	
15	Compensation to the subject	
16	Significant findings will be provided to the subject upon request	
17	Approximate number of subjects in the study	
18	Right to have a copy of the consent form	

Comments: