Behavioral Health Study Informed Consent Waiver Request

1. Study Information

Study II	tle	
Principa	Il Investigator	
Institutio	n/Department	
Contact	Information	
Protocol	Number (if applicable)	
2. De	scription of the Study	
Provide a brief summary of the research project:		
3. Re	quest for Waiver of Informed Consent	
Describe the reason for requesting an informed consent waiver:		
	stification for Waiver (45 CFR 46.116 Criteria)	
1.	Explain how the research involves no more than minimal risk to participants:	
2.		
	Explain why the waiver will not adversely affect the rights and welfare of participants:	
	Explain why the waiver will not adversely affect the rights and welfare of participants:	
	Explain why the waiver will not adversely affect the rights and welfare of participants:	
3.	Explain why the waiver will not adversely affect the rights and welfare of participants: Explain why the research could not practicably be carried out without the waiver:	
3.		

4. Explain, if appropriate, how pertinent information will be provided to participants after participation:

5. Additional Information			
Include any	y additional information relevant to the waiver request:		
6. Cert	tification		
Name of P	Principal Investigator		
Date			