

STUDY SUBMISSION FORM: ADDITIONAL STUDY LOCATION			
1.	Sponsor:	Protocol No.:	
2.	Principal Investigator:		
A.	Name of additional study location:		
B.	Address of additional study location :		
C.	Site Phone:		
D.	Do you want this location's phone number listed on the consent form?	<input type="checkbox"/> No <input type="checkbox"/> Yes	
E.	Contact person:	Email:	
F.	Approximate distance from the main site: miles		
G.	Will the Principal Investigator be physically supervising at this location?	<input type="checkbox"/> No <input type="checkbox"/> Yes	
3.	The following Sub-investigator(s) will be working out of this location:		
4.	Nature of the additional facility:		
A.	<input type="checkbox"/> Medical office	<input type="checkbox"/> Research Facility	<input type="checkbox"/> Imaging center
	<input type="checkbox"/> Hospital*	<input type="checkbox"/> Clinic	<input type="checkbox"/> Nursing home
	<input type="checkbox"/> University*	<input type="checkbox"/> Other:	
<i>* Include letter from appropriate official acknowledging Alpha IRB's review of this study.</i>			
B.	Does the site have adequate resources, including staff to conduct this study? <input type="checkbox"/> Yes <input type="checkbox"/> No - if no, explain:		
C.	What equipment and resources are available at this site to treat emergencies? (e.g. ACLS certified staff, crash cart, access to 911, etc.):		
D.	Name of the nearest emergency facility to be used in the event of an emergency: Distance of emergency facility from this study site:		
5.	Informed Consent:		
A.	Will subjects be consented at this site? <input type="checkbox"/> No <input type="checkbox"/> Yes		
	If yes, who will explain the study to subjects and obtain consent at this site? <input type="checkbox"/> PI <input type="checkbox"/> Sub-I <input type="checkbox"/> Research Coordinator <input type="checkbox"/> Other:		
6.	Additional comments:		
Principal Investigator Signature:			Date: