

## PATIENT INFORMATION FOR CLINICAL STUDIES

First Name	Middle Initia	l Last Na	me		
SS#	Date of Birth		Age		
Home address		City	ZIP Cod	de	
Home Phone	Home PhoneWork Phone		Mobile Phone		
Employer	Occupation				
Parent/Guardian Name (for minors)		Relationship			
Contact in case of emergency		Phone			
Where did you learn about this study?		Primary Physician's Name			
to understand the specifics about to a copy of the signed consent. The stort time and travel, if any, will be visits completed, unless other arra. If enrolled in the clinical trial, during the study or thereafter, I a notify the office of my decision. If study participation involves no cost.	regulations and good clinical prosent prior to any procedures at the study. My decision to partic screening visit alone is performed provided at my final study visiting mements are made and explain, I must adhere to the follow up m free to ask them. I also know any changes occur in my health st to me. Costs of other treatments.	ractices standards, and after reading in ipate can be delay ed at no cost and we cor thereafter and ned to me.  schedule of visits at that I can stop study, medications, or a sents not study relations.	t I may ask as man ed if I need additio without reimbursen prorated dependinand procedures. If ady participation a address, I will notified are my respons	y questions as I need nal time. I will receiv nent. Reimbursement of of the number of I have questions t any time and will fy the office. The sibility.	
Would you like to be contact	red for future studiesyes	no <b>Email addres</b> s	5		
Participant's Signature			Date		
Parent/Guardian's Signature	gnature (For minors)		Date	Rev 06-01-15	