UMCIRB #	PI:
-	Iedical Center Institutional Review Board r Waiver or Alteration of Authorization
 Select the types Protected Health Information ([]] Billing records 	PHI) to be collected: [D] Hospital/medical records (in and out-patient)
[] Mental Health records	[D] Lab, pathology and/or radiology results
[]] Physician/clinic records	[]] PHI previously collected for research purposes
[]] SODM Axium records	[]] Other:
2. Select the responses below for all the identifier	s to be captured in the research study:
[]] Postal address	[]] Health plan numbers []] Telephone
[]] Account /medical record number	[] Name [] Internet P
[D] Certificate/license number	[D] Name of employers [D] Web university
[D] Name of relatives	[D] Photographic images [D] Fax numb
[D] E-mail address	[D] Finger or voice prints
[]] Any device or vehicle identifiers an	d serial numbers, including license plate numbers
	arga data data of daath all agas over 80
[D] Date of birth, admission date, disch	arge date, date of death, an ages over 89

[D] Telephone number []] Internet Protocol (IP) address number [D] Web universal resource locator (URL) [] Fax number

mitted to be collected in reliance on a waiver ed, the study subject must be given a written disclosure which (i) states that providing social security number is not required; and (ii) describes the purpose for which the social security number will be used.

3. Select the response below on how participant's Protected Health Information (PHI) is protected against improper use or disclosure:

[D] Research team members will sign a Confidentiality Agreement
 [D] The information will not be shared unless it is stripped of all <u>18 identifiers</u>

[]] The information will be shared with a random code as outlined in the research protocol.

4. Explain the data management measures to protect the confidentiality of participant's data such as storage and access issues, including (i) safeguards for storage of any identifiers on computer workstations; and (ii) safeguards for storage of any identifiers on laptop computers, flash drives or any other portable electronic device, as applicable.

5.	Data will be stripped of all <u>identifiers</u> upon completion of:
	[D] subject participation [D] data analysis
	[D] FDA approval [D] specimen processing
	[]] other (please specify):
	OR
	Identifiers will be retained indefinitely because:
	[]] the study is longitudinal
	[]] of federal requirements (specify):
	[] other (specify):
6.	Provide any additional explanations on why the use/disclosure of PHI involves no more than minimal risk to participant privacy

7. Explain why the participant's written Authorization cannot be attained and, therefore, research cannot be practicably carried out without the Waiver or Alteration of Authorization. If required elements of a valid HIPAA Authorization are being omitted from the Authorization process, please list them and justify their exclusion.

8. Select the response or explain why research can not practicably be conducted without the participant's PHI.
 []] PHI is needed to identify eligibility for the study
 []] PHI is the forms of the study (a g an original studies)

[]] PHI is the focus of the study (e.g. – epidemiological studies)

[D] Other (specify):

If there is additional information that cannot fit within the fields on this form, or ancillary documentation related to this request, please attach it to this form.

I verify that protected health information will not be re-used or disclosed to any other person or entity, except as required by law, research oversight, per the terms of a valid HIPAA authorization or those uses outlined above. I will only collect the information as specified above, and limit access to that information to the greatest extent possible as previously described. I will destroy the identifier at the earliest opportunity consistent with the conduct of the research as specified above. Additionally, a comprehensive listing of individuals whose PHI will be accessed as part of this review shall be documented through completing electronic accounting of disclosures within the electronic health record (EHR).

Principal Investigator's signature

References: 45 CFR 160 and 164, Standards for Privacy of Individually Identifiable Health Information; Final Rule NC Gen. Stat. Sect. 132-1.10 (Social Security Numbers and Personal Identifying Information)

Date