

FWA #: **FWA00001380**
Institution: **Upstate Affiliate Organization (UAO) dba Greenville Health System**
Expires: **09/02/2021**

OMB No. 0990-0278
Approved for use through August 31, 2017

Federalwide Assurance (FWA) for the Protection of Human Subjects

1. Institution Filing Assurance

Legal Name: **Upstate Affiliate Organization (UAO) dba Greenville Health System**
City: **Greenville** State/Province: **SC** Country: **USA**

2. Institutional Components

List below all components over which the Institution has legal authority that operate under a different name. Also list with an asterisk (*) any alternate names under which the Institution operates.

NOTE: The Signatory Official signing this Assurance must be legally authorized to represent the Institution providing this Assurance and all components listed below.

Name of Component or Alternate Names Used	City	State (or Country if Outside U.S.)	
Greenville Mem Med Campus	GREENVILLE	SC	A
Ctr for Family Med	GREENVILLE	SC	A
Children's Hosp	GREENVILLE	SC	A
Cross Creek Surgery Ctr	GREENVILLE	SC	A
Patewood Med Campus	GREENVILLE	SC	A
Greer Med Campus	GREER	SC	A
Simpsonville Med Campus	SIMPSONVILLE	SC	A
Traveler's Rest Med Campus	TRAVELER'S REST	SC	A
Baptist Easley Hospital	EASLEY	SC	A
Greenville Health System Cancer Institute - Greenville CCOP- International Drive	GREENVILLE	SC	A
Greenville Health System Cancer Institute-Seneca	SENECA	SC	A
Greenville Health System Cancer Institute-Andrews	GREENVILLE	SC	A
Greenville Health System Cancer Institute-Butternut (Grove)	GREENVILLE	SC	A
Greenville Health System Cancer Institute-Easley	EASLEY	SC	A
Greenville Health System Cancer Institute-Faris	GREENVILLE	SC	A
Greenville Health System Cancer Institute -Greer	GREER	SC	A
Greenville Health System Cancer	SPARTANBURG	SC	A

Institute-Spartanburg Laurens County Memorial Hospital	CLINTON	SC	A
Oconee Medical Center	SENECA	SC	A
Greenville Health System Cancer Institute - BI-LO Charities Children's Cancer Center	GREENVILLE	SC	A
Greenville Health System Cancer Institute-Laurens	Laurens	SC	A

3. Statement of Principles

This Institution assures that all of its activities related to human subjects research, regardless of the source of support, will be guided by the following statement of principles governing the institution in the discharge of its responsibilities for protecting the rights and welfare of human subjects of research conducted at or sponsored by the institution. (indicate below)

The Belmont Report

4. Applicability

(a) This Assurance applies whenever this Institution becomes engaged in human subjects research conducted or supported by any U.S. federal department or agency that has adopted the U.S. Federal Policy for the Protection of Human Subjects (also known as the Common Rule), unless the research is otherwise exempt from the requirements of the Common Rule or the department or agency conducting or supporting the research determines that the research shall be conducted under a separate assurance.

(b) Optional: This Institution elects to apply the following to all of its human subjects research regardless of the source of support, except for research that is covered by a separate assurance:

The Common Rule (see section 3 of the Terms of the FWA for a list of U.S. federal departments and agencies that have adopted the Common Rule and the applicable citations to the Code of Federal Regulations)

5. Assurance of Compliance with the Terms of the Federalwide Assurance

(a) This Institution assures that whenever it engages in research to which this Assurance applies, it will comply with the Terms of the Federalwide Assurance (contained in a separate document on the Office for Human Research Protections (OHRP) website).

6. Designation of Institutional Review Boards (IRBs)

This Institution assures that it will rely upon only IRBs registered with OHRP for review of research to which this FWA applies. This institution (a) designates the following internal IRB(s) for review of research under this Assurance; or (b) does not have an internal IRB and designates the following external IRB for review of all research to which this FWA applies or, if multiple external IRBs are relied upon, the following external IRB that reviews the largest percentage of research to which this FWA applies.

NOTE: Institutions designating internal IRBs do not need to designate any of the external IRBs upon which it relies.

HHS IRB Registration Number	Name of IRB as Registered with HHS	Is the IRB Internal or External to the Institution?
IRB00000027	Medical U South Carolina IRB #1	E
IRB00000028	Medical U South Carolina IRB #2	E
IRB00000240	U of SOUTH CAROLINA IRB #1	E
IRB00000481	CLEMSON U IRB #1	E
IRB00000533	Western IRB #1-5, 16, 17 & #35 - All U.S. Panels	E
IRB00000715	Palmetto Health IRB #1 - PH IRB #1	E
IRB00000781	National Cancer Inst Central IRB #1 (Adult)	E
IRB00000805	Greenville Hosp System IRB #1	I

IRB00000873	Greenville Hosp System IRB #2	I
IRB00000915	Palmetto Health IRB #2 - PH IRB #2	E
IRB00001113	U.S. Oncology, Inc. IRB #1 - IRB	E
IRB00001130	U.S. Oncology, Inc. IRB #2 - IRB	E
IRB00001369	Spartanburg Regional Healthcare System IRB #1 - 1	E
IRB00001377	Medical U South Carolina IRB #3	E
IRB00001385	Anderson Area Med Ctr IRB #1	E
IRB00002227	Greenville Hosp System IRB #3	I
IRB00003223	Self Regional Healthcare IRB #1	E
IRB00003226	Quorum Review Inc. IRBs 1-4	E
IRB00004296	National Cancer Inst Central IRB #2 (Pediatric)	E

7. Human Protections Administrator (e.g., Human Subjects Administrator or Human Subjects Contact Person)

First Name: **Christopher** Middle Initial: **C** Last Name: **Wright**
 Degrees or Suffix: **MD** Institutional Title: **Medical Director, Office of Human Research Protection**
 Institution: **Greenville Health System**
 Telephone: **864 455-8997** FAX: **864 455-6766** E-Mail: **CWright2@ghs.org**
 Address: **Health Science Administration Building**
 701 Grove Road
 City: **Greenville** State/Province: **SC** Country: **USA**

8. Signatory Official (i.e., Official Legally Authorized to Represent the Institution)

I have read and agree to the Terms of the Federalwide Assurance.

I recognize that providing research investigators, IRB members and staff, and other relevant personnel with appropriate initial and continuing education and training about human subject protections will help ensure that the requirements of this Assurance are satisfied.

Acting officially in an authorized capacity on behalf of this Institution and with an understanding of the Institution's responsibilities under this Assurance, I assure protections for human subjects as specified above. The IRB(s) that this institution relies upon will comply with the Terms of the Federalwide Assurance when reviewing research covered by this Assurance and possess appropriate knowledge of the local context in which this Institution's research will be conducted.

All information provided with this Assurance is up-to-date and accurate. I am aware that false statements could be cause for invalidating this Assurance and may lead to other administrative or legal action.

Signature: **Spence Taylor MD**

Date: **09/01/2016**

First Name: **Spence** Middle Initial: Last Name: **Taylor**

Degrees or Suffix: **MD** Institutional Title: **President, Greenville Health System**

Institution: **Greenville Health System**

Telephone: **864 455-7992** FAX: **864 455-8404** E-Mail: **staylor@ghs.org**

Address: **Health Science Administration Building**
701 Grove Road

City: **Greenville** State/Province: **SC** Country: **USA**

9. FWA Approval

The Federalwide Assurance for the Protection of Human Subjects for Institutions Within the United States submitted to HHS by the above Institution is hereby approved.

Assurance Number: **FWA00001380**

Expiration Date: **09/02/2021**

Signature of HHS Approving Official: **Gail Holloway**

Date: **09/02/2016**

According to the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMB control number. The valid OMB control number for this information collection is 0990-0278 . The time required to complete this information collection is estimated to average 30 minutes per response, including the time to review instructions, gather the data needed, and complete and review the information collection. If you have comments concerning the accuracy of the time estimate(s) or suggestions for improving this form, please write to: U.S. Department of Health & Human Services, OS/OCIO/PRA, 200 Independence Ave., S.W., Suite 336-E, Washington D.C. 20201, Attention: PRA Reports Clearance