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About the USPHS Syphilis Study

Where the Study Took Place

The study took place in Macon County, Alabama, the county seat of Tuskegee referred to as the "Black Belt" because of its rich soil and vast number of black sharecroppers who were the economic backbone of the region. The research itself took place on the campus of Tuskegee

What it Was Designed to Find Out

The intent of the study was to record the natural history of syphilis in Blacks. The study was called the "Tuskegee Study of Untreated Syphilis in the Negro Male." When the study was initiated there were no proven treatments for the disease. Researchers told the men participating in the study that they were to be treated for "bad blood." This term was used locally by people to describe a host of diagnosable ailments including but not limited to anemia, fatigue, and syphilis.

Who Were the Participants

A total of 600 men were enrolled in the study. Of this group 399, who had syphilis were a part of the experimental group and 201 were control subjects. Most of the men were poor and illiterate sharecroppers from the county.

What the Men Received in Exchange for Participation

The men were offered what most Negroes could only dream of in terms of medical care and survivors insurance. They were enticed and enrolled in the study with incentives including: medical exams, rides to and from the clinics, meals on examination days, free treatment for minor ailments and guarantees that provisions would be made after their deaths in terms of burial stipends paid to their survivors.

Treatment Withheld

There were no proven treatments for syphilis when the study began. When penicillin became the standard treatment for the disease in 1947 the medicine was withheld as a part of the treatment for both the experimental group and control group.

How/Why the Study Ended

On July 25, 1972 Jean Heller of the Associated Press broke the story that appeared simultaneously both in New York and Washington, that there had been a 40-year nontherapeutic experiment called "a study" on the effects of untreated syphilis on Black men in the rural south.

Between the start of the study in 1932 and 1947, the date when penicillin was determined as a cure for the disease, dozens of men had died and their wives, children and untold number of others had been infected. This set into motion international public outcry and a series of actions initiated by U.S. federal agencies. The Assistant Secretary for Health and Scientific Affairs appointed an Ad Hoc Advisory Panel, comprised of nine members from the fields of health administration, medicine, law, religion, education, etc. to review the study.

While the panel concluded that the men participated in the study freely, agreeing to the examinations and treatments, there was evidence that scientific research protocol routinely applied to human subjects was either ignored or deeply flawed to ensure the safety and well-being of the men involved. Specifically, the men were never told about or offered the research procedure called informed consent. Researchers had not informed the men of the actual name of the study, i.e. "Tuskegee Study of Untreated Syphilis in the Negro Male," its purpose, and potential consequences of the treatment or non-treatment that they would receive during the study. The men never knew of the debilitating and life threatening consequences of the treatments they were to receive, the impact on their wives, girlfriends, and children they may have conceived once involved in the research. The panel also concluded that there were no choices given to the participants to quit the study when penicillin became available as a treatment and cure for syphilis.

Reviewing the results of the research the panel concluded that the study was "ethically unjustified." The panel articulated all of the above findings in October of 1972 and then one month later the Assistant Secretary for Health and Scientific Affairs officially declared the end of the Tuskegee Study.

Class-Action Suit

In the summer of 1973, Attorney Fred Gray filed a class-action suit on behalf of the men in the study, their wives, children and families. It ended a settlement giving more than \$9 million to the study participants.

The Role of the US Public Health Service

In the beginning of the 20th Century, the U.S. Public Health Service (PHS) was entrusted with the responsibility to monitor, identify trends in the heath of the citizenry, and develop interventions to treat disease, ailments and negative trends adversely impacting the health and

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The PHS began working with Tuskegee Institute in 1932 to study hundreds of black men with syphilis from Macon County, Alabama.

Compensation for Participants

As part of the class-action suit settlement, the U.S. government promised to provide a range of free services to the survivors of the study, their wives, widows, and children. All living participants became immediately entitled to free medical and burial services. These services were provided by the Tuskegee Health Benefit Program, which was and continues to be administered by the Centers for Disease Control and Prevention in their National Center for HIV, STD and TB Prevention.

1996 Tuskegee Legacy Committee

In February of 1994 at the Claude Moore Health Sciences Library in Charlottesville, VA, a symposium was held entitled "Doing Bad in the Name of Good?: The Tuskegee Syphilis Study and Its Legacy." Resulting from this gathering was the creation of the Tuskegee Syphilis Study Legacy Committee which met for the first time in January 18th & 19th of 1996. The committee had two goals; (1) to persuade President Clinton to apologize on behalf of the government for the atrocities of the study and (2) to develop a strategy to address the damages of the study to the psyche of African-Americans and others about the ethical behavior of government-led research; rebuilding the reputation of Tuskegee through public education about the study, developing a clearinghouse on the ethics of scientific research and scholarship and assembling training programs for health care providers. After intensive discussions, the Committee's final report in May of 1996 urged President Clinton to apologize for the emotional, medical, research and psychological damage of the study. On May 16th at a White House ceremony attended by the men, members of the Legacy Committee and others representing the medical and research communities, the apology was delivered to the surviving participants of the study and families of the deceased.

Staff Directory Campus Map



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