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The New York Times

Syphilis Victims in U.S. Study Went Untreated for 40 Years

By JEAN HELLER
The Associated Press

WASHINGTON, July 25—For 40 years the United States Public Health Service has conducted a study in which human beings with syphilis, who were induced to serve as guinea pigs, have gone without medical treatment for the disease and a few have died of its late effects, even though an effective therapy was eventually discovered.

The study was conducted to determine from autopsies what the disease does to the human body.

Officials of the health service who initiated the experiment have long since retired. Current officials, who say they

have serious doubts about the morality of the study, also say that it is too late to treat the syphilis in any surviving participants.

Doctors in the service say they are now rendering whatever other medical services they can give to the survivors while the study of the disease's effects continues.

Dr. Merlin K. DuVal, Assistant Secretary of Health, Education and Welfare for Health and Scientific Affairs, expressed shock on learning of the study. He said that he was making an immediate investigation.

The experiment, called the Tuskegee Study, began in 1932 with about 600 black men,



The Belmont Report-1979

- ☞ Report of the meeting of the National Commission for the Protection of Human Subjects of Biomedical and Behavioral Research at the Belmont Conference Center at the Smithsonian
- ☞ By this time IRB have been established in most institutions and having difficulties implementing all the rules and directives from Nuremberg and Helsinki

The Belmont Report, issued in 1979 by the National Commission for the Protection of Human Subjects of Biomedical and Behavioral Research, provides the ethical framework for the Federal Regulations designed to protect human research subjects.

The Belmont Report is available in English, Chinese, Spanish, and French. Click the language button to return here to continue.

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The Belmont Report

Office of the Secretary

Director/Policy and Guidelines for the Protection of Human Subjects of Research

The National Commission for the Protection of Human Subjects of Biomedical and Behavioral Research

April 18, 1979

DEPARTMENT OF HEALTH, EDUCATION AND WELFARE
OFFICE OF THE SECRETARY

CONFIDENTIAL - On July 12, 1979, the National Research Act (P.L. 96-359) was signed into law, creating the National Commission for the Protection of Human Subjects of Biomedical and Behavioral Research. One of the purposes of the Commission is to study and report on the ethical principles that should underlie research involving human subjects and to develop guidelines which should be followed by all researchers who conduct research involving human subjects. In carrying out this study, the Commission has been directed to consider the relationship between individual and collective research and the responsibility of researchers to society. On the role of government in the development of the Commission, the Department of Health, Education and Welfare has issued guidelines for the development of the Commission and its responsibilities. The National Commission for the Protection of Human Subjects of Biomedical and Behavioral Research is a Federal agency established by Executive Order 12046, dated February 18, 1977, and is located at the Department of Health, Education and Welfare, Washington, D.C. 20460.

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