Hansen’s Disease

Feared For Millenia

Since the earliest days of civilization the world over, leprosy has been regarded as a contagious, mutilating and incurable scourge. Humankind’s reaction to this particular disease has left behind a terrible black mark on world history and our collective memory – one of disfigurment, rejection and expulsion from society.

A Misunderstood Disease

Leprosy has been one of humankind’s most feared diseases, yet it is one of the least communicable diseases, with only about 5% of the world’s population even being susceptible. For millennia, people did not understand the disease and its transmission. Its appearance, seemingly out of nowhere, made many think the disease was punishment from God for being physically and spiritually unclean. Such beliefs have caused leprosy sufferers to be shunned by the rest of society and forced to live in isolation.

Today, we know the disease to be caused by a bacteria transmitted through direct person-to-person contact over a prolonged period of time. The disease attacks the nerves, causing severe skin and eye damage. The incubation period is much longer than for other sicknesses. Months, or even years, can pass before symptoms begin to appear.

Untreated patients suffer from sores on the skin, the reabsorption of bone and cartilage by the body accompanied by a loss of feeling and paralysis in the limbs. Over time, these effects can lead to the body’s permanent disfigurement. Eventually the weakened body’s immune system is unable to fight off other sicknesses, resulting in death.

A Sickness is Criminalized

When or how leprosy came to Hawai‘i is unknown, but it appears in records as early as the 1830s. Hawaiians, having no immunities to introduced diseases, were particularly vulnerable to infection. By the mid-1800s, Hawaiians suffered death and disfigurement at alarming rates. Fearing further spread of the disease, the Kingdom of Hawai‘i set aside land for the purpose of confining leprosy patients.

Beginning in 1865, police and district justices were required to arrest any persons suspected of having the sickness. Kalihi Hospital and Detention Station in Honolulu evaluated “suspects”. Those with advanced cases were sent to Kalawao, a settlement on the isolated peninsula protruding from Molokai island’s north shore.

Families were torn apart by the policy. People fled or hid family from authorities out of fear they would be taken away, never to be seen again. The biblical shame associated with having a diseased family member caused many families to disown sick relatives. Many sick people denied their own family members in order to protect them from being suspected as potential carriers.

King Kamehameha V signed into law the “Act to Prevent the Spread of Leprosy” on January 3, 1865. It remained in effect for 103 years.
The Cause Identified

On the other side of the world, in 1873 Norway, Dr. Gerhard Armaturu Hansen made a breakthrough discovery. He identified the cause of leprosy in his laboratory -- a simple bacteria which attacks the nerves. The discovery of this microorganism was the first step in finding an effective treatment. Leprosy is now called Hansen's Disease in an effort to battle the centuries-old stigma and also to honor its discoverer's contribution to medical science.

Hope for a Cure

During the years following Hansen's discovery many treatments for the disease were explored in Hawaii. Experimental treatments were commonplace throughout the Hawaiian Islands and rest of the world. Father Damien explored the therapeutic benefits of Japanese Goto baths. The bathing regime involved daily immersions in a hot bath with herbs, coupled with the taking of herbal medicines. Though the baths did bring relief they were not a cure.

In an effort to find a cure for the disease the United States government established a research and treatment facility. In July 1909, the U.S. Leprosy Investigation Station (seen above) opened its doors. The state-of-the-art facility was established to treat patients and conduct scientific investigations into finding a cure. No expense was spared on the 68,000 square foot campus, which included flushable toilets, electricity, and refrigeration units. However, only nine patients volunteered to participate in the experimental treatments and be confined to hospital grounds. Such a small number of participants did not bode well for research and experimentation. Similar research was also being done at the same time at Kalihi Hospital in Honolulu and eventually all research was moved there. The U.S. Leprosy Investigation Station was abandoned in 1913.

In the early 1900s chaulmoogra oil was the primary treatment for the disease. The oil, extracted from seeds of the chaulmoogra tree had been used throughout Southeast Asia in Hansen's Disease treatments. The oil was administrated externally and by injection. Many of the disease's effects appeared to disappear and it was thought to be the source for a possible cure. However, by the 1920s not enough cases responded to the treatment and hope for a cure faded. It was not until the mid-20th century that a cure for Hansen's Disease was finally found and brought to the Hawaiian Islands.

The End of Isolation

In 1941, Dr. Guy Faget used a newly developed sulfone drug, called Promin, to treat patients at the U.S. Public Health Service National Leprosarium in Carville, Louisiana. After several months of treatment with the new drug, the disease's physical effects and capacity for transmission to others were completely arrested. A cure had finally been found.

Almost thirty years would pass before Hawaii's isolation policy was finally abolished in 1969. More than 8,000 people died at Kalaupapa during the 103 year period of isolation. For many people of Hawaiian descent, a visit to Kalaupapa is a chance to research and reconnect with family that may have been "lost" for generations.

Today at Kalaupapa and in Honolulu, people living with Hansen's Disease continue to receive treatment from the State of Hawaii Department of Health. Patients still living at Kalaupapa now choose to do so. During their lives they have seen major advances in medical knowledge, drug treatments, and social attitudes regarding the disease.

The World Health Organization now estimates there are 1.15 million registered cases of Hansen's Disease in over 55 different countries. Around the world today an on-going effort to end discrimination against peoples living with Hansen's Disease is underway through public education, community acceptance, and patient demands for their human rights.