

Vogt-Koyanagi-Harada disease is a rare disorder of unknown origin that affects many body systems, including as the eyes, ears, skin, and the covering of the brain and spinal cord (the meninges). The most noticeable symptom is a rapid loss of vision. There may also be neurological signs such as severe headache, vertigo, nausea, and drowsiness. Loss of hearing, and loss of hair (alopecia) and skin color may occur along, with whitening (loss of pigmentation) of the hair and eyelashes (poliosis). Vogt-Koyanagi-Harada disease is a rare disease that affects males and females in equal numbers. The disorder is more prevalent in Oriental, Hispanic, and American Indian populations than in people who trace their ancestry to northern Europe. Onset typically occurs at around 30 or 40 years of age, but cases have been reported among children as young as four years old.