Vogt-Koyanagi-Harada disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Vogt-Koyanagi-Harada disease. ORPHA:3437

Vogt-Koyanagi-Harada disease is a bilateral, chronic, diffuse granulomatous panuveitis typically characterized by serous retinal detachment and frequently associated with neurological (meningitis), auditory, and dermatological alterations.