Stargardt disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Stargardt disease. ORPHA:827

Stargardt disease, also known as Stargardt 1 (STGD1), is an autosomal recessive form of retinal dystrophy that is usually characterized by a progressive loss of central vision associated with irregular macular and perimacular yellow-white fundus flecks, and a so-called ''beaten bronze'' atrophic central macular lesion.