Open Peer Review on Qeios

Holt-Oram Syndrome

National Cancer Institute

Source

National Cancer Institute. <u>Holt-Oram Syndrome</u>. NCI Thesaurus. Code C125592.

A rare, autosomal dominant inherited syndrome caused by mutations in the TBX5 gene. It is characterized by skeletal abnormalities in the upper limbs and heart abnormalities.