

Open Peer Review on Qeios

Holt-Oram syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Holt-Oram</u> <u>syndrome</u>. ORPHA:392

Holt-Oram syndrome (HOS) is the most common form of heart-hand syndrome (see this term) and is characterized by skeletal abnormalities of the upper limbs and mild-tosevere congenital cardiac defects.

Qeios ID: KDAZMO · https://doi.org/10.32388/KDAZMO