

Open Peer Review on Qeios

## Thomas syndrome

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Thomas</u> <u>syndrome</u>. ORPHA:3316

Thomas syndrome is characterised by renal anomalies, cardiac malformations and cleft lip or palate. It has been described in six patients. Transmission was suggested to be autosomal recessive.

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