

Open Peer Review on Qeios

Matthew-Wood syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Matthew-Wood syndrome</u>. ORPHA:2470

Matthew-Wood syndrome is a rare clinical entity including as main characteristics anophthalmia or severe microphthalmia, and pulmonary hypoplasia or aplasia.

Qeios ID: EYPB68 · https://doi.org/10.32388/EYPB68