## Open Peer Review on Qeios

## **RHYNS syndrome**

## INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>RHYNS</u> <u>syndrome</u>. ORPHA:140976* 

RHYNS syndrome is characterised by the association of retinitis pigmentosa, hypopituitarism, nephronophthisis, and skeletal dysplasia.