## Open Peer Review on Qeios

## Pierson Syndrome

National Cancer Institute

## Source

National Cancer Institute. <u>Pierson Syndrome</u>. NCI Thesaurus. Code C128145.

An autosomal recessive disorder caused by mutation(s) in the LAMB2 gene, encoding laminin subunit beta-2. It is characterized by congenital nephrotic syndrome with diffuse mesangial sclerosis and distinct ocular abnormalities.