## Open Peer Review on Qeios

## Fraser Syndrome

National Cancer Institute

## Source

National Cancer Institute. Fraser Syndrome. NCI Thesaurus. Code C118436.

A rare, autosomal recessive inherited disorder caused by mutations in the FRAS1, FREM2, or GRIP1 genes. It is characterized by the presence of cryptophthalmos, cutaneous syndactyly, and genitourinary abnormalities.