## Open Peer Review on Qeios

## **CALFAN Syndrome**

National Cancer Institute

## Source

National Cancer Institute. <u>CALFAN Syndrome</u>. NCI Thesaurus. Code C159655.

A condition caused by biallelic mutation(s) in the SCYL1 gene, encoding N-terminal kinaselike protein. It is characterized by peripheral neuropathy, cerebellar atrophy, ataxia, and recurrent episodes of liver failure.