

Open Peer Review on Qeios

## Feingold Syndrome

National Cancer Institute

## Source

National Cancer Institute. <u>Feingold Syndrome</u>. NCI Thesaurus. Code C74987.

A rare autosomal dominant syndrome caused by mutations in the MYCN oncogene. It is characterized by microcephaly, limb abnormalities, esophageal and/or duodenal atresia.

Qeios ID: QISO2X · https://doi.org/10.32388/QISO2X