# CAD-CDG

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. CAD-CDG. ORPHA:448010*

CAD-CDG is a rare congenital disorder of glycosylation caused by mutations in the CAD gene and characterized by epileptic encephalopathy, global developmental delay, normocytic anemia and anisopoikilocytosis. Loss of acquired skills in early childhood is present and natural disease course can be lethal in early childhood.