# Chromosome 2q37 Deletion Syndrome

National Cancer Institute

## Source

*National Cancer Institute. Chromosome 2q37 Deletion Syndrome. NCI Thesaurus. Code C129021.*

A syndrome of high phenotypic variability caused by contiguous gene deletions in 2q37. The inheritance is autosomal dominant. The condition may be characterized by brachydactly type E; mental retardation; short stature; and other skeletal, cardiovascular, and neurologic manifestations.