Trichorhinophalangeal Syndrome Type II

National Cancer Institute

Source

National Cancer Institute. Trichorhinophalangeal Syndrome Type II. NCI Thesaurus. Code C75118.

A rare, usually sporadic and less frequently familiar syndrome caused by deletions on the long arm of chromosome 8. It is characterized by distinctive facial appearance (sparse hair, pear-shaped nose, and large ears), multiple exostoses, redundant skin, and mental retardation.