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Pseudoachondroplasia

National Cancer Institute

Source

National Cancer Institute. *Pseudoachondroplasia*. NCI Thesaurus. Code C118635.

A rare, autosomal dominant inherited disorder caused by mutations in the COMP gene. It is characterized by short stature, short arms and legs, waddling walk, osteoarthritis, and limited range of motion at the elbows and hips.