Andersen-Tawil syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Andersen-Tawil syndrome. ORPHA:37553

Andersen's syndrome (AS) is a rare disorder characterized by periodic muscle paralysis, prolongation of the QT interval with a variety of ventricular arrhythmias (leading to predisposition to sudden cardiac death) and characteristic physical features: short stature, scoliosis, low-set ears, hypertelorism, broad nasal root, micrognathia, clinodactyly, brachydactyly and syndactyly.