# Dysgenezja gonad typu XY i anomalie towarzyszące Kod Orpha: 1770 Kod OMIM: 233430

## Opis choroby *

**Definicja**
A rare syndrome with 46,XY disorder of sex development characterized by mild developmental delay and streak gonads associated with short stature, cardiac, renal, musculoskeletal, and ectodermal abnormalities (the latter including scalp defects and unusual hair whorls), and dysmorphic facial features (such as preauricular pits, short columella, and small nares). There have been no further descriptions in the literature since 1980.
**Dane**

**Klasyfikacja**
Zespół wad wrodzonych

**Kod ORPHA**
1770

**Kod OMIM**
233430

**Kod ICD10**
Q99.1

**Kod ICD11**
-

*Źródło

orphanet

## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.