## Opis choroby *

Definicja

A multiple congenital anomalies syndrome characterized by wormian bones, dextrocardia and short stature due to a growth hormone deficiency. Additional manifestations that have been reported include brachycamptodactyly, kidney hypoplasia, bilateral cryptorchidism, midshaft hypospadias, imperforate anus/anorectal agenesis, body asymmetry, mild developmental delay, hemimegalencephaly and facial dysmorphism (hypotelorism, downslanting palpebral fissures, low-set and posteriorly angulated ears, depressed nasal bridge, and microstomia).

Dane

**Klasyfikacja**
Zespół wad wrodzonych

Synonimy
Stratton-Parker syndrome
Zespół Strattona i Parkera

**Kod ORPHA**
2863

**Kod OMIM**
185120

**Kod ICD10**
Q87.1

**Kod ICD11**
-

---

*Źródło

orphanet