

# Zespół McDonough

Kod Orpha: 2471 Kod OMIM: 248950

## Opis choroby \*

### Definicja

McDonough syndrome is a rare, multiple congenital anomalies/dysmorphic syndrome characterized by facial dysmorphsim (prominent superciliary arcs, synophrys, strabismus, large, anteverted ears, large nose, malocclusion of teeth), delayed psychomotor development, intellectual disability and congenital heart defects (e.g. pulmonic stenosis, patent ductus arteriosus, atrial septal defect). Additional features include thorax deformation (pectus excavatum/carinatum), kyphoscoliosis, diastasis recti and cryptorchidism. There have been no further descriptions in the literature since 1984.

### Dane

#### Klasyfikacja

Zespół wad  
wrodzonych

**Kod ORPHA**  
2471

**Kod OMIM**  
248950

**Kod ICD10**  
Q87.8

**Kod ICD11**  
LD2F.1Y

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[\\* Źródło](#)

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## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

Dostępna na stronie [www.orphanet.pl](http://www.orphanet.pl)