

Opis choroby *

Definicja

A rare otorhinolaryngologic disease characterized by total or partial anosmia at birth. The anosmia is caused by a defect in the development of the olfactory bulbs or by replacement of the olfactory epithelium by respiratory epithelium. Isolated congenital anosmia is found in some parents of individuals with Kallman syndrome.

Dane

Klasyfikacja

Choroba

Kod ORPHA

88620

Kod OMIM

107200

Kod ICD10

Q07.8

Kod ICD11

CA0Y

*Źródło

orphanet