Opis choroby *

Definicja

A rare, genetic male infertility due to a sperm disorder characterized by the absence of a measurable amount of spermatozoa in the ejaculate (azoospermia), or a number of sperm in the ejaculate inferior to 15 million/mL (oligozoospermia), resulting from a mutation in a single gene known to cause azoo- or oligo-spermia. Sperm morphology may be normal.

Dane

Klasyfikacja
Choroba

Kod ORPHA
399805

Kod OMIM
618086

Kod ICD10
N46

Kod ICD11
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*Źródło

orphanet