

Opis choroby *

Definicja

Maternal uniparental disomy of chromosome 6 is an uniparental disomy of maternal origin characterized by intrauterine growth retardation. Homozygosity for a recessive disease mutation for which only a mother is a carrier may lead to other phenotypes.

Dane

Klasyfikacja	Synonimy
Zespół wad wrodzonych UPD(6)mat	UPD(6)mat

Kod ORPHA	Kod OMIM	Kod ICD10
96181	-	Q99.8

Kod ICD11
LD45.0

*Źródło

orphanet