

## Opis choroby \*

### Definicja

A rare frontonasal dysplasia characterized by a craniofacial phenotype comprising frontal bossing with high anterior hairline, ptosis, hypertelorism, epicanthus inversus, flat nasal bridge, and broad nasal tip. Large anterior fontanelle, sagittal synostosis, and cranial base anomalies have also been described.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych SIX2-related FND	SIX2-related FND
	SIX2-related FND

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
488437	-	Q75.8

**Kod ICD11**  
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### \*Źródło

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