

## Opis choroby \*

### Definicja

A rare, severe, circulatory system disease characterized by premature, diffuse, severe atherosclerosis (including the aorta and renal, coronary, and cerebral arteries), sensorineural deafness, diabetes mellitus, progressive neurological deterioration with cerebellar symptoms and photomyoclonic seizures, and progressive nephropathy. Partial deficiency of mitochondrial complexes III and IV in the kidney and fibroblasts (but not in muscle) may be associated. There have been no further descriptions in the literature since 1994.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Atherosclerosis-hearing loss-diabetes-epilepsy-nephropathy syndrome
	Zespół Feigenbauma, Bergerona i Richardsona
	Feigenbaum-Bergeron-Richardson syndrome

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
1192	209010	I70.9

**Kod ICD11**  
LD2H.Y

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### \*Źródło

orphanet