

## Opis choroby \*

### Definicja

A rare congenital anomalies syndrome characterized by a variable spectrum of ulnar defects, mammary and apocrine gland hypoplasia and genital anomalies. The most frequent signs include fifth finger and dental anomalies, delayed puberty and mammary hypoplasia. Short stature and obesity are common.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Pallister ulnar-mammary syndrome
	UMS
	Zespół łokciowo-sutkowy Pallistera
	Zespół Schinzela
	Schinzal syndrome
	UMS

**Kod ORPHA**  
3138

**Kod OMIM**  
181450

**Kod ICD10**  
Q71.8

**Kod ICD11**  
LD2F.1Y

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### \*Źródło

orphanet