## Opis choroby \*

Definicja

Piebaldism is a rare congenital pigmentation skin disorder characterized by the presence of hypopigmented and depigmented skin areas (leukoderma) on various parts of the body, preferentially on the forehead, chest, abdomen, upper arms, and lower extremities, that are associated with a white forelock (poliosis), and in some cases with hypopigmented and depigmented eyebrows and eyelashes.

Dane

Klasyfikacja

Choroba

Kod ORPHA 2884

**Kod OMIM** 172800

Kod ICD10 E70.3

Kod ICD11 EC23.2Y

\*Źródło

orphanet