

## Opis choroby \*

### Definicja

A rare form of familial hypoaldosteronism characterized by adult onset of subnormal plasma aldosterone with elevated plasma renin activity, hyperkalemia, metabolic acidosis, and hypotension. Signs and symptoms are typically mild, and affected individuals may be clinically asymptomatic and diagnosed only after biochemical screening.

### Dane

#### Klasyfikacja

Podtyp kliniczny

#### Synonimy

Late-onset familial hyperreninemic  
hypoaldosteronism  
Łagodny niedobór syntazy aldosteronu  
Mild aldosterone synthase deficiency

#### Kod ORPHA

556037

#### Kod OMIM

-

#### Kod ICD10

E27.4

#### Kod ICD11

-

---

#### \*Źródło

orphanet