

## Opis choroby \*

### Definicja

A form of pseudohypoaldosteronism type 1 characterized by mild mineralocorticoid resistance that is restricted to the kidneys and that usually improves in early childhood. Typical presentation is in the neonatal period with weight loss, failure to thrive, vomiting and dehydration in association with hyponatremia, hyperkalemia and metabolic acidosis as well as elevated aldosterone and renin levels.

### Dane

#### Klasyfikacja

Podtyp kliniczny

#### Synonimy

Autosomal dominant PHA1

Autosomalny dominujący

pseudohypoaldosteronizm typu 1

Autosomal dominant pseudohypoaldosteronism  
type 1

Renal PHA1

#### Kod ORPHA

171871

#### Kod OMIM

177735

#### Kod ICD10

N25.8

#### Kod ICD11

GB90.41

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#### \*Źródło

orphanet