

## Opis choroby \*

### Definicja

A rare, primary form of mineralocorticoid resistance characterized by mild to profound salt wasting either restricted to the kidney (renal pseudohypoaldosteronism type 1), or generalized affecting many organs (generalized pseudohypoaldosteronism type 1). Clinical presentation is in the neonatal period with failure to thrive, vomiting and dehydration with biochemical findings of hyperkalaemia, metabolic acidosis and, elevated plasma aldosterone and renin concentration.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

PHA type 1

PHA type 1

PHA1

#### Kod ORPHA

756

#### Kod OMIM

264350

#### Kod ICD10

N25.8

#### Kod ICD11

GB90.41

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#### \*Źródło

orphanet