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
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Kod ICD10
E27.4

Kod ICD11
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*Źródło

orphanet