

Opis choroby *

Definicja

A rare heritable form of primary aldosteronism (PA) that is characterized by early-onset severe hypertension, non- glucocorticoid-remediable hyperaldosteronism, overproduction of 18-oxocortisol and 18-hydroxycortisol, and profound hypokalemia.

Dane

Klasyfikacja

Choroba

Synonimy

FH-III

FH3

FH-III

Rodzinny hiperaldosteronizm typu 3

FH3

Familial hyperaldosteronism type 3

Kod ORPHA

251274

Kod OMIM

613677

Kod ICD10

E26.0

Kod ICD11

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*Źródło

orphanet