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