Opis choroby *

Definicja

Familial glucocorticoid deficiency (FGD) is a group of primary adrenal insufficiencies characterized clinically by neonatal hyperpigmentation, hypoglycemia, failure to thrive, and recurrent infections, and biochemically by glucocorticoid deficiency without mineralocorticoid deficiency.

Dane

Klasyfikacja
Choroba

Kod ORPHA
361

Kod OMIM
617825

Kod ICD10
E27.1

Kod ICD11
5A74.Y

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*Źródło

orphanet