

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

*Źródło

orphanet