Familial glucocorticoid deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial glucocorticoid deficiency. ORPHA:361

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.