

Disease

Congenital adrenal hyperplasia due to 21-hydroxylase deficiency^{1, 2}

Gene

CYP21A2

Inheritance	AR
Current Treatments	Hydrocortisone, 9α-fludrohydrocortisone
Evidence for Treatment	guideline ³
Non-Molecular Testing Availability	yes
Non-Molecular Test	17-hydroxyprogesterone level ¹
Disease Frequency (per 100,000)	6.67 ⁴
Genes Included in Disease Frequency	CYP21A2
Experimental / Investigational Therapies	https://clinicaltrials.gov/ct2/results?cond=Congenital+Adrenal+Hyperplasia&term=&cntry=&state=&city=&

References

- ¹ <https://www.ncbi.nlm.nih.gov/books/NBK1171/>
- ² <https://pubmed.ncbi.nlm.nih.gov/32966723/>
- ³ <https://pubmed.ncbi.nlm.nih.gov/30272171/>
- ⁴ <https://ghr.nlm.nih.gov/condition/21-hydroxylase-deficiency#genes>