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Multiplier Hack 1.0 Server, Multiplier Hack 1.1. Congenital adrenal hyperplasia (CAH) is a group of autosomal recessive disorders caused by an inherited enzymatic defect in the adrenal cortex. Untreated patients may have developmental defects, including both sexual and skeletal defects. CAH is classified into four types according to the enzyme defect in the zona fasciculata, zona glomerulosa, or zona reticularis. The proper diagnosis of CAH is based on high levels of 17-hydroxyprogesterone, 17-hydroxypregnenolone, 17-hydroxycorticosterone, or androsterone, and increased levels of the 11-deoxycortisol or 21-deoxycortisol derivative in a 24-hour urine sample. Untreated patients with CAH may also have other clinical abnormalities including hypotension, hypoglycemia, and hyperpigmentation. The increased androgen levels may result in irregular growth of genitalia in male patients. Individuals with CAH also may have an increased risk of developing certain types of cancer, particularly in the adrenal glands. The CAH gene is located at 2p22.3. Sixteen mutations (A-P) in the CYP21A2 gene of CAH have been identified to date (Margeirsdottir I et al., 1999). N-ethylmaleimide-sensitive fusion protein attachment protein receptor type 1 (NAP-1) and NAP-1-associated protein 1 (NAP-1P) both have high homology to NAP-1. The NAP-1 and NAP-1P proteins were identified in response to glutathione-S-transferase-NAP-1 fusion protein. They are receptors for the Ca²⁺-dependent signaling molecule annexin II, calreticulin
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