

Effective Date: October 20, 2025

TEST CHANGE

Adrenocorticotropic Hormone

0070010, ACTH

Specimen Requirements:	
Patient Preparation:	Morning collection (7 a.m. to 10 a.m.) is preferred.
Collect:	Lavender (<u>K2-EDTA</u>), <u>K2EDTA</u>) or Pink (<u>K2-EDTA</u>), or <u>K3-EDTA</u> . Collection tube must be siliconized glass or plastic.
Specimen Preparation:	Separate from cells ASAP or within 2 hours of collection. Transfer 1 mL plasma to an ARUP Standard Transport Tube and freeze immediately. (Min: 0.5 mL)
Transport Temperature:	CRITICAL FROZEN. Separate specimens must be submitted when multiple tests are ordered.
Unacceptable Conditions:	Serum, heparinized plasma, tissue or urine. Grossly hemolyzed specimens.
Remarks:	
Stability:	After separation from cells: Ambient: 3 hours; Refrigerated: 4 hours; Frozen: 10 weeks (No freeze/thaw cycles.)
Methodology:	Quantitative Electrochemiluminescent Immunoassay (ECLIA)
Performed:	Sun-Sat
Reported:	Within 24 hours
Note:	No reference intervals established for p.m. collections.
CPT Codes:	82024

CPT Codes: 8202

New York DOH Approval Status: This test is New York DOH approved.

Interpretive Data:

Reference interval based on samples collected between 7 a.m. and 10 a.m. No reference intervals established for p.m. collections. Pediatric reference values are the same as adults (Acta Paediatr Scand 1981;70:341-345). This assay measures intact ACTH 1-39; some types of synthetic ACTH and ACTH fragments are not detected by this assay.

Reference Interval:

Effective August 5, 2019

7.2<u>?-?</u>--63.3 pg/mL (a.m. draws)



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