

## **TEST CHANGE**

Human Immunodeficiency Virus Types 1 and 2 (HIV-1/2) Antibody Differentiation, Supplemental		
2013107, HIV AB SUP		
Specimen Requirements:		
Patient Preparation:		
Collect:	Lavender (EDTA) or pink (K2EDTA). Also acceptable: Serum separator tube (SST).	
Specimen Preparation:	Separate from cells ASAP or within 2 hours of collection. Transfer 1 mL serum or plasma into an ARUP standard transport tube dedicated only for HIV testing. (Min: 0.5 mL) Remove particulate material.	
Transport Temperature:	Frozen.	
Unacceptable Conditions:	Specimens containing particulate material. Severely hemolyzed or heat-inactivated specimens.	
Remarks:		
Stability:	After separation from cells: Ambient: 48 hours; Refrigerated: 7 days; Frozen: <u>8 months Indefinitely</u> (avoid repeated freeze/thaw cycles)	
Methodology:	Qualitative Immunoassay	
Performed:	Varies	
Reported:	1-2 days	
Note:	For use ONLY when patient has a repeatedly reactive third- or fourth-generation HIV screen test result. This test discriminates between HIV-1 and HIV-2 antibodies. Results for each type are reported. This test is for use as the antibody differentiation test in the specific multitest algorithm. If results are negative or indeterminate, this test does NOT reflex to a nucleic acid test. A multitest algorithm is recommended by the Centers for Disease Control and Prevention (CDC) and the Clinical Laboratory Standards Institute (CLSI) for the diagnosis of HIV (refer to http://arupconsult.com/content/human- immunodeficiency-virus).	
CPT Codes:	86701; 86702	
New York DOH Approval Status:	This test is New York DOH approved.	



Interpretive Data:

This test should not be used for blood donor screening, associated reentry protocols, or for screening human cells, tissues, and cellular- and tissue-based products (HCT/P).

Reference Interval:

Test Number		Reference Interval
	HIV-1 Antibody	Negative
	HIV-2 Antibody	Negative