

HIV Rapid Testing MPEP December 2005
Panel and Vial Designations, CDC Donor Bulk Numbers,
CDC HIV Rapid Test Results and Donor HIV Status

Panel Letter	Vial Label	CDC Donor Bulk Number	CDC Test Result ^{1,3}	Donor HIV Status	Laboratory Interpretation ² and/or Results	
					Test Result	Interpretation
A	A1	18	Positive (S)	Infected	_____	_____
	A2	14	Positive (S)	Infected	_____	_____
	A3	5	Negative	Uninfected	_____	_____
	A4	10	Positive (W)	Infected	_____	_____
	A5	16	Positive (W)	Infected	_____	_____
	A6	19	Positive (W)	Infected	_____	_____
B	B1	19	Positive (W)	Infected	_____	_____
	B2	10	Positive (W)	Infected	_____	_____
	B3	14	Positive (S)	Infected	_____	_____
	B4	18	Positive (S)	Infected	_____	_____
	B5	5	Negative	Uninfected	_____	_____
	B6	16	Positive (W)	Infected	_____	_____
C	C1	5	Negative	Uninfected	_____	_____
	C2	18	Positive (S)	Infected	_____	_____
	C3	16	Positive (W)	Infected	_____	_____
	C4	14	Positive (S)	Infected	_____	_____
	C5	19	Positive (W)	Infected	_____	_____
	C6	10	Positive (W)	Infected	_____	_____
D	D1	19	Positive (W)	Infected	_____	_____
	D2	14	Positive (S)	Infected	_____	_____
	D3	10	Positive (W)	Infected	_____	_____
	D4	16	Positive (W)	Infected	_____	_____
	D5	18	Positive (S)	Infected	_____	_____
	D6	5	Negative	Uninfected	_____	_____

¹ The CDC result was obtained after pre-shipment testing for the presence of HIV-1 Antibody with all commercially available HIV Rapid Testing kits licensed by the Food and Drug Administration (FDA) and with selected FDA-licensed Enzyme Immunoassay (EIA) kits. The CDC result is consistent with the manufacturers' criteria for interpretation of results.

² Laboratory Interpretation space (to be completed by participant laboratory) provided to facilitate comparison of participant laboratory result with CDC result.

³ Strong (S) and Weak (W) designations are based on qualitative observations of the colorimetric test results for reactive samples.