

HIV Rapid Testing MPEP September 2009
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	1	Positive (W)	Infected	_____	_____
	A2	2	Positive (W)	Infected	_____	_____
	A3	3	Positive (S)	Infected	_____	_____
	A4	1*	Positive (W)	Infected	_____	_____
	A5	4	Positive (S)	Infected	_____	_____
	A6	5	Negative (N)	Uninfected	_____	_____
B	B1	2	Positive (W)	Infected	_____	_____
	B2	3	Positive (S)	Infected	_____	_____
	B3	1	Positive (W)	Infected	_____	_____
	B4	4	Positive (S)	Infected	_____	_____
	B5	5	Negative (N)	Uninfected	_____	_____
	B6	1*	Positive (W)	Infected	_____	_____
C	C1	3	Positive (S)	Infected	_____	_____
	C2	1	Positive (W)	Infected	_____	_____
	C3	4	Positive (S)	Infected	_____	_____
	C4	5	Negative (N)	Uninfected	_____	_____
	C5	1*	Positive (W)	Infected	_____	_____
	C6	2	Positive (W)	Infected	_____	_____
D	D1	1	Positive (W)	Infected	_____	_____
	D2	4	Positive (S)	Infected	_____	_____
	D3	5	Negative (N)	Uninfected	_____	_____
	D4	1*	Positive (W)	Infected	_____	_____
	D5	2	Positive (W)	Infected	_____	_____
	D6	3	Positive (S)	Infected	_____	_____

* Duplicate donors

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.