Centers for Disease Control and Prevention Model Performance Evaluation Program

Human T-Lymphotropic Virus Types I and II (HTLV-I/II) Testing

Figures Used to Summarize the Participant Laboratory Results for the May 24, 1999 Shipment

U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention
Public Health Practice Program Office
Division of Laboratory Systems
Atlanta, Georgia 30341-3717





Use of trade names is for identification only and does not constitute endorsement by the U.S. Department of Health and Human Services.

Report of the May 24, 1999 Human T-lymphotropic Virus Types I and II (HTLV-I/II) Performance Evaluation Sample Testing Results Provided by Participant Laboratories in the Model Performance Evaluation Program, Centers for Disease Control and Prevention (CDC).

The production of this report was coordinated in CDC by:

Public Health Practice Program Office	Edward L. Baker, M.D., M.P.H., Director
Division of Laboratory Systems	Robert Martin, Ph.D., Director
Laboratory Practice Assessment Branch	Thomas L. Hearn, Ph.D., Acting Chief

The material in this report was developed and prepared by:

Information about this report should be addressed to the Model Performance Evaluation Program by calling (770) 488-8091 or (770) 488-8098.

Centers for Disease Control and Prevention (CDC) Model Performance Evaluation Program Human T-Lymphotropic Virus Types I and II (HTLV-I/II) Testing May 24, 1999 Participant Laboratory Shipment

Table 1

Panel Letter	Vial Label	CDC Donor Number	CDC Contractor Result ²	Laboratory Interpretation ¹			
				INIT. ³ F	INAL⁴	WB	IIF
			_				
Α	A1	4	Negative				
	A2,A6	1	Negative				
	A3,A5	2	Positive, HTLV-II				
	A4	3	Positive, HTLV-I	_			
В	B1,B3	1	Negative				
	B2	4	Negative				
	B4,B6	2	Positive, HTLV-II				
	B5	3	Positive, HTLV-I				

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

The CDC result was obtained after composite EIA, WB, and RIPA testing, by CDC contractor, and employing the interpretation criteria of the Public Health Service Working Group.

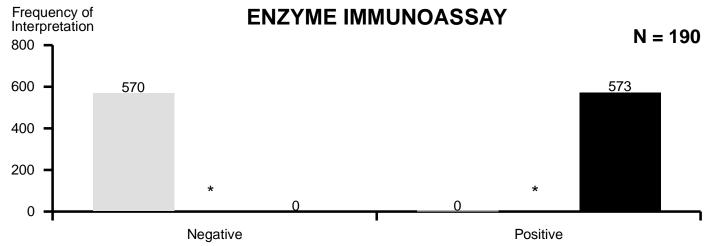
Initial EIA interpretation

Final EIA interpretation

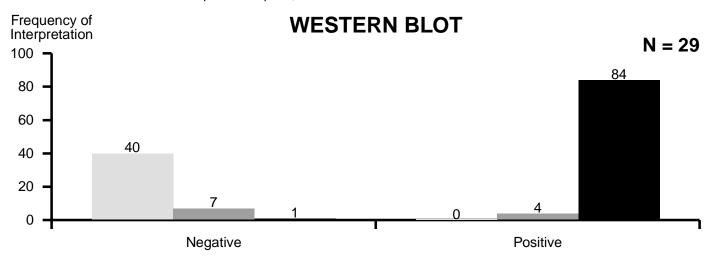
SUPPLEMENTAL INFORMATION FOR COMPREHENDING THE NUMBERS USED TO LABEL FIGURES IN THIS REPORT

The "N=" that appears on each graph represents the number of laboratories that reported results. For some graphs, laboratories reported results using more than one test; therefore, the number of results may exceed the actual number of laboratories providing reports. In figures 1-7 and 10, the vertical axis is labeled either as frequency or percentage of results; in figures 8 and 9, this axis is labeled as percentage of reports. However, in all figures, the number appearing directly above or within each bar represents a frequency of results only.

Figure 1. Frequency of HTLV-I/II antibody test result interpretations, by sample type (reactivity), for enzyme immunoassay (EIA), Western blot (WB), and indirect immunofluorescence (IIF), reported by participant laboratories for the May 24, 1999 shipment



* = Indeterminate is not an EIA interpretation option; these areas have been left blank



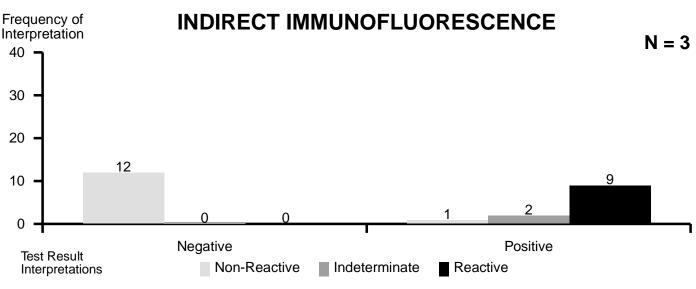


Figure 2. Percentage of HTLV-I/II participant laboratories, by laboratory type, that reported EIA, WB, and IIF results to the CDC for the May 24, 1999 shipment

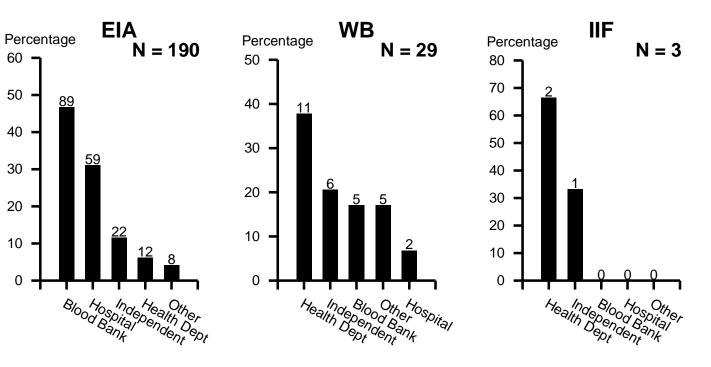


Figure 3. Combination of HTLV-I/II antibody tests reported by participant laboratories for the May 24, 1999 shipment

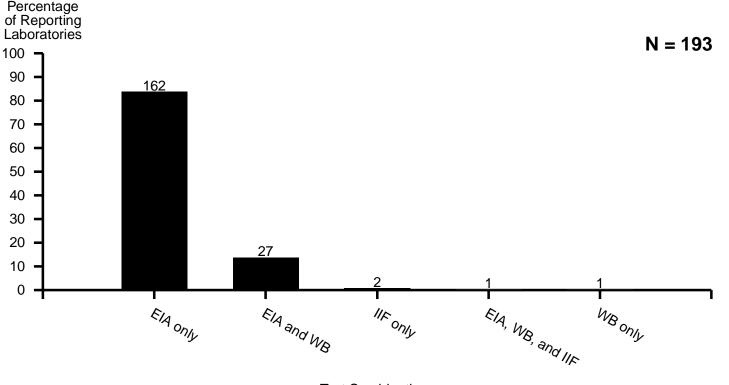
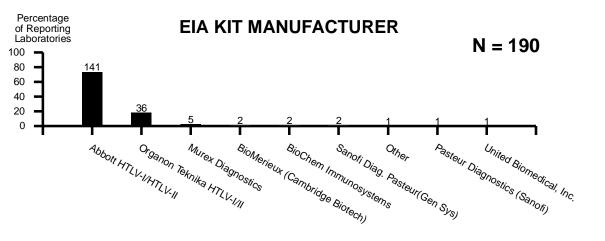
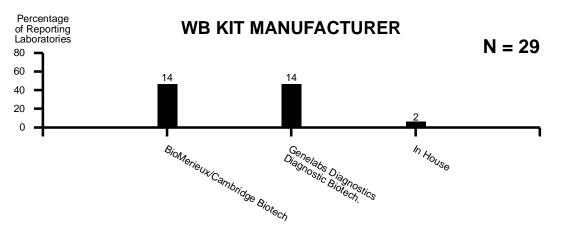


Figure 4. Types of HTLV-I/II antibody test kits used for enzyme immunoassay, Western blot, and indirect immunofluorescence, as reported by participant laboratories to the CDC for the May 24, 1999 shipment





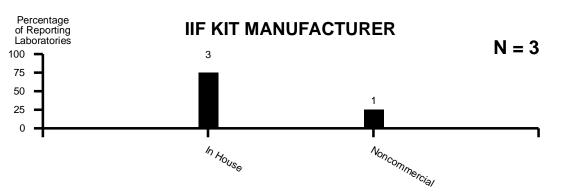
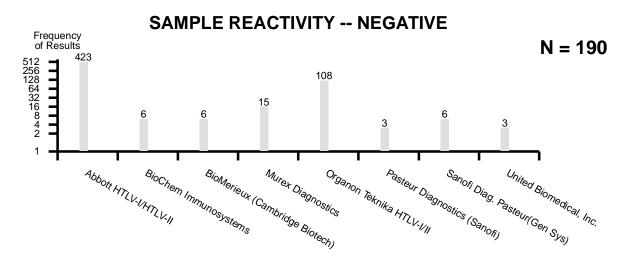
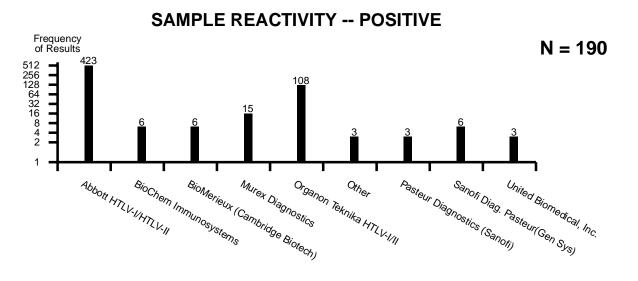


Figure 5. Enzyme immunoassay HTLV-I/II antibody test results, by kit manufacturer, reported by participant laboratories for the May 24, 1999 shipment

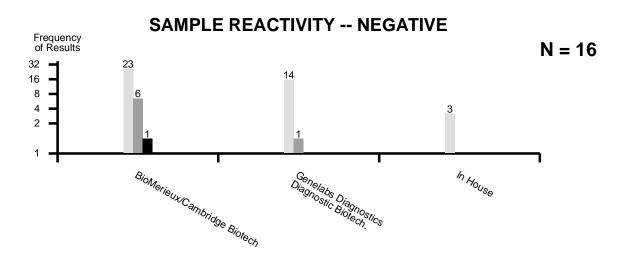


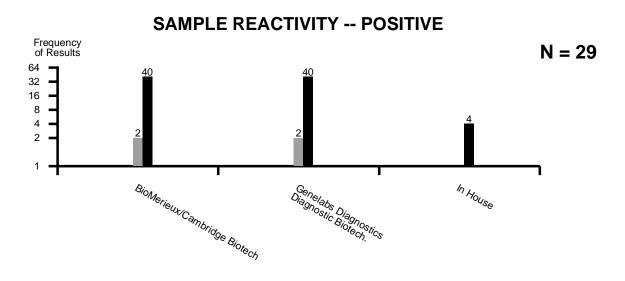


Test Result Interpretations

Non-Reactive Reactive

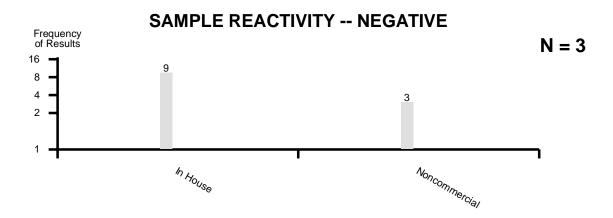
Figure 6. Western blot HTLV-I/II antibody test results, by kit manufacturer, reported by participant laboratories for the May 24, 1999 shipment

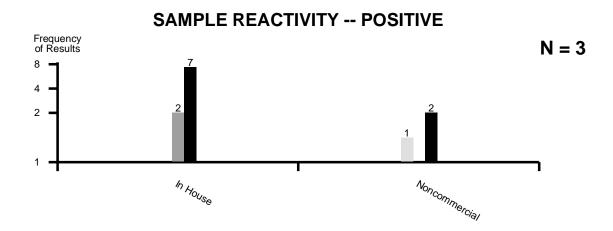




Test Result

Figure 7. Indirect immunofluorescence HTLV-I/II antibody test results, by kit manufacturer, reported by participant laboratories for the May 24, 1999 shipment





Test Result

Figure 8. Western blot HTLV-I/II antibody band patterns reported to CDC by participant laboratories for the May 24, 1999 shipment

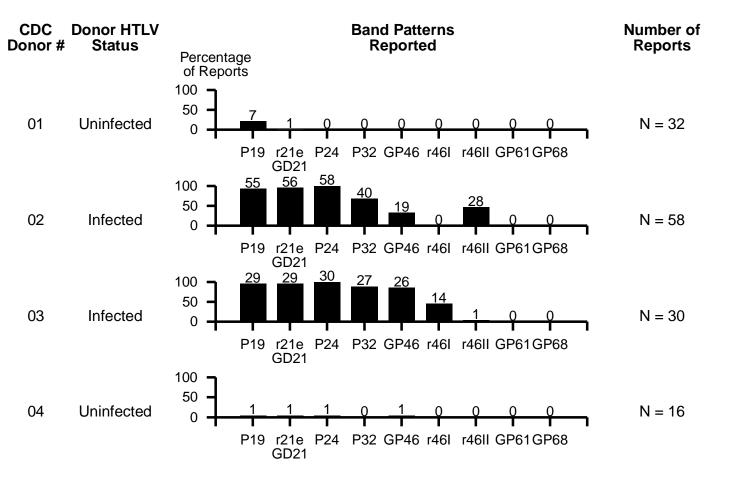


Figure 9. Fluorescence intensity patterns, of HTLV-I/II-infected cells, for IIF results reported to CDC by participant laboratories for the May 24, 1999 shipment

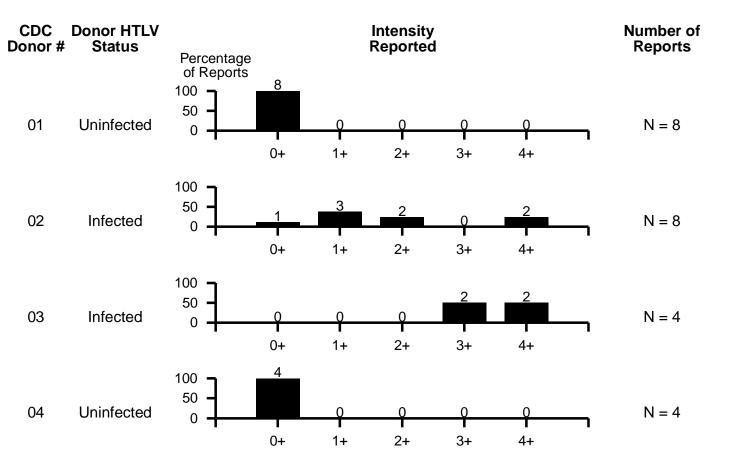
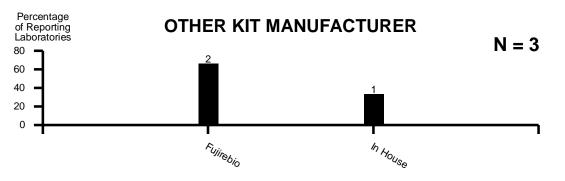
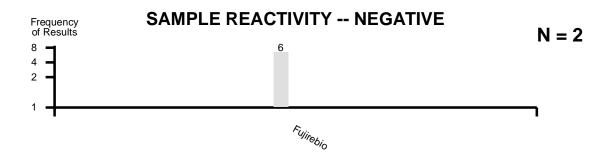
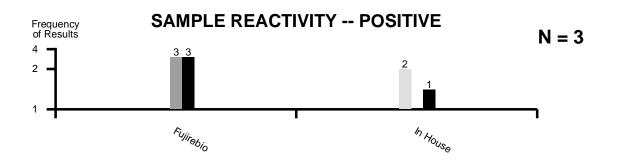


Figure 10. Types of 'Other' HTLV antibody test kits used and results reported by participant laboratories to the CDC for the May 24, 1999 shipment







Test Result Interpretations

■ Non-Reactive ■ Indeterminate ■ Reactive