■ Test Procedure



Read the results within next 10 minutes

■ Interpretation of Results

Note: Results should not be read beyond 10 minutes







Invalid



One dot at C

Two dots at T & C

Dot only at T

No Dot

Ordering Information

Product	Code No.	Pack size
Signal	51FT100-10	10 Tests
	51FT100-50	50 Tests
	51FT100-60	100 Tests



THE MANUFACTURING SITE'S QMS IS CERTIFIED FOR ISO 13485:2003, ISO 9001:2008

Comprises of former IVD business undertaking of Span Diagnostics Ltd.)
Plot No. 336, 338, 340, Road No. 3, G.I.D.C., Sachin - 394 230, (Surat), INDIA.
Telephone: 0261-239 7712 to 15, Facsimile: 0261-239 7719, Website: www.arkray.co.in

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Flow through

immunodot test format for detection of

Antibodies to HIV I+2 in serum or plasma

■ Fact File

100% sensitivity and 100% specificity of test results with cocktail of highly purified, recombinant antigens comprising of immunodominant region of envelope antigens gp41 & gp36 for both HIV 1 & 2.

Detects all subtypes of HIV-1 including Group M - subtypes A, B, C and Group O.

Antibodies developed are of high avidity enhancing delectability in **early seroconversion*** stage, as early as **9 days** after the first bleed.

*Seroconversion sensitivity demonstrates the ability to detect antibodies at the lowest level during early infection.



^{*}As per evaluation at NARI using universally accepted BBI USA Seroconversion Panel Panel member represents sample taken from one person over a period of time. It represents evolution of disease maker from non-reactive to reactive.

Salient Features



Visual two dot assay with inbuilt control dot to validate successful completion of assay procedure gives results in 10 minutes.



Unique cassette design with handle for maximum user safety and convenience along with self dropping bottles ensuring maximum ease of use.



The optimised mixture of recombinant antigens of the most conserved immunodominant region of both HIV-1 & HIV-2, immobilised on membrane, enables detection of low titer & low affinity antibodies in its early seroconversion stage.



Optimally designed test protocol with proprietory washing buffer ensures highest specificity by removing all non specific interfering proteins.



Proprietory conjugate containing uniform sized gold particles ensures prominent appearance of dots for unambigious interpretation of test results.

■ Evaluations & Approval*



Approved for use in Blood Banks by Directorate General of Health Service (DGHS).

NARI

Sensitivity: 100% Specificity: 100%

* National AIDS Research Institute (NARI), Pune.



Sensitivity: 100% Specificity: 100%

* Y. R. Gaitonde Center for AIDS Research and Education (YRG CARE), Chennai.

*Data on Arkray file