abcam

Product datasheet

HIV1 p24 ELISA Kit ab218268

SimpleStep ELISA

18 References 画像数 15

製品の概要					
製品名	HIV1 p24 ELISA Kit				
検出方法	Colorimetric				
再現性					Intra-Assay(同時再現性)
	サンプル	Ν	平均值	SD	CV%
	Overall	3			4.5%
					Inter-Assay(日差再現性)
	サンプル	Ν	平均值	SD	CV%
	Overall	5			4.7%
サンプルの種類	Cell culture supernatan Plasma	t, Serum, Cel	l culture extracts, Tis	sue Extracts, Hep	Plasma, EDTA
アッセイタイプ	Sandwich (quantitative)			
検出感度	1.1 pg/ml				
検出範囲	4.69 pg/ml - 300 pg/ml				
添加回収試験					特定サンプルでの回収試験
	サンプルの種類		平均%	測定範	西田
	Serum		105	94% -	117%
	Cell culture extracts		107	103%	- 113%
	Cell culture media		102	94% -	107%

102

98

1h 30m

Hep Plasma

EDTA Plasma

100% - 103%

93% - 103%

ステップ 種交差性 製品の概要

One step assay

交差種: Human

HIV1 p24 ELISA Kit (ab218268) is a single-wash 90 min sandwich ELISA designed for the quantitative measurement of HIV1 p24 protein in human serum, plasma, cell culture supernatant, and cell and tissue extract samples. It uses our proprietary SimpleStep ELISA® technology. Quantitate HIV1 p24 with 1.1 pg/mL sensitivity.

Lentivirus vectors based on HIV-1 have gained popularity as a tool for cell and gene therapy, owing to their ability to carry and integrate a high volume of transgenes in a range of dividing and non-dividing cell types. Pseudo-lentiviral particles are typically produced from 293T cells and harvested from the medium 48-78hrs post-transfection. To ensure that the pseudoviral medium is viable, and to control the number of copies of integrated viral constructs per target cell, the viral titre needs to be determined before transduction experiments. Our HIV1 p24 SimpleStep ELISA[®] can be used to rapidly and sensitively determine the titre of HIV1 particles in range of sample types.

SimpleStep ELISA® technology employs capture antibodies conjugated to an affinity tag that is recognized by the monoclonal antibody used to coat our SimpleStep ELISA® plates. This approach to sandwich ELISA allows the formation of the antibody-analyte sandwich complex in a single step, significantly reducing assay time. See the SimpleStep ELISA® protocol summary in the image section for further details. Our SimpleStep ELISA® technology provides several benefits:

- -Single-wash protocol reduces assay time to 90 minutes or less
- -High sensitivity, specificity and reproducibility from superior antibodies
- -Fully validated in biological samples
- -96-wells plate breakable into 12 x 8 wells strips

A 384-well SimpleStep ELISA® microplate (<u>ab203359</u>) is available to use as an alternative to the 96-well microplate provided with SimpeStep ELISA® kits.

ASSAY SPECIFICITY This kit recognizes both native and recombinant HIV-1 p24 protein in serum, EDTA and heparin plasma, cell culture supernatant, and cell and tissue extract samples only. Urine, saliva, and milk samples have not been tested with this kit.

CROSS REACTIVITY Recombinant HIV-1 Gag protein was prepared at 50 ng/mL and 1 ng/mL and assayed for cross reactivity. 1% cross-reactivity was observed.

特記事項 HIV1 p24 (capsid) protein is essential for HIV-1 viral replication and for the HIV-1 infection of nondividing cells. HIV1 p24 proteins form viral capsid that encapsulates the genomic HIV1 RNA. HIV1 p24 concentration in host plasma is commonly used as indicator of viral load. Upon the viral infection, the development of anti-HIV1 p24 host humoral responses leads to immune complex formation and reduction of the free HIV1 p24 concentration in circulation.

> Abcam has not and does not intend to apply for the REACH Authorisation of customers' uses of products that contain European Authorisation list (Annex XIV) substances. It is the responsibility of our customers to check the necessity of application of REACH Authorisation, and any other relevant authorisations, for their intended uses.

試験プラットフォーム

Pre-coated microplate (12 x 8 well strips)

製品の特性

保存方法

Store at +4°C. Please refer to protocols.

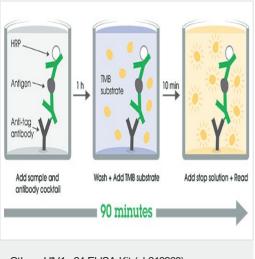
内容	1 x 96 tests
10X HIV1 p24 Capture Antibody	1 x 600µl
10X HIV1 p24 Detector Antibody	1 x 600µl
10X Wash Buffer PT (ab206977)	1 x 20ml
50X Cell Extraction Enhancer Solution (ab193971)	1 x 1ml
5X Cell Extraction Buffer PTR (ab193970)	1 x 10ml
Antibody Diluent 5BI	1 x 6ml
HIV1 p24 Lyophilized Recombinant Protein	2 vials
Plate Seals	1 unit
Sample Diluent 50BS	1 x 20ml
Sample Diluent NS (ab193972)	1 x 50ml
SimpleStep Pre-Coated 96-Well Microplate (ab206978)	1 unit
Stop Solution	1 x 12ml
TMB Development Solution	1 x 12ml

関連性

HIV1 performs highly complex orchestrated tasks during the assembly, budding, maturation and infection stages of the viral replication cycle. During viral assembly, the proteins form membrane associations and self-associations that ultimately result in budding of an immature virion from the infected cell. Gag precursors also function during viral assembly to selectively bind and package two plus strands of genomic RNA. Capsid protein p24 probably forms the conical core of the virus that encapsulates the genomic RNA-nucleocapsid complex.

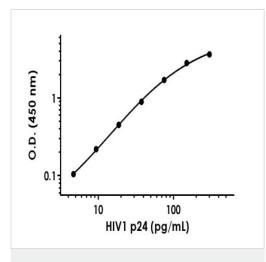
細胞内局在

Membrane



SimpleStep ELISA technology allows the formation of the antibodyantigen complex in one single step, reducing assay time to 90 minutes. Add samples or standards and antibody mix to wells all at once, incubate, wash, and add your final substrate. See protocol for a detailed step-by-step guide.





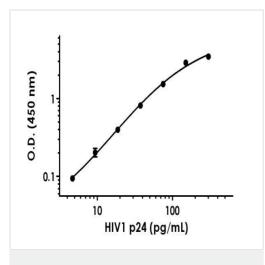
Example of HIV1 p24 standard curve in Sample Diluent 50BS

Background-subtracted data values (mean +/- SD) are graphed.

Standard Curve Measurements					
Concentration	O.D 4	Mean			
(pg/mL)	1	2	O.D		
0	0.1495	0.1478	0.149		
4.69	0.257	0.249	0.253		
9.38	0.370	0.364	0.367		
18.75	0.593	0.605	0.599		
37.5	1.040	1.039	1.039		
75	1.863	1.839	1.851		
150	2.901	3.031	2.966		
300	3.762	3.801	3.782		

Raw data values are shown in the table

Raw data values for Example of HIV-1 p24 standard curve in Sample Diluent 50BS.



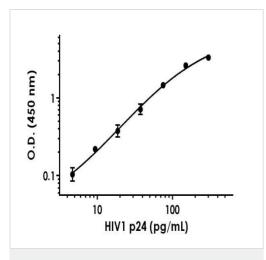
Example of HIV1 p24 standard curve in Sample Diluent NS

Background-subtracted data values (mean +/- SD) are graphed.

Standard Curve Measurements					
Concentration	O.D 4	Mean			
(pg/mL)	1	2	O.D		
0	0.082	0.081	0.081		
4.69	0.174	0.178	0.176		
9.38	0.304	0.267	0.285		
18.75	0.460	0.498	0.479		
37.5	0.892	0.898	0.895		
75	1.633	1.598	1.616		
150	3.017	2.917	2.967		
300	3.581	3.499	3.540		

Raw data values are shown in the table

Raw data values for example of HIV-1 p24 standard curve in Sample Diluent NS.

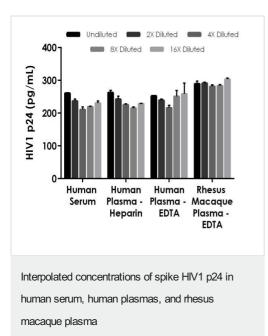


Example of HIV1 p24 standard curve in 1X Cell Extraction Buffer PTR Background-subtracted data values (mean +/- SD) are graphed.

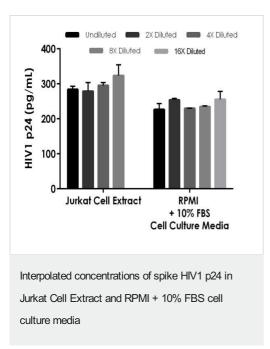
Standard Curve Measurements					
Concentration	O.D 4	Mean			
(pg/mL)	1	2	O.D		
0	0.066	0.065	0.065		
4.69	0.185	0.160	0.172		
9.38	0.283	0.290	0.286		
18.75	0.491	0.399	0.445		
37.5	0.860	0.701	0.780		
75	1.604	1.445	1.525		
150	2.824	2.528	2.676		
300	3.407	3.328	3.367		

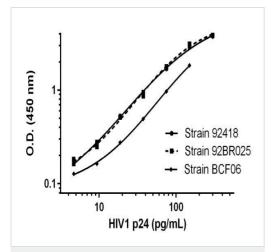
Raw data values are shown in the table

Example of HIV-1 p24 standard curve in 1X Cell Extraction Buffer PTR



The concentrations of HIV1 p24 were measured in duplicates, interpolated from the HIV1 p24 standard curves and corrected for sample dilution. Undiluted samples are as follows: human serum 100% (neat), human plasma (heparin) 100% (neat), human plasma (EDTA) 100% (neat), and rhesus macaque plasma (EDTA) 100% (neat). The interpolated dilution factor corrected values are plotted (mean +/- SD, n=2).





Serial dilutions of recombinant HIV1 p24 were prepared within the working range of the assay and assayed for reactivity. The concentrations of HV1 p24 were measured in duplicates, interpolated from the HV1 p24 standard curves and corrected for sample dilution. Undiluted samples are as follows: Jurkat cell extract 100 µg/mL, RPMI + 10% FBS cell culture media 100% (neat). The interpolated dilution factor corrected values are plotted (mean +/- SD, n=2).

Serial dilutions of recombinant HIV1 p24 (group M, subtype B, strain 92418), HIV1 p24 (group M, subtype C, strain 92BR025) and HIV1 p24 (group O, strain BCF06) were prepared within the working range of the assay and assayed for reactivity. O.D. values within the linear range of each protein are graphed.

Sample Type	Average % Recovery	Range (%)	
100% Human Serum	133	115 - 159	
100% Human Plasma Heparin	122	117 - 128	
100% Human Plasma EDTA	113	103 - 125	

Three concentrations of purified HIV-1 p24 protein were spiked in duplicate to the indicated biological matrix and treated with the Acid Treatment Protocol to evaluate signal recovery in the working range of the assay. The signals of the same concentrations of purified HIV-1 p24 protein spiked in duplicate to Sample Diluent 50BS and treated with the Acid Treatment Protocol were taken as 100%.

Acid treatment recovery

Sample Type	Average % Effect	Range (%)
100% Human Serum (donor 1)	63	62-63
100% Human Serum (donor 2)	65	63-66
100% Human Serum (donor 3)	70	69-71
100% Human Serum (donor 4)	60	60-60
100% Human Serum (pooled)	59	59-59
100% Human Plasma Heparin (pooled)	70	68-72
100% Human Plasma EDTA (pooled)	69	68-70

Single concentrations of purified HIV-1 p24 protein were spiked in duplicate to the indicated biological matrix and treated with the Acid Treatment Protocol or mock acid treated to evaluate the effect of the Acid Treatment Protocol on the signal. The signals of single concentrations of purified HIV-1 p24 protein spiked in duplicate to the indicated biological matrix and mock acid treated were taken as 100%.

Acid treatment effect

Sample Type	Average % Recovery	Range (%)
100% Human Serum	16	14-17
100% Human Plasma - Heparin	14	14-15
100% Human Plasma - EDTA	10	10-10
100% Rhesus Macaque Plasma EDTA	16	14-19

Acid treatment recovery of synthetic complexes

Single concentrations of purified HIV-1 p24 protein were spiked to the indicated biological matrix, pre-incubated with unlabeled capture and detector antibodies or unrelated antibodies and treated with the Acid Treatment Protocol to evaluate the signal recovery of p24 complexed with interfering antibodies by Acid Treatment Protocol. The signals of single concentrations of purified HIV-1 p24 protein spiked to the indicated biological matrix, preincubated with unrelated antibodies and treated with the Acid Treatment Protocol were taken as 100%. There were no signals if the single concentrations of purified HIV-1 p24 protein were spiked to the indicated biological matrix, pre-incubated with the unlabeled capture and detector antibodies and mock acid treated, indicating that the unlabeled capture and detector antibodies efficiently formed synthetic immune complexes with HIV-1 p24 and completely blocked the signals.

Dilution Factor	Interpolated value	100% Human Serum	100% Human Plasma (Heparin)	100% Human Plasma (EDTA)	100% Rhesus Macaque Plasma (EDTA)
المعائل بالمعط	pg/mL	261	263	252	289
Undiluted	% Expected value	100	100	100	100
2	pg/mL	118	121	120	146
	% Expected value	91	92	95	101
4	pg/mL	52.7	56.3	54	70.4
	% Expected value	81	96	86	97
8	pg/mL	27.4	26.9	31.4	35.4
	% Expected value	84	82	99	98
16	pg/mL	14.4	14.3	16.2	19.0
	% Expected value	88	87	103	105

Purified HIV-1 p24 was spiked into biological samples and diluted in a 2-fold dilution series in Sample Diluent 50BS.

Linearity of dilution – spiked Purified HIV-1 p24 in Human serum, plasma (heparin, citrate, EDTA) and rhesus macaque EDTA plasma.

Dilution Factor	Interpolated value	100% RPMI + 10% FBS Cell Culture Media	100 µg/mL <u>Jurkat</u> Cell Extract
lladilutad	pg/mL	227	284
Undiluted	% Expected value	100	100
2	pg/mL	127	139
	% Expected value	112	98
4	pg/mL	57.4	73.8
	% Expected value	101	104
8	pg/mL	29.3	40.4
	% Expected value	104	114
16	pg/mL	16.0	NL
	% Expected value	113	NL

Linearity of dilution – spiked purified HIV-1 p24 in FBS cell culture media and Jurkat Cell Extract Purified HIV-1 p24 was spiked into biological samples and diluted in a 2-fold dilution series in Sample Diluent NS for cell culture media and 1X Cell Extraction Buffer PTR for cell extract samples.

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