



Nucleic Acid Extraction Kit

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(Magnetic Bead Method)



Molecular

Nucleic Acid Extraction Kit (Magnetic Bead Method)

High quality DNA / RNA applied to PCR, DNA Cloning, NGS and etc.

Our reagents can be used for nucleic acid isolation of multiple of sample. They are pre-filled and ready-to-use which can be easily load into analyzer to render security and ease of our end-user.



Easy operation, rapid extraction

Only one step washing

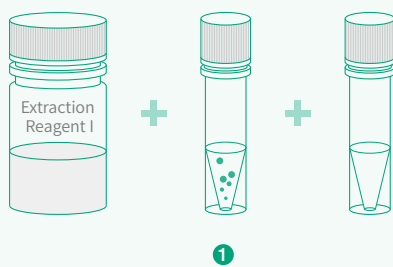
9 min for 32 samples

Extract once and get DNA and RNA meanwhile, meeting your needs for multiple index detection.

Manual operation(A-200)

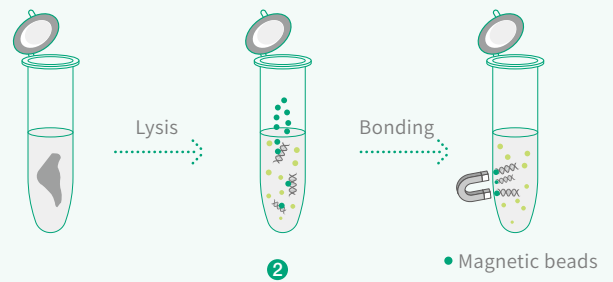
• Pretreatment

500 μ L(Extraction Reagent I) + 4 μ L(Magnetic Beads Solution) + 15 μ L(Proteinase K), mix into [Working Solution].



• Lysate

500 μ L [Working Solution] + 200 μ L sample, mix well, lyse at 55 $^{\circ}$ C for 4 min, absorbed by magnetic separator for 1 min, and discard the supernatant.



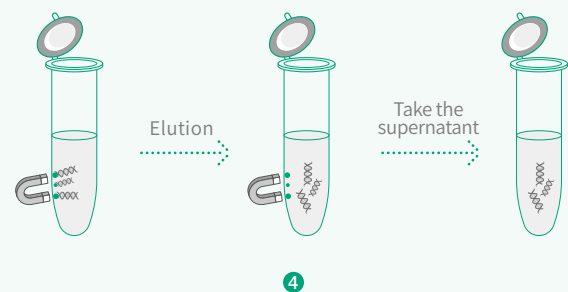
• Rinsing

add 600 μ L [Isolation Reagent II], mix well, absorbed by magnetic separator for 1min, and discard the supernatant.



• Elution

add 50~100 μ L [Elution Buffer], elute at 80 $^{\circ}$ C for 2 min, absorbed by magnetic separator for 30s, reserve the supernatant.



Efficient isolation, reliable performance

High repeatability

Good linear correlation

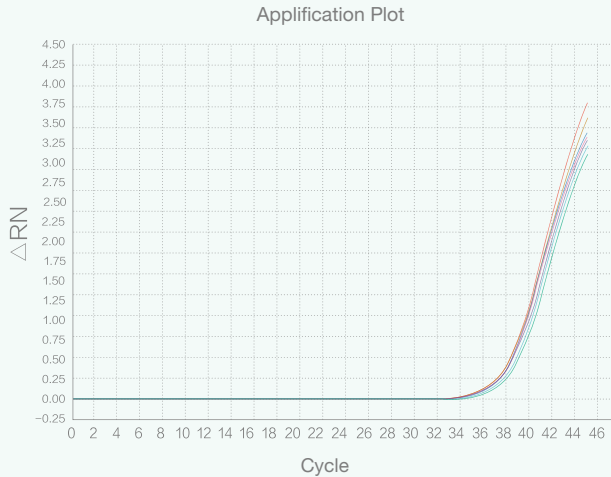


Figure 3-1 Amplification Curve of HBV Reference Material (10 IU/mL)

200 μL 10 IU/mL diluted HBV reference material from WHO (NIBSC code: 10/264) was isolated by the kit to get 50 μL analyte. The analyte was detected by HBV diagnosis kit 10 times. Positive rate is 100%, as shown in Figure 3-1

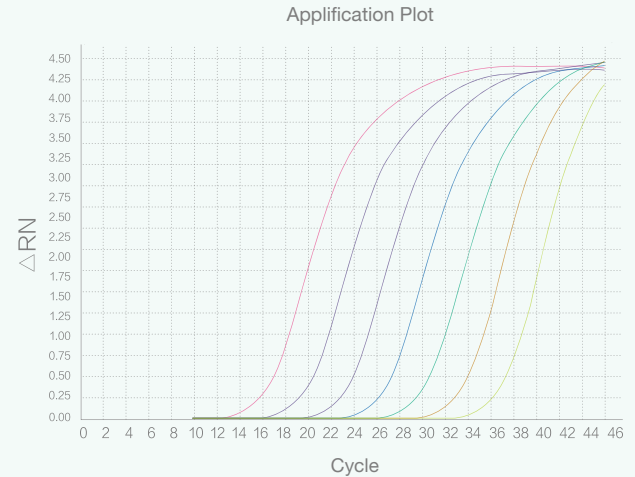


Figure 3-2 Amplification Curve of HCV Reference Material (25 IU/mL)

200 μL 25 IU/mL diluted HCV reference material (5th WHO International Standard for HCV NAT, NIBSC code: 14/150) was isolated by the kit to get 50 μL analyte. The analyte was detected by HBV diagnosis kit 10 times. Positive rate is 100%, as shown in Figure 3-2

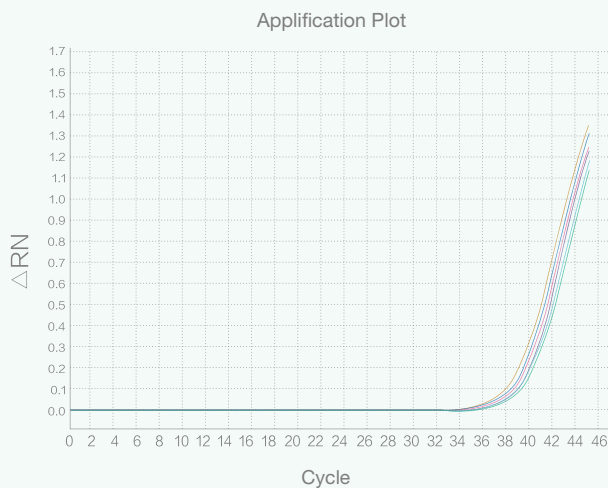


Figure 3-3 Amplification Curve of DNA Pseudoviridae

The DNA Pseudoviridae with a concentration of 5×10^8 IU/mL was diluted with negative serum to 5×10^7 IU/mL, 5×10^6 IU/mL, 5×10^5 IU/mL, 5×10^4 IU/mL, 5×10^3 IU/mL, 5×10^2 IU/mL and 50 IU/mL. They were determined after isolation. The results were shown in Figure 3-3

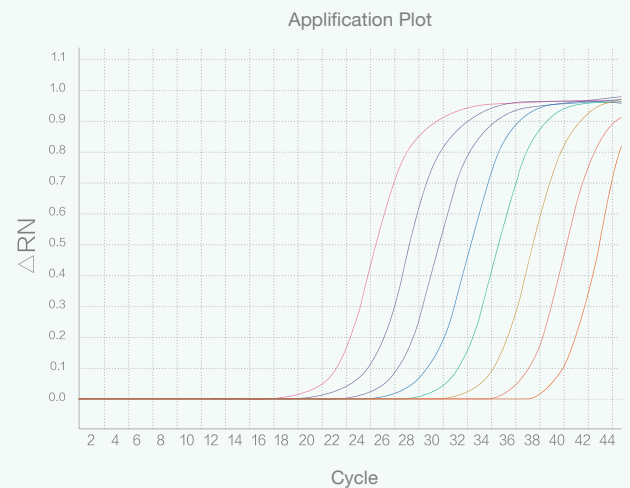
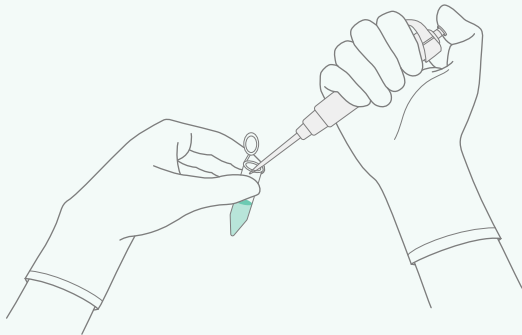


Figure 3-4 Amplification Curve of RNA Pseudoviridae

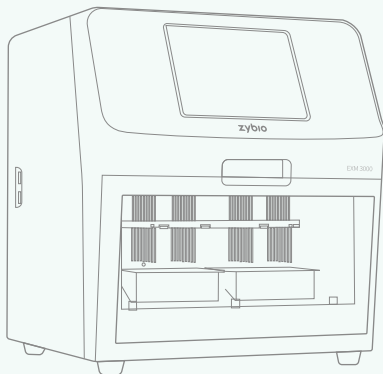
The RNA Pseudoviridae with a concentration of 5×10^7 IU/mL was diluted with negative serum to 5×10^6 IU/mL, 5×10^5 IU/mL, 5×10^4 IU/mL, 5×10^3 IU/mL, 5×10^2 IU/mL. They were determined after isolation. The results were shown in Figure 3-4

Flexible extraction method



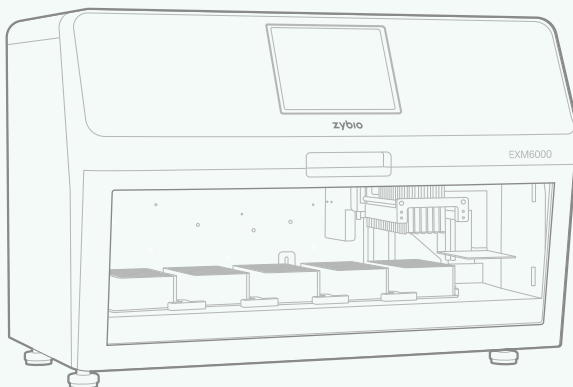
Manual Operation

Up to 16 samples for per test;
Extraction time : 10-15 min;
Only one step washing;



Nucleic Acid Isolation System EXM3000

Up to 32 samples for per test;
Extraction time : 9 min;
Prefilled and ready-to-use;



Nucleic Acid Isolation System EXM6000

Up to 96 samples for per test;
Extraction time : 12 min;
Low cross-contamination risk;

Performance parameter

Sample Types: Liquid samples such as serum, plasma, nasopharyngeal swab, cell preservation solution, tissue fluid, urine and secretions.

Extraction Time: 9 - 12 min

Recovery: $\geq 90\%$

Repeatability: $CV \leq 2\%$

Subsequent Use: qPCR, hybridization

Specifications

Reagent Kit	Application	Sample size	Model	Packing Specifications
Viral Nucleic Acid Kit	pathogen infection, pathogen resistance	200 μ L	A-200	32 T/Kit 96 T/Kit
		200 μ L	B-200	8 T/Kit 16 T/Kit 32 T/Kit
		200 μ L	T-200	32 T/Kit 96 T/Kit



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