



Viga Quantitative HBV Molecular Diagnostic Kit

Store at -15°C to -30°C in darkness

100 rxn

Cat NO: MD003055

25 rxn

Cat NO: MD003056

By ROJE

Edition, 02/2022

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Kit Content

Viga Quantitative HBV Molecular Diagnostic Kit

Components	25 Preps	100
		Preps
Pro HBV Mix	220 µl	875µl
QR-ROMAX, 4X	160 μΙ	625µl
IC	125 μΙ	500µl
HBV *QS1(1×10 ⁴ IU/μl)	65µl	250μΙ
HBV *QS2(1×10 ³ IU/μl)	65μΙ	250μΙ
HBV *QS3(1×10 ² IU/μl)	65μΙ	250μΙ
HBV *QS4(1×10 ¹ IU/μl)	65μΙ	250μΙ
Water (Molecular Biology grade)	1500μΙ	1500μΙ

Recommended Starting Material

DNA Sample Requirement

Before starting, add 2-5 ml blood sample into a tube containing EDTA. After plasma isolation and DNA extraction, utilize 10µl of whole prepared sample in Real Time-PCR.

Before start

Before first use, make sure about intactness and completeness of kit contents and regents Avoiding utilizing samples other than human plasma to prevent incorrect IVD examination results.

Misusing the regents may lead to contamination and invalid results.

Use RNAase/DNAase free tip sampler with filter.

Buffer preparation

Master mix preparation

The total volume of prepared sample used in this test is $10 \mu l$. Refer table2. Required information for preparing tubes is available in tables 3,4,5. If you use internal control, refer to the following information in the handbook.

Notice: prepare Master mix just for single-use and avoid adding QR-

ROMAX to Pro HBV mix if you do not need to test.

Table1: Reagents preparation per one single reaction (DNA isolation efficiency and PCR inhibition is controlled by adding internal control in purification stage).

Required component	volume
Pro HBV Mix	8.75µl
QR-ROMAX, 4X	6.25µl
Purified DNA	10µl

Table 2: Required volumes for standard tubes

Standards	Volume per tube	Pro HBV Mix + QR- ROMAX, 4X per reaction
QS1	10µl	15µl
QS2	10µl	15µl
QS3	10µl	15µl
QS4	10µl	15µl



Table 3: Required volumes for every single test tubes

Volume per	Pro HBV Mix	
tube of an	+ QR-	
unknown	ROMAX, 4X	
sample	per reaction	
10μΙ	15µl	

Table 4: Required volumes for negative control tubes

Volume per tube of water*	Pro HBV Mix + QR- ROMAX, 4X per reaction	
10μΙ	15µl	

Table 5: Thermal profile for Real Time PCR

stage	Temperature	Incubation Time	Cycle Numbers
Pre- Denatura tion	95 °C	5 min	1
Denatura tion Annealin g and Extensio n	95 °C 58°C	10 sec 60 sec	5
Denatura tion Annealin g and acquisitio n on channel Green and Yellow	95 °C 58°C	10 sec 60 sec	40

Protocol

Take out each component from the kit and place them on a benchtop. Allow the reagents to equilibrate to room temperature, then briefly vortex each tube for later use. Before use, vortex briefly. Utilize 10 µl of whole prepared sample in Real-Time PCR.

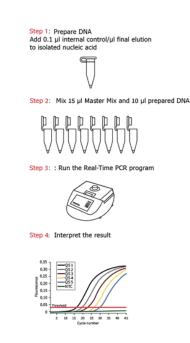


Figure1: Addition of internal control during DNA amplification in PCR. In this case, Master mix involves Pro HBV Mix and QR-ROMAX,4×.

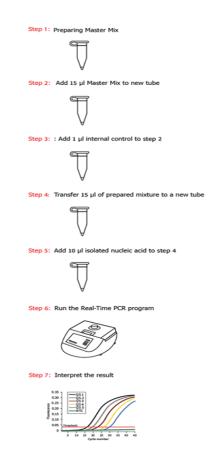


Figure2: Addition of internal control to Master mix. Notice that there is not any addition of internal control on the purification stage. In this case, Master mix involves Pro HBV Mix and QR-ROMAX,4×. For more information, refer Master mix preparation.