

CYTOMEGALOVIRUS DNA CONTROL

MBC016: Purified DNA of Cytomegalovirus.

LOT SPECIFICATIONS:

Microorganism: Cytomegalovirus.

Preparation: Grown in MRC-5 infected cells.

COMPONENTS:

Lyophilized genomic DNA of Cytomegalovirus (1×10^4 - 1.5×10^4 copies/ μ l once reconstituted). DNA quantification has been performed with real-time PCR from Stratagene (ref. Mx3005P).

TECHNICAL DATA:

Presentation: Lyophilized.

Instructions for reconstitution:

- 1.-Centrifuge the lyophilized genomic DNA for 1 min at 1000 g.
- 2.-Add 100 μ l of sterile bidistilled water and mix until completely resuspended. The concentration will be 1×10^4 - 1.5×10^4 copies/ μ l once reconstituted.
- 3.-Shake with vortex for 30 seconds to dissolve and homogenize completely.

Extract preparation: SDS and proteinase K treatment, phenol/chloroform extraction and ethanol precipitation.

Stability and storage:

Special shipping conditions are not required as product is lyophilized. Reconstitute upon receipt. Store between -5°C and -40°C after reconstitution.

The DNA solution should be aliquoted in order to avoid repeated freezing and thawing.

By following the instructions for use, the product is stable until the expiry date stated on the labels.

Recommendations and precautions:

This product is not for diagnostic use.

This product is for research use only.

This lot of DNA was prepared from Cytomegalovirus. The material should be handled and disposed of as potentially infectious. Observe the local regulations for clinical waste disposal.

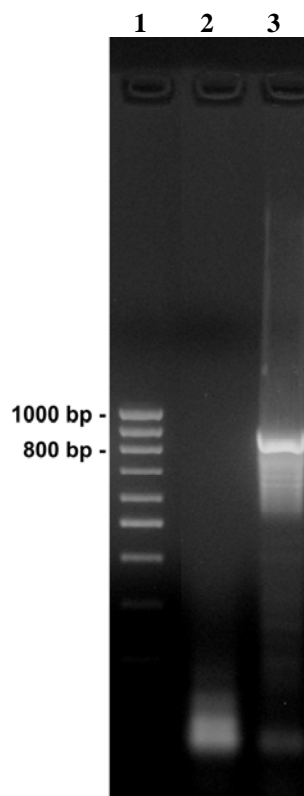
Nucleic acids should not be repeatedly frozen and thawed. It is recommended to reconstitute and aliquote the control upon reception.

FOR INFORMATION USE ONLY

Not to be used for performing the assay. Refer to the insert accompanying the kit.

The dilutions should be done just before use. Frozen or dilutions containing less than 1000 copies/ μ l is not recommended as copy numbers can be lost.

PCR analysis of DNA control: PCR analysis was performed with a specific oligo pair on purified Cytomegalovirus DNA. The reaction produced a 860 bp fragment. It was visualized on a 2% agarose gel using ethidium bromide staining. The gel photograph is shown below:



Line	Sample
1	Molecular size marker (100 bp)
2	Negative control
3	860 bp amplified fragment

For inquiries please contact:
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FOR RESEARCH USE ONLY

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