

CHLAMYDOPHILA PNEUMONIAE DNA CONTROL

MBC011: Purified DNA of *Chlamydomphila pneumoniae*.

LOT SPECIFICATIONS:

Microorganism: *Chlamydomphila pneumoniae*.

Preparation: Purified from Hep-2 infected cells by sonication and differential centrifugation.

COMPONENTS:

Lyophilized genomic DNA of *Chlamydomphila pneumoniae* (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 minute at 1000 g.
- 2.-Add 100 μ l of sterile bidistilled water and mix until completely reconstituted. 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.

Shipping and storage:

Special transport conditions not required. 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 *Chlamydomphila pneumoniae*. The material should be handled and disposed of as potentially infectious. Observe the local regulations for clinical waste disposal.

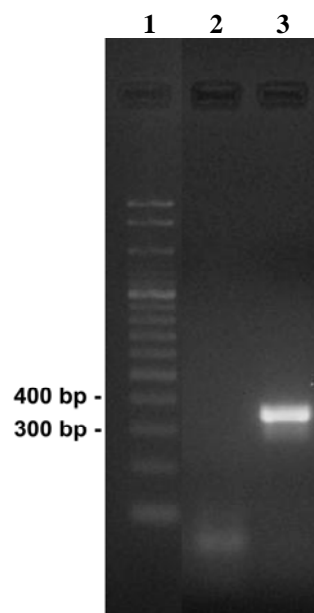
Nucleic acids should not be repeatedly frozed and thawed. It is recommended to reconstitute and aliquote the control upon reception.

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

FOR INFORMATION USE ONLY

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

PCR analysis of DNA control: PCR analysis was performed with a specific oligo pair on purified *Chlamydomphila pneumoniae* DNA. The reaction produced a 368 bp fragment. It was visualized on a 2% agarose gel using ethidium bromide staining. The gel photograph is shown below:



Line	Simple
1	Molecular size marker (100 bp)
2	Negative control
3	368 bp amplified fragment

For inquiries please contact:
customerservice@vircell.com

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FOR RESEARCH USE ONLY

Manufacturer: VIRCELL, S.L. Pza. Dominguez Ortiz 1. Polígono Industrial Dos de Octubre.18320 Santa Fe *GRANADA* SPAIN* Tel.+34.958.441264* Fax+34.958.510712
http://www.vircell.com