

INSTRUCTIONS FOR USE

Basement Membrane Positive Control (neat)

C009N/C009N-0.5

1ml/0.5ml

INTENDED USE

The Bio-Diagnostics Basement membrane positive control is intended for use in indirect immunofluorescent antibody (IFA) tests to detect autoantibodies in human serum. This positive control material has been manufactured so that it is ready for use.

SUMMARY AND EXPLANATION

This control has been found to be strongly IgG positive on monkey Oesophagus slides by giving brilliant yellow-green fluorescence (between 3+ and 4+) when used neat.

PRINCIPLE OF THE TEST

The positive control and human sera are incubated on monkey oesophagus slides. The primary test reaction involves circulating antibodies present in the patient's serum, which attach to their homologous antigens. A rinsing period is followed by a secondary reaction using a FITC anti-human globulin conjugate. The antigen surface is then thoroughly rinsed free of unbound conjugate and viewed under an appropriate fluorescent microscope to visually identify various morphological patterns of nuclear fluorescence. With a positive reaction, a bright line of fluorescence is observed on the basement membrane of the oesophagus tissue.

WARNINGS AND PRECAUTIONS

1. This positive control has been tested and found to be negative or non-reactive for STS, HBsAg, HBV by PCR (NAT), HIV 1/2 antibody, HIV by PCR (NAT), HCV antibody and HCV PCR (NAT). However, these tests cannot guarantee the absence of infectious agents. All human components should be handled with appropriate care.
2. The control contains 0.01% thiomersal as a preservative. Although this is at a low concentration, these reagents should be considered toxic. They should not be ingested or allowed to come into contact with either the skin or the mucous membranes.
3. Do not use control beyond its expiry date.
4. For *in vitro* diagnostic use.

MATERIALS PROVIDED / STORAGE & STABILITY

CONTROL + Basement membrane positive control: C009N / C009N-0.5

The control must be stored at 2-8°C upon receipt. Check label for expiry date.

ADDITIONAL MATERIALS REQUIRED BUT NOT PROVIDED

Substrate section slides as appropriate

FITC conjugate as appropriate

Mounting Medium (Bio-Diagnostics R005).

Phosphate Buffered Saline (PBS) (Bio-Diagnostics R002).

Test tubes and rack or microtitre system Disposable pipettes Coverslips

Staining Dish and Slide Forceps Moisture Chamber Distilled Water

Volumetric Flask (500 ml) Fluorescence Microscope Paper Towels – lint free

All reagents required are available from Bio-Diagnostics Ltd - see catalogue for details.

KEY FOR OTHER SYMBOLS

Used in this instruction leaflet and on accompanying product packaging:

 Manufacture **RFU** Ready for use

 Temperature limitation **IVD** *In vitro* diagnostic medical device

TEST INSTRUCTIONS

This control is supplied ready for use for testing on Bio-Diagnostics slides. Follow the recommended testing method for the particular slide type.

QUALITY CONTROL

Positive and negative serum controls must be included in each day's testing to confirm reproducibility, sensitivity and specificity of the test procedure.

TEST LIMITATIONS

No diagnosis should be based upon a single serologic test result since various host factors must be taken into consideration.

EC REP

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