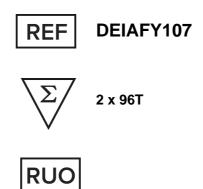




Influenza A Virus Nucleoprotein Antigen ELISA Kit



This product is for research use only and is not intended for diagnostic use.

For illustrative purposes only. To perform the assay the instructions for use provided with the kit have to be used.

Creative Diagnostics

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Cat: DEIAFY107

PRODUCT INFORMATION

Intended Use

Influenza A Virus Nucleoprotein Antigen ELISA Kit is a qualitative procedure for the detection of Influenza A nucleoprotein antigen in complex sample matrices derived from both human and veterinary sources.

General Description

Influenza viruses can be divided into three classes, A, B, and C, largely based upon conserved antigenic differences in the internal nucleoprotein. Influenza A virus, typically encountered more frequently than types B and C, and associated with the majority of serious epidemics, can be further subdivided into strains or subtypes based on antigenic differences in the external hemagglutinin proteins (H1-H16) and neuraminidase proteins (N1-N9).

A variety of wild waterfowl appear to be the predominant natural reservoir for Influenza A viruses and subtypes representing most of the hemagglutinin and neuraminidase combinations can be found circulating in these birds. Historically, human influenza virus infections have been associated with H1N1, H2N2, and H3N2 subtypes of influenza A, although a recent (1997) and significant outbreak in Hong Kong was identified as an H5N1 subtype. This outbreak was not only significant because it resulted in 18 human infections and 6 deaths, but it also represented the first known demonstration of avian influenza virus transmission to humans.

While influenza A subtype identification is extremely important (vaccine production, epidemiology), the rapid and accurate differentiation of influenza A from influenza B and C and other respiratory agents in humans and animals is also important (treatment and biosecurity).

The Influenza A nucleoprotein antigen ELISA kit is a capture ELISA for the qualitative measurement of influenza A virus nucleoprotein antigen and can be completed in less than 1.5 h and contains only one wash step. In addition, the test kit incorporates proprietary diluents that are designed to prevent the development of nonspecific signal derived from complex sample matrix effects and/or the nonspecific adsorption of reactive test components which result in improvements in both sensitivity and specificity.

Reagents And Materials Provided

- 1. IAV Antigen Capture Plate (96 tests)-2 ea.
- 2. Sample Preparation Reagent (1x)-12 ml
- 3. IAV Positive Control (1x)-1 ml
- 4. IAV Negative Control (1x)- 2 x 1.5 ml
- 5. Wash Buffer (20x)-30 ml
- 6. IAV Detection Antibody, HRP-labeled (1x)-22 ml
- 7. Chromagen Solution (1x)-22 ml
- 8. Stop Solution (1x)-22 ml
- 9. Sample Dilution Tray-2 ea.

Storage

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Store all kit components at 2-8°C. Crystal formation may occur in the wash buffer concentrate during prolonged storage at 2-8°C.The crystals can be re-dissolved by swirling the bottle in warm tap water.

Assay Procedure

- 1. Remove the kit components from storage and allow to warm to room temperature.
- 2. Determine the number of test wells needed. Use one well for each sample. In addition, include one well for the Positive Control and three wells for the Negative Control.
- 3. To begin the assay, transfer 50 μl of Sample Preparation Reagent to the appropriate number of wells in the dilution tray provided.
- 4. Add 200 µl of each sample, positive control, and negative control to the IAV Sample Preparation Reagent. Mix by pipetting up and down several times.
- 5. Transfer 100 µl of sample or control to the appropriate wells of the IAV Antigen Capture Plate.
- 6. Cover the plate and incubate for 30 min at room temperature on a plate shaker set at moderate speed.
- 7. Add 100 µl of 1x IAV Detection Antibody to each well, cover the plate and incubate for 45 min on a plate shaker using the same settings (Step 6).
- 8. Wash the wells 6x with at least 300 µl/well 1x Wash Buffer.
- Add 100 µl of Chromagen to each well and incubate for 10 min on a plate shaker.
- 10. Stop the reaction by the addition of 100 μ l of Stop Solution.
- 11. Shake the plate for 10-15 sec to ensure that the reaction is uniformly stopped and then read the plate in a plate reader using a 450 nm filter.

Quality Control

- 1. All negative control absorbance values should be ≤ 0.250 .
- 2. The positive control absorbance value should be ≥ 0.500 .
- 3. The calculated value for the positive control/cut-off should be ≥ 2 .
- 4. To determine the cut-off value, calculate the mean of the three negative control absorbance values and multiply this value by 2.

Interpretation Of Results

To interpret the results for a given sample, divide the absorbance value for the sample by the cut-off value. Calculated sample values that are >1.1 are considered reactive. Calculated sample values that are <0.9 are considered nonreactive. Calculated sample values that are ≥ 0.9 and ≤ 1.1 are considered equivocal.

Specificity

N/A

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