

Avian Leukemia Virus Antigen Rapid Test

Catalog No.: JIA405

INTENDED USE

The Avian Leukemia Virus Antigen Rapid Test is a lateral flow immunochromatographic assay for the qualitative detection of Avian Leukemia virus antigen (ALV Ag) from avian serum, plasma, the tissues of lesion or secretions from cloaca.

Assay Time: 5-10 minutes

Specimen: serum, plasma, the tissues of lesion or secretions

PRINCIPLE

The Avian Leukemia Virus Antigen Rapid Test is based on sandwich lateral flow immunochromatographic assay. The test device has a testing window for the observation of assay running and result reading. The testing window has an invisible T (test) zone and a C (control) zone before running the assay. When the treated sample was applied into the sample hole on the device, the liquid will laterally flow through the surface of the test strip and react with the pre-coated monoclonal antibodies. If there is ALV antigen in the specimen, a visible T line will appear. The C line should always appear after a sample is applied, which indicates a valid result. By this means, the device can accurately indicate the presence of ALV antigen in the specimen.

REAGENTS AND MATERIALS

- 20 Test devices, with disposable droppers
- 20 vials of Assay buffer
- 20 Swabs
- 1 Products Manual

STORAGE AND STABILITY

The kit can be stored at room temperature (4-30°C). The test kit is stable through the expiration date (18 months) marked on the package label. **DO NOT FREEZE.** Do not store the test kit in direct sunlight.

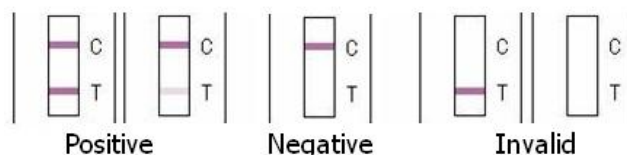
TEST PROCEDURE

- Collect avian serum or plasma into a centrifuge tube. Make sure the specimen is clean and transparent. Transfer 3 drops (around 100 μ L) of the specimen into the assay buffer tube and mix them well. Use the mixture in the assay.
- If the specimen is the tissues of lesion, collect a part of the tissue into the assay buffer. Agitate and crush the tissue and make the tissue mixed as much as possible in the assay buffer. Use the supernatant mixture in the assay.
- If the specimen is secretions from cloaca, collect the bird's cloacal secretions or feces with the cotton swab and insert the swab into the provided assay buffer tube.

Agitate it to get efficient sample extraction.

- Allow all materials, including specimen and test device, recover to 15-25°C before running the assay.
- Take out the test device from the foil pouch and place it horizontally.
- Suck the treated sample extraction from the assay buffer tube and place 3 drops into the sample hole "S" of the test device.
- Interpret the result in 5-10 minutes. Result after 15 minutes is considered as invalid.

INTERPRETATION OF RESULTS



- **Positive (+):** The presence of both "C" line and zone "T" line, no matter T line is clear or vague.
- **Negative (-):** Only clear C line appear. No T line.
- **Invalid:** No colored line appears in C zone. No matter if T line appears.

PRECAUTIONS

- All reagents must be at room temperature before running the assay.
- Do not remove test cassette from its pouch until immediately before use.
- Do not use the test beyond its expiration date.
- The components in this kit have been quality control tested as standard batch unit. Do not mix components from different lot numbers.
- All specimens are of potential infection. It must be strictly treated according to the rules and regulations by local states.

LIMITATION

The Avian Leukemia Virus Antigen Rapid Test is for in vitro veterinary diagnosis use only. All result should be considered with other clinical information available with veterinarian. It is suggested to apply a further confirmative method such as RT-PCR when positive result was observed.



J&G Biotech Ltd

326 Cleveland Road, London

England E18 2AN, United Kingdom

PN:20221128JG405