



# SZABO SCANDIC

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## Produktinformation



Forschungsprodukte & Biochemikalien



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Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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**Rotavirus Antigen Lateral Flow Assay Kit**

**Catalog No:** E-AD-C098

40T

This manual must be read attentively and completely before using this product.

If you have any problems, please contact our Technical Service Center for help.

Toll-free: 1-888-852-8623    Tel: 1-832-243-6086    Fax: 1-832-243-6017

Email: [techsupport@elabscience.com](mailto:techsupport@elabscience.com)

Website: [www.vetassay-elab.com](http://www.vetassay-elab.com)

Please kindly provide us the lot number (on the outside of the box) of the kit for more efficient service.

## Test principle

This kit applies the principle of Immunochromatography assay. The sample will move together with the colloidal gold marker along the chromatography membrane. If Rotavirus (RV) Antigen exist in the samples, it will combine with the colloidal gold marker to combine with the RV antibody, then the detection line will appear a color. Otherwise, it will not show the color reaction.

## Kit components

Item	Specification
Detection Card (with disposable dropper)	40T
Cotton Swab	1 package
Sample Diluent	40 vials
Manual	1 copy

Note: All reagent bottle caps must be tightened to prevent evaporation and microbial pollution.

## Notes

1. FOR RESEARCH USE ONLY. Please read the manual carefully before use, changes of operation may result in unreliable results.
2. Do not use product out of date or in a broken aluminum foil, it is disposable and cannot be used repeatedly.
3. The detection card should be brought to room temperature before opening after take it out from the refrigerator. The opening detection card should be used as soon as possible.
4. Do not use water, PBS or serum of other animals or other 'not required by sample' liquid as the negative control.
5. The tested sample should be fresh and clear. Avoid of using samples of turbidity, polluted, high hemolysis or abnormal viscous.
6. Avoid of touching the chromatography membrane of the sample well and test well.
7. The waste of experiment should be considered as contaminant, and must be properly handled according to the local regulations.
8. Each reagent is optimized for use in the E-AD-C098. Do not substitute reagents from any other manufacturer into the test kit. Do not combine reagents from other E-AD-C098 with different lot numbers.

## Storage and expiry date

**Storage:** Store at 2-30°C. With cool and dry environment, avoid freeze.

**Expiry date:** expiration date is on the packing box.

## Sample preparation

1. **Feces:** Use cotton swab to collect fecal samples that have just been excreted, or directly from the rectum. If rotavirus screening is performed on animals without diarrhea symptoms, rectal sampling should be used. It is not recommended to collect dry fecal samples, because collecting dry fecal samples often results in false positives due to excessive sample collection.



**Note:** Correct sample collection

2. Insert the swab into the **Sample Diluent** tube and mix the swab until the sample has been dissolved into the assay diluent (Approximately 10 sec).
3. Wait for 1 minute to settle down the large particles.

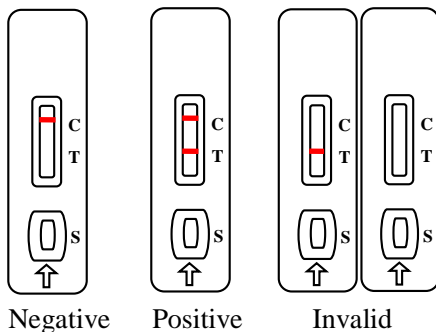
## Assay procedure

Allow all kit components and sample to reach room temperature prior to testing.

1. Tear the aluminum foil bag of the detection card and take out the detection card, and put it on a smooth, clean table.
2. Take the sample with the disposable dropper, add 5 drop (about 100 $\mu$ L) of sample solution to the sample well vertically and slowly.
3. Incubate for 10 to 15 minutes and then judge the results immediately.

## Judgment of result

1. **Negative:** Only the control line region (C) shows a line in the observation well.
2. **Positive:** Both the test line region (T) and the control line region (C) show a line in the observation well.
3. **Invalid:** No line shows in the observation well of the control line region (C).



### **Interpretation of the results**

1. The negative result reveals that there is no RV antigen in the sample. If there is a corresponding acute symptom, then RV infection cannot be excluded.
2. The positive result reveals that there is RV antigen in the sample. It might be infected with RV, and the result should be combined with other methods to analyze.

### **Limitations**

1. This kit can only be used for qualitative detection of RV-Ag in swine and cattle.
2. The detection results of this kit are only for reference. For confirmation of the result, please combine the symptoms and other methods of detection, this detection cannot be used as the only criteria for result.