VITASSAY

Rotavirus+Adenovirus+ Astrovirus+Norovirus

Rapid test for the simultaneous qualitative detection of rotavirus, adenovirus, astrovirus and norovirus in human stool samples.

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For professional in vitro diagnostic use only.

INTENDED USE

Vitassay Rotavirus + Adenovirus + Astrovirus + Norovirus is a rapid, immunochromatographic assay for the simultaneous qualitative detection of rotavirus, adenovirus, astrovirus and norovirus in human stool samples.

Simple, non-invasive and highly sensitivity immunoassay to make a presumptive diagnosis of rotavirus, adenovirus, astrovirus and/or norovirus infection.

INTRODUCTION

Acute gastroenteritis (AGE) is a common cause for hospital/clinic visits and hospitalisations in children as well as adults. It is characterised by diarrhoea, vomiting and abdominal pain that may lead to hypovolemic shock and dehydration, and subsequently death in severe cases.

Watery diarrhea occurs several times a day. Rotavirus, adenovirus, astrovirus and norovirus infection occasionally leads to severe dehydration in infants and children. Symptoms of dehydration include lethargy, dry, cool skin, absence of tears when crying, dry mouth, sunken eye and extreme thirst.

In general, the symptoms begin 1 to 2 days following infection with a virus that causes gastroenteritis and may last from 1 to 10 days, depending on which virus causes the illness (Rotavirus 3 days, Adenovirus 5-8 days and Astrovirus 3 days).

Children in developing countries are particularly at risk of both morbidity and mortality. Worldwide, gastroenteritis affects 3 to 5 billion children each year, and accounts for 1.5 to 2.5 million deaths per year or 12% of all deaths among children less than 5 years of age. In developed countries AGE accounts for 300 deaths per year .

Four major viral pathogens have been associated with AGE. These include three RNA viruses (rotavirus, norovirus, and astrovirus) and one DNA virus (enteric adenovirus). Saporovirus, Aichivirus and members of enterovirus group also cause AGE but are less common. These viruses are transmitted by the fecal-oral route, via contact with contaminated hands, surfaces and objects

PRINCIPLE

Vitassay Rotavirus + Adenovirus +Astrovirus+ Norovirus is a qualitative immunochromatographic assay to make a presumptive diagnosis of rotavirus, adenovirus, astrovirus and/or norovirus infection.

Strip A: The test line zone of the nitrocellulose membrane is precoated with monoclonal antibodies against rotavirus.

Strip B: The test line zone of the nitrocellulose membrane is precoated with monoclonal antibodies against adenovirus.

Strip C: The test line zone of the nitrocellulose membrane is precoated with monoclonal antibodies against astrovirus.

Strip D: The test line zone of the nitrocellulose membrane is precoated with monoclonal antibodies against norovirus.

During the process, the sample reacts with the antibodies against rotavirus (strip A) and/or adenovirus (strip B) and/or astrovirus (strip C), and/or norovirus (strip D) forming conjugates. The mixture moves upward on the membrane by capillary action. If the sample is rotavirus positive, antibodies present on the membrane (test line) capture the conjugate complex and a red line will be visible in the strip A, if the sample is adenovirus positive, antibodies present on the membrane (test line) capture the conjugate complex and a red line will be visible in strip B, if the sample is astrovirus positive, antibodies present on the membrane (test line) capture the conjugate complex and a red line will be visible in strip C and if the sample is norovirus positive, antibodies present on the membrane (test line) capture the conjugate complex and a red line will be visible in strip D. Although the sample is positive or negative, the mixture continues to move across the membranes and the green control line always appears (for all the strips).

The presence of these green lines (in the control zone (C)) indicates that sufficient volume is added; proper flow is obtained and serves as an internal control for the reagents.

PRECAUTIONS

- For professional in vitro use only.
- Do not use after expiration date.
- Do not use the test if its pouch is damaged.
- Specimens should be considered as potentially hazardous and handle in the same manner as an infectious agent. A new test must be used for each sample to avoid contaminations errors.
- Tests should be discarded in a proper biohazard container after testing.
- Reagents contain preservatives. Avoid any contact with the skin or mucous membrane. Consult safety data sheet, available on request.
- Components provided in the kit are approved for use with the Vitassay Rotavirus + Adenovirus + Astrovirus+ Norovirus. Do not use any other commercial kit component.
- Follow Good Laboratory Practices, wear protective clothing, use disposal gloves, goggles and mask. Do not eat, drink or smoke in the working area.

STORAGE AND STABILITY

Store as packaged in the sealed pouch either at refrigerated or room temperature (2-30°C/35.6-86°F).



The test is stable until the expiration date printed on the sealed pouch.

The test must remain in the sealed pouch until use.

Do not freeze.

MATERIALS

MATERIAL PROVIDED	MATERIAL REQUIRED BUT NOT PROVIDED
10 tests/kit Vitassay Rotavirus + Adenovirus + Astrovirus+Norovirus	Specimen collection container.Disposable gloves.Timer.Spatula.
Instructions for use. 10 vials with diluent for sample dilution.	

SPECIMEN COLLECTION

Stool samples should be collected in clean and dry containers. Collect sufficient quantity of feces: 1-2 g or mL for liquid samples.

The samples can be stored in the refrigerator $(2-8^{\circ}C/36-46.4^{\circ}C)$ for 1-2 days prior to testing. For longer storage, maximum 1 year, the specimen must be kept frozen at $-20^{\circ}C/4^{\circ}F$. The samples will be brought to room temperature before to testing.

Homogenise stool sample as thoroughly as possible prior to preparation.

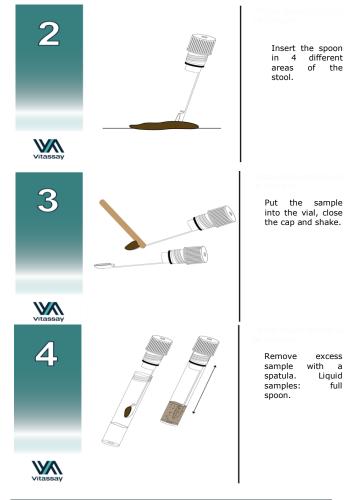
SPECIMEN PREPARATION

- 1 Remove the cap of the vial with diluent for sample dilution (figure 1) and use the spoon to collect sufficient sample quantity. For solid stool, insert the spoon in 4 different areas of the stool sample (figure 2), remove any excess sample with a spatula (figure 3), and place the spoon cap back into the vial for sample dilution (figure 4). For liquid stool, take a spoonful of the sample (figure 3) and transfer it into the vial for sample dilution.
- 2.Close the vial for sample dilution tightly and shake it to dilute and mix the sample with the diluent (figure 4).





Vial for sample dilution



PROCEDURE

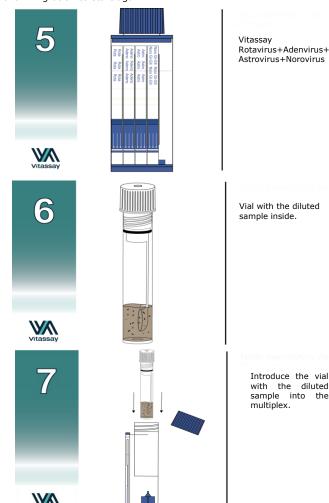
Allow the test, stool sample, controls and diluent to reach room temperature (15-30 $^{\circ}$ C/59-86 $^{\circ}$ F) prior to testing. Do not open pouches until the performance of the assay.

- Shake the vial with the sample vigorously to obtain a good sample dilution.
- 2. Remove the **Vitassay Rotavirus + Adenovirus + Astrovirus+ Norovirus** from its sealed bag just before using it (figure 5).
- 3. Take the vial for sample dilution containing the diluted sample (figure 6), place it inside the multiplex tube (figure 7). Screw the

cap of the multiplex tube tightly (figure 8). The bottom of the vial for sample dilution will break and the diluent+sample solution reaches the sample zone of the strips (figure 9).

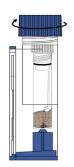
 Leave the multiplex tube vertically on a flat surface and read the results at 10 minutes. Do not read the test results later than 10 minutes.

If the test does not run due to solid particles (the sample is not homogenized), migration process can stop on one or more strips. In this case, tap the end of the multiplex tube on hard surface to allow migration to start again.









Close the cap and the bottom of the diluent vial will break.

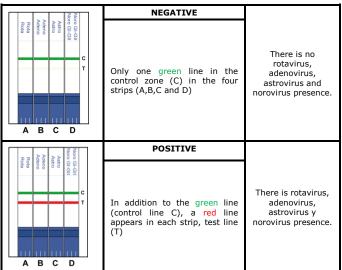


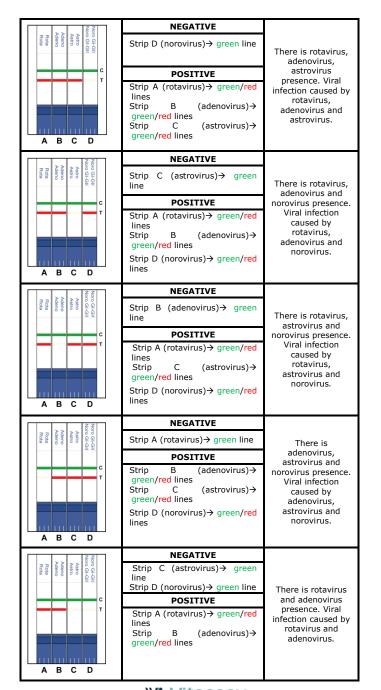


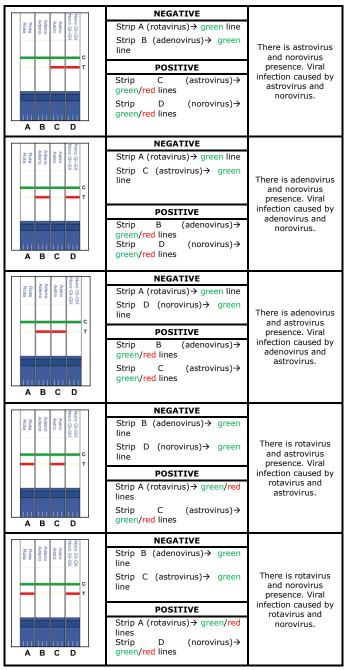
Reaction takes place. Read results at 10 minutes.

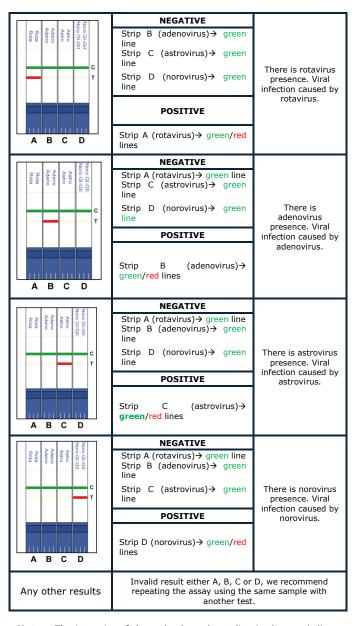
INTERPRETATION OF THE RESULTS

Strip A: rotavirus, Strip B: adenovirus, Strip C: astrovirus and Strip D: norovirus









Notes: The intensity of the red coloured test line in the result line region (T) will vary depending on the concentration of antigens in the specimen.

Positive results detailed in the above table should be followed up with additional confirmatory diagnostic procedures.

Single or dual simultaneous virus infections are more frequent than triple or quadruple.

Invalid results: Total absence of any control coloured lines (green) indicates an invalid result, regardless of the appearance or not of the test lines (red). Wrong procedural techniques or deterioration of the reagents are mostly the main reasons for control line failure. Review the procedure and repeat the assay with a new test. If the problem persists, discontinue using the kit and contact your local distributor.

QUALITY CONTROL

Internal procedural controls are included in **Vitassay Rotavirus+Adenovirus+Astrovirus+Norovirus**. Green lines appearing in the in the results window are internal controls, which confirm sufficient specimen volume and correct procedural technique.

LIMITATIONS

- Vitassay Rotavirus + Adenovirus + Astrovirus+Norovirus must be carried out within 2 hours of opening the sealed bag.
- An excess of stool sample could cause wrong results (brown bands appear). Dilute the sample with the diluent and repeat the test.
- The intensity of test line may vary depending on the concentration of antigens.
- After one week of infection, the number of viruses in faeces is decreasing, making the sample less reactive. Stool samples should be collected within one week of the onset symptoms.
- The use of other samples different from human fecal samples has not been established.
- The quality of Vitassay Rotavirus + Adenovirus + Astrovirus
 + Norovirus depends on the quality of the sample; Proper fecal specimens must be obtained.
- Positive results determine the presence of rotavirus, adenovirus, astrovirus and/or norovirus in fecal samples. A positive result should be followed up with additional laboratory techniques to confirm the results. A confirmed infection should only be made by a physician after the evaluation of all clinical and laboratory findings and must be based in the correlation of the results with further clinical observations.
- Negative results should not be considered as conclusive; it is
 possible that the concentration of antigen is lower than the
 detection limit value. If symptoms or situation still persist a
 rotavirus, adenovirus, astrovirus and/or norovirus determination
 should be carried out with another technique.

EXPECTED VALUES

Viral agents are the most common causes of childhood gastroenteritis over the world.

Enteric viruses have been recognized as the most significant etiological agents of the disease and yet, four categories of viruses are being considered clinically relevant: group A rotavirus (family Reoviridae), norovirus (family Caliciviridae), adenovirus and astrovirus. Several studies proved co-infections in infants 46% with acute watery diarrhoea.

Single infection cases were detected in 335 (34%) of the 973 study children, whereas mixed virus infections were detected in 32 (3.3%) of the same study children. Norovirus was associated in 21 of 32 (66%) mixed infections cases, including 5 premature cases. The most frequent dual gastrointestinal infections were rotavirus and norovirus (50% of 32), adenovirus and rotavirus (16%), rotavirus and astrovirus (13% of 32), norovirus and adenovirus (9% of 32), and norovirus and astrovirus (3%). Surprisingly, we also detected 3 cases of triple gastrointestinal tract infections, 2 of which occurred in two premature children, that were not associated with disease severity criteria such as dehydration or longer duration of hospitalization (not shown). Multiple infections appeared to be detected more significantly during autumnal periods, when norovirus was epidemic (P < 0.001). We observed that the number of *rotavirus* infections appeared to be significantly higher from March through May (P < 0.001) and that *norovirus* and adenovirus infections were detected more frequently during fall and winter months, respectively (P = 0.016 and P = 0.002) in hospitalized children.

PERFORMANCE CHARACTERISTICS

Clinical sensitivity and specificity

An evaluation was performed using **Vitassay Rotavirus + Adenovirus + Astrovirus + Norovirus** and other commercial test (Ridascreen®Rotavirus ELISA Test, r-Biopharm).

Results were as follows:

		Ridascreen®Rotavirus ELISA Test		
		Positive	Negative	Total
Vitassay Rotavirus +	Positive	18	1	19
Adenovirus + Astrovirus+Norovirus	Negative	0	43	43
Rotavirus	Total	18	44	62

Vitassay Rotavirus + Adenovirus + Astrovirus+Norovirus (Rotavirus) vs Ridascreen®Rotavirus ELISA Test					
Sensitivity	Specificity	PPV	NPV		
>99%	98%	>94%	>99%		



And evaluation was performed using **Vitassay Rotavirus + Adenovirus + Astrovirus + Norovirus** and PCR.

Results were as follows:

		PCR		
		Positive	Negative	Total
Vitassay Rotavirus +	Positive	7	0	7
Adenovirus + Astrovirus+Norovirus	Negative	0	52	52
Adenovirus	Total	7	52	59

Vitassay Rotavirus + Adenovirus + Astrovirus+Norovirus (Adenovirus) vs PCR					
Sensitivity	Specificity	PPV	NPV		
>99%	>99%	>99%	>99%		

And evaluation was performed using **Vitassay Rotavirus + Adenovirus + Astrovirus+Norovirus** and an Elisa assay (Ridasscreen@Astrovirus Test, r-Biopharm).

		Ridascreen®Astrovirus Test		us Test
_		Positive	Negative	Total
Vitassay Rotavirus+Adenovirus+	Positive	16	0	16
Astrovirus+Norovirus	Negative	1	11	12
Astrovirus	Total	17	11	28

Vitassay Rotavirus+Adenovirus+Astrovirus+Norovirus (Astrovirus) vs Ridascreen®Astrovirus Test					
Sensitivity	Specificity	PPV	NPV		
>94%	>99%	>99%	>92%		

And evaluation was performed using **Vitassay Rotavirus + Adenovirus + Astrovirus + Norovirus** and other commercial test (Simple Norovirus, Operon).

Results were as follows:

		Simple Norovirus		
		Positive	Negative	Total
Vitassay Rotavirus +	Positive	2	0	2
Adenovirus + Astrovirus+Norovirus	Negative	0	48	48
Norovirus GI	Total	2	48	50

Vitassay Rotavirus + Adenovirus + Astrovirus+Norovirus (Norovirus GI) vs Simple Norovirus					
Sensitivity	Specificity	VPP	VPN		
>99%	>99%	>99%	>99%		

		Simple Norovirus		
_		Positive	Negative	Total
Vitassay Rotavirus + Adenovirus + Astrovirus+Norovirus	Positive	10	0	10
	Negative	0	48	48
Norovirus GII	Total	10	48	58

Vitassay Rotavirus + Adenovirus + Astrovirus+Norovirus (Norovirus GII) vs Simple Norovirus					
Sensitivity	Specificity	VPP	VPN		
>99%	>99%	>99%	>99%		

The results showed that **Vitassay Rotavirus + Adenovirus + Astrovirus+Norovirus** has a high sensitivity and specificity to detect rotavirus, adenovirus, astrovirus and norovirus (GI and GII).

Cross reactivity

No cross reactivity was detected against other gastrointestinal pathogens that are occasionally present in feces.

Adenovirus (Strip A, C and	Escherichia coli	Salmonella paratyphi	
D)	O157:H7	Заппопена рагасурт	
Astrovirus (Strip A, B and	Giardia lamblia	Salmonella	
D)	Giardia lambila	typhimurium	
Campylobacter coli	Hepatitis A	Salmonella typhi	
Campylobacter jejuni	Helicobacter pylori	Shigella boydii	
Clostridium difficile	Listeria monocytogenes	Shigella dysenteriae	
Cryptosporidium parvum	Norovirus (Strip A, B	Shiqella flexneri	
стургозронашт рагчит	and C)	Singena nexiteri	
Enterovirus	Rotavirus (Strip B, C	Shigella sonnei	
Litteroviius	and D)	Siligella sollilei	
Entamoeba hystolitica	RSV	Staphylococcus aureus	
Escherichia coli 0111	Salmonella enteritidis	Yersinia enterocolitica	

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SYMBOLS FOR IVD COMPONENTS AND REAGENTS

IVD	in vitro diagnostic device	*	Keep dry
Ţ i	Consult instructions for use	X	Temperature limitation
\subseteq	Use by	ш	Manufacturer
LOT	Batch code	\sum_{n}	Contains sufficient for <n> test</n>
DIL	Sample diluent	REF	Catalogue number



