



CoproStrip™ C. difficile

GDH + Toxin A + Toxin B

A rapid, one step test for the simultaneous qualitative detection of *Clostridium difficile* Glutamate Dehydrogenase (GDH), Toxin A and Toxin B antigens in human faeces.

Instruction Manual

Test kit for 20 determinations
(Catalog No.41220)

For professional *in vitro* diagnostic use only
Store at 2-30°C. **Do Not Freeze**



Savyon® Diagnostics Ltd.

3 Habosem St. Ashdod 7761003

ISRAEL

Tel.: +972.8.8562920

Fax: +972.8.8523176

E-mail: support@savyondiagnosics.com

Intended Use: The CoproStrip™ C. difficile GDH + Toxin A + Toxin B is a rapid chromatographic immunoassay combo card for the simultaneous qualitative detection of *Clostridium difficile* Glutamate Dehydrogenase (GDH), Toxin A and Toxin B in human faeces that aids in the diagnosis of *C.difficile* infection.

SUMMARY AND EXPLANATION:

The gram-positive anaerobic bacillus *Clostridium difficile* is the leading causative agent of antibiotic-associated diarrhea and pseudomembranous colitis (1). This pathogen is capable of causing disease that could be severe or fatal if not diagnosed on time and treated. Exposure to antibiotics is the major risk factor for *C. difficile* infection. Infection can develop if the normal gastrointestinal flora is disrupted by antibiotic therapy and a person acquires toxin-producing *C. difficile*, typically via the fecal-oral route (2). *C. difficile*'s key virulence factors are toxin A and toxin B (3, 4). These toxins show high sequence and functional homology. Toxin A has been described as a tissue damaging enterotoxin which attracts neutrophils and monocytes and toxin B as a potent cytotoxin that degrades the colonic epithelial cells (5). Most virulent strains produce both toxins, however, toxin A negative/toxin B positive strains are also capable of causing disease (6, 7). *Clostridium difficile* Glutamate Dehydrogenase (GDH) is an enzyme produced in large quantities by all toxigenic and non-toxigenic strains, making it an excellent marker for the organism (8, 9).

PRINCIPLE OF THE PROCEDURE

The CoproStrip™ C. difficile GDH + Toxin A + Toxin B is a qualitative immunoassay for detection of *Clostridium difficile* Glutamate Dehydrogenase (GDH), Toxin A and Toxin B in human fecal specimens.

The membrane of the Test A is pre-coated with monoclonal antibodies against of *Clostridium difficile* (GDH) antigen, the membrane of the Test B is pre-coated with monoclonal antibodies against Toxin A of *Clostridium difficile* antigens and the membrane of the Test C is pre-coated with monoclonal antibodies against Toxin B of *Clostridium difficile* antigens on the test lines region. During testing, the sample reacts with the red colored particles coated with anti-GDH antibodies in the Test A and/or with anti-Toxin A antibodies in the Test B and/ or with anti-Toxin B antibodies in the Test C, which were pre-dried on the test strips. The mixture moves upward on the membrane by capillary action. In the case of a positive result in the Test A the specific antibodies present on the membrane will react with the mixture conjugate and generate one red colored line. In the case of a positive result in the Test B the specific antibodies present on the membrane will react with the mixture conjugate and generate one red colored line. In the case of a positive result in the Test C the specific antibodies present on the membrane will react with the mixture conjugate and generate one red colored line. The mixture continues to move across the membrane to the immobilized antibody places in the control band region. A green colored band always appears in the control lines and serves as verification that sufficient volume was added, that proper flow was obtained and as an internal control for the reagents.

MATERIALS PROVIDED

- CoproStrip™ C. difficile GDH+ Toxin A+ Toxin B devices
- Instructions for use
- Sample collection vial with buffer

MATERIALS NOT PROVIDED

- Specimen collection container
- Disposable gloves
- Timer

WARNING AND PRECAUTIONS

- For professional *in vitro* diagnostic use only.
- Do not use after expiration date.
- The test should remain in the sealed pouch until use.
- Do not use the test if pouch is damaged.
- Follow Good Laboratory Practices, wear protective clothing, use disposal gloves, do not eat, drink or smoke in the area.
- All the specimens should be considered potentially hazardous and handled in the same manner as an infectious agent.
- The test should be discarded in a proper biohazard container after testing.
- The test must be carried out within 2 hours of opening the sealed aluminium pouch.

STORAGE AND STABILITY

Store as packaged in the sealed pouch either at refrigerated or room temperature (2-30°C). The test is stable through the expiration date printed on the sealed pouch. The test must remain in the sealed pouch until use. Do not freeze.

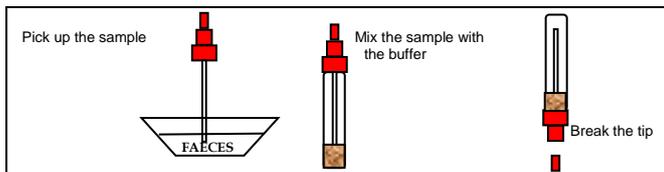
SPECIMEN COLLECTION AND PREPARATION

Collect sufficient quantity of feces (1-2 g or mL for liquid sample). Stool samples should be collected in clean and dry containers (no preservatives or transport media). The samples can be stored in the refrigerator (2-8°C) for 24 hours prior to testing. For longer storage the specimen must be kept frozen at -20°C. In this case, the sample will be totally thawed, and brought to room temperature before testing.

To process the collected stool samples (see illustration 1):

Use a separate specimen collection vial for each sample. Unscrew the cap of the vial and introduce the stick into the faecal specimen to pick up the sample (approx. 125 mg). Close the vial with the buffer and stool sample. Vortex the vial for 15 seconds in order to assure good sample dispersion. For liquid stool samples, aspirate the fecal specimen with a dropper and add 125uL into the specimen collection vial with buffer.

Illustration 1



PROCEDURE

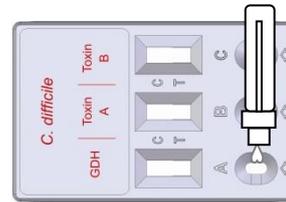
Allow the tests, stool samples and buffer to reach to room temperature (15-30°C) prior to testing. Do not open pouches until ready to perform the assay.

1. Remove the Device from its sealed pouch and use it as soon as possible.
2. Shake the specimen collection vial to assure good sample dispersion. Break off the tip of the vial.
3. Use a separate device for each sample. Dispense exactly 4 drops or 100 uL into each of the specimen wells (i.e. GDH well, Toxin A well, and Toxin B well, marked as S). Start the timer.
4. Read the results at **10 minutes** after dispensing the sample.

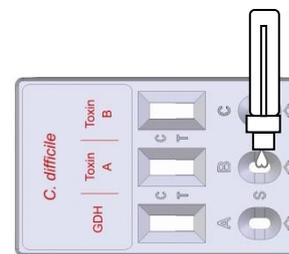
If the test does not run due to solid particles, stir the sample added in the sample window (S) with the stick. If it doesn't work, dispense a drop of diluents until seeing the liquid running through the reaction zone.

Illustration 2

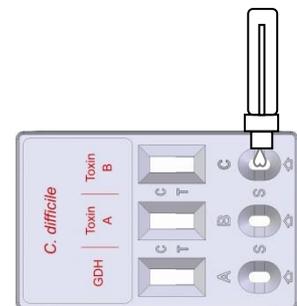
Add 4 drops of the diluted mixture in each circular window



GDH Test- well A



Toxin A Test- well B

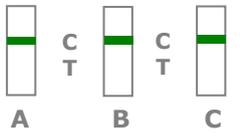
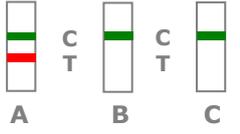
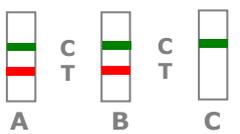
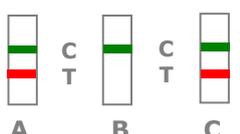
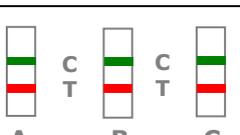
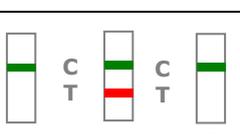
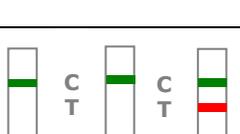
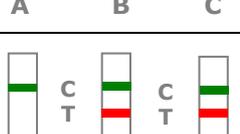


Toxin B Test- well C

INTERPRETATION OF RESULTS

Note: The test red line intensity may vary between the different test strips. Appearance of any intensity of the red line indicates a positive result

Illustration 3

	Indicated Result	Interpretation
1.		<p>GDH, Toxin A and Toxin B of are negative.</p> <p>No <i>C. difficile</i> infection.</p>
2.		<p>GDH positive. Toxin A and Toxin B are negative.</p> <p>Indication of <i>C. difficile</i> infection.</p>
3.		<p>GDH and Toxin A are positive. Toxin B is negative.</p> <p>Indication of <i>C. difficile</i> infection. Further analysis of the sample by CE/FDA cleared assay is recommended.</p>
4.		<p>GDH and Toxin B are positive. Toxin A is negative.</p> <p><i>C. difficile</i> infection.</p> <p>Indication of <i>C. difficile</i> infection.</p>
5.		<p>GDH, Toxin A and Toxin B are positive.</p> <p>Indication of <i>C. difficile</i> infection.</p>
6.		<p>GDH and Toxin B are negative. Toxin A is positive.</p> <p>Repeat the test using a fresh sample. If result is again positive for Toxin A and negative for GDH, further analysis of the sample by CE/FDA cleared assay is recommended</p>
7.		<p>GDH and Toxin A are negative. Toxin B is positive.</p> <p>Repeat the test using a fresh sample. If results are again positive for Toxin B and negative for GDH, further analysis of the sample by CE/FDA cleared assay is recommended</p>
8.		<p>Toxin A and Toxin B are positive. GDH is negative.</p> <p>Repeat the test using a fresh sample. If results are again positive for Toxin B and negative for GDH, further analysis of the sample by CE/FDA cleared assay is recommended</p>
9.	Any other result	<p>Invalid result: either A, B or C, repeat the test using a fresh sample. If results remain the same further analysis of the sample by CE/FDA cleared assay is recommended</p>

INVALID: Total absence of the green control band in one, two or the three Tests (A/B/C) regardless the appearance or not of the red test lines in one or both Tests (A/B/C). Notes: insufficient specimen volume, incorrect procedural techniques or deterioration of the reagents are mostly the main reasons for control lines failure. Review the procedure and repeat the assay using a new test. If the symptoms or situation still persists, discontinue using the test kit and contact your local distributor. See illustration 3 above.

NOTES ON THE INTERPRETATION OF RESULTS

The intensity of the red coloured band in the result line region (T) will vary depending on the concentration of antigens in the specimen. However, neither the quantitative value, nor the rate of increase in antigens can be determined by this qualitative test.

QUALITY CONTROL

Internal procedural controls are included in the test in the form of green lines appearing in the control line region (C). It confirms sufficient specimen volume and correct procedural technique.

LIMITATIONS OF THE PROCEDURE

1. The CoproStrip™ C. difficile GDH + Toxin A+ Toxin B test will only indicate the presence of GDH, Toxin A and/or B in the specimen (qualitative detection). Neither the quantitative value nor the rate of increase in antigens concentration can be determined by this test.
2. An excess of sample could lead to incorrect results (brown bands appear). In such a case, dilute the sample with the buffer and repeat the test.
3. Some stool samples can decrease the intensity of the control bands.
4. The test must be carried out within 2 hours of opening the sealed aluminum pouch.
5. If the test result is negative and clinical symptoms persist, additional testing using other clinical methods is recommended. A negative result does not at any time preclude the possibility of infection caused by *Clostridium difficile*.
6. This test provides a presumptive diagnosis of infection caused by *Clostridium difficile*. All results must be interpreted together with other clinical information and laboratory findings available to the physician and should be followed up with additional laboratory techniques.

PERFORMANCE CHARACTERISTICS

SENSITIVITY AND SPECIFICITY

A study using stool samples from patients with diarrhea showed the following results: The sensitivity and specificity results using the CoproStrip™ C. difficile GDH + Toxin A + Toxin B test in comparison with other commercial immunoassays test (IC test: C. DIFF QUIK CHEK Complete® TechLab) were: Sensitivity >99% and specificity >99%.

ANALYTICAL SENSITIVITY

The cut-off for the assay was established at concentrations of 0.5 ng/ml for toxin A, 0.78 ng/ml for toxin B, and 0.39 ng/ml for glutamate dehydrogenase (GDH).

CROSS-REACTIVITY

An evaluation was performed to determine the cross reactivity of CoproStrip™ C. difficile GDH+Toxin A + Toxin B Device. There is no cross reactivity with common gastrointestinal microorganisms occasionally present in feces.

- *Campylobacter spp*
- *E. coli O157:H7*
- *Listeria monocytogenes*

- *Helicobacter pylori*
- *Shigella spp.*
- *Salmonella spp*
- *Yersinia spp*
- *Yersinia enterocolitica*

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Savyon® Diagnostics Ltd.

3 Habosem St. Ashdod 7761003
ISRAEL

Tel.: +972.8.8562920

Fax : +972.8.8523176

E-mail: support@savyondiagnosics.com



European Authorized Representative: Obelis s.a.

Boulevard Général Wahis 53, B-1030 Brussels

Tel: +32.2.732.59.54 Fax: +32.2.732.60.03

E-mail: mail@obelis.net

Symbols for IVD components and Reagents			
	Manufacturer		For <i>in vitro</i> diagnostic use only
	Authorized representative		Consult instructions for use
	Contains sufficient for <n> tests		Keep dry
	Catalogue Code		Temperature limitation
	Lot Number		Use by
	Sample diluent		