Clostridium difficile GDH ELISA Kit

Catalog Number KA3381
96 assays
Version: 03

Intended for research use only
Table of Contents

**Introduction** .................................................................................................................. 3
  Intended Use ....................................................................................................................... 3
  Background ......................................................................................................................... 3
  Principle of the Assay ......................................................................................................... 3

**General Information** .................................................................................................... 5
  Materials Supplied ............................................................................................................. 5
  Storage Instruction ............................................................................................................. 5
  Materials Required but Not Supplied ................................................................................ 5
  Precautions for Use .......................................................................................................... 6

**Assay Protocol** ............................................................................................................. 7
  Reagent Preparation .......................................................................................................... 7
  Sample Preparation .......................................................................................................... 7
  Assay Procedure .............................................................................................................. 8

**Data Analysis** ............................................................................................................... 10
  Calculation of Results ...................................................................................................... 10
  Performance Characteristics ........................................................................................... 10

**Resources** .................................................................................................................... 12
  References ......................................................................................................................... 12
  Plate Layout ...................................................................................................................... 13
Introduction

Intended Use

*Clostridium difficile* GDH ELISA Kit is an Enzyme-Linked Immunosorbent Assay (ELISA) for use as a screening test to detect *Clostridium difficile* glutamate dehydrogenase (GDH) in human fecal specimens from individuals suspected of having *C. difficile* disease. The test does not distinguish between toxinogenic and nontoxinogenic *C. difficile* strains and is intended to aid in the detection of *C. difficile*. Additional tests should be conducted to detect *C. difficile* toxins.

Background

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). All strains of *C. difficile* produce high levels of GDH (8, 9). Therefore, *C. difficile*'s GDH enzyme is considered a very good antigen marker for detection of this organism. The *Clostridium difficile* GDH ELISA Kit is a highly specific and sensitive detection kit for GDH in stool specimens. A positive result confirms the presence of *C. difficile* and a negative result indicates its absence. A separate test should be performed to confirm the presence of *C. difficile* toxins in the GDH positive samples.

Principle of the Assay

- Break-apart microwells are coated with *C. difficile* GDH specific monoclonal antibodies.
- A horseradish peroxidase (HRP) conjugated monoclonal anti-GDH antibody is added to the antibody-coated microwells.
- Fecal samples are diluted in sample diluent and added to the microwells. In this step *C. difficile* GDH is marked by the HRP conjugated antibody and immobilized by the coated antibody.
- Unbound conjugate is removed by washing.
- TMB-substrate is added, the substrate is hydrolyzed by the peroxidase and yields a blue solution of the reduced substrate.
- Upon the addition of the stop solution, the blue color turns yellow and should be read by an ELISA reader at a wavelength of 450/620 nm.
The absorbance is proportional to the level of *C. difficile* GDH in the sample.

**Summary of Procedure**

**Manual procedure:**

Wells of microtiter plate coated with anti-*C. difficile* GDH antibodies

↓

Add 50 μl of HRP-Conjugate (Ready to Use)

↓

Add 100 μl of Negative Control (Sample Diluent), 100 μl of Positive Control and 100 μl of diluted specimens

↓

Cover plate and incubate 50 min at 37º C at 100% humidity

↓

Wash 5 times with Wash Buffer (300 μl)

↓

Add 100 μl of TMB-Substrate

↓

Cover plate and incubate 15 min at room temperature

↓

Add 100 μl of Stop Solution

↓

Read absorbance at OD 450/620 nm

↓

Interpret results

*Automation procedure:

Please follow the following recommendations to ensure the high quality results of the test when using an automation procedure

✓ Fill the plate in three sets (up to 4 strips at a time). Add the HRP-Conjugate solution to the first set followed by adding the diluted samples before moving to the next set.

✓ 30 minutes sample incubation at 37 ºC.
General Information

Materials Supplied

List of component

<table>
<thead>
<tr>
<th>Component</th>
<th>Amount</th>
</tr>
</thead>
<tbody>
<tr>
<td>Microtiter plate coated with anti-GDH monoclonal antibody: 96 break-apart wells (8x12) coated with monoclonal antibody specific for GDH, packed in an aluminum pouch containing a desiccant card.</td>
<td>96 (8x12) wells</td>
</tr>
<tr>
<td>Concentrated Wash Buffer (20x): PBS - Tween buffer</td>
<td>100 ml</td>
</tr>
<tr>
<td>Sample Diluent: A ready-to-use buffer solution. Contains less than 0.05% Proclin as preservative. The Diluent is also to be used as the negative control solution (see Assay Procedure)</td>
<td>50 ml x 2</td>
</tr>
<tr>
<td>HRP-Conjugate (green): A ready-to-use solution containing Horseradish peroxidase (HRP) conjugated anti-GDH monoclonal antibody. Contains less than 0.05% Proclin as preservative.</td>
<td>7 ml</td>
</tr>
<tr>
<td>Positive Control: A ready to use solution containing recombinant GDH protein. Contains less than 0.05% Proclin as preservative.</td>
<td>2.5 ml</td>
</tr>
<tr>
<td>TMB-Substrate: A ready to use solution contains 3,3',5,5' tetramethylbenzidine as a chromogen and peroxide as a substrate.</td>
<td>16 ml</td>
</tr>
<tr>
<td>Stop Solution: A ready to use solution. Contains 1M H₂SO₄.</td>
<td>16 ml</td>
</tr>
<tr>
<td>Disposable plastic pipettes</td>
<td>100 pc</td>
</tr>
<tr>
<td>Plate cover</td>
<td>1 unit</td>
</tr>
</tbody>
</table>

Storage Instruction

✔ The expiration date of the kit is given on the label. Expiration dates for each component are listed on individual labels. The kit should be stored between 2° and 8°C and should be returned to the refrigerator as soon as possible after use. Exposure of originally stoppered or sealed components to ambient temperature for a few hours will not cause damage to the reagents. DO NOT FREEZE!

✔ Unused strips must be resealed in the aluminum pouch with the desiccant card, by rolling the open end and sealing tightly with tape over the entire length of the opening.

Materials Required but Not Supplied

✔ Clean test tubes for dilution of human samples.
✔ Adjustable micropipettes, or multichannel pipettes (50-200 and 200-1000μl ranges) and disposable tips.
✔ Disposable plastic/wooden collectors or teaspoons.
✔ One liter volumetric flask.
✓ One 50ml volumetric cylinder.
✓ Wash bottle.
✓ Absorbent paper.
✓ Vortex mixer.
✓ A 37°C water bath with a lid, or a moisture chamber placed in a 37°C incubator.
✓ ELISA-reader with 450/620nm filter.
✓ Distilled or double deionized water.
✓ For Automation use: A centrifuge equipped with a rotor compatible with sample tubes to be used in the automation machine.

Precautions for Use

✓ Precautions
  • Reagents should be brought to room temperature before use.
  • When handling assay wells, avoid scratching the bottom of the wells because this may result in elevated absorbance readings.
  • Stool samples, microassay wells, micropipette tips and disposable stool collectors and tubes should be handled and disposed of as potentially biohazards after use. Wear gloves when doing the test.
  • Unused microassay wells must be replaced in the re-sealable pouch with the desiccant to protect them from moisture.
  • TMB-Substrate solution is an irritant material to skin and mucous membranes. Avoid direct contact.
  • Diluted sulfuric acid (1M H₂SO₄) is an irritant agent for the eyes and skin. In case of contact with eyes, immediately flush area with water and consult a physician).

✓ Test Limitations
  • Inability to detect GDH in fecal samples may not preclude actual disease but may be caused by other factors such as incorrect sampling handling or storage of stools. It is also possible that GDH levels are below the kit’s limit detection. The Clostridium difficile GDH ELISA Kit detects GDH in stool at levels of ≥ 1 ng/well.
  • The stability of C. difficile GDH in stool samples may be affected. Therefore, it is important to keep samples at 2-8°C soon after collection. Samples that are not analyzed within 48 hours may be frozen and thawed.
  • Some samples may give low detection levels. This could be caused by a number of reasons such as the presence of bacteria at low levels, or by factors in the feces that interfere with GDH or the test. Under these conditions it is recommended to retest samples using fresh specimen.
Assay Protocol

Reagent Preparation

1. Bring all components and specimens to be tested to room temperature. Determine the total number of specimens to be tested. In addition to the specimens, the following must be included in each test: one well of Negative Control (Use Sample Diluent for this purpose) and one well of Positive Control.

2. Withdraw the microtiter plate from its aluminum pouch by cutting one end near the seal. Leave the required number of strips (according to the number of specimens to be tested) in the 96 well frame.

3. Dilute the Concentrated Wash Buffer 1/20 with double-deionized or distilled water. For example, in order to prepare one liter of Wash Buffer, add 50 ml of the Concentrated Wash Buffer to 950 ml of double-deionized or distilled water.

Sample Preparation

✓ Stool Collection

1. Standard collection and handling procedures used in-house for fecal specimens or culture are appropriate.

2. Preserved stool: The test has not been confirmed with specimens after fixation. (e.g. in 10% formalin, Sodium Acetate Formalin (SAF), or Polyvinyl Alcohol (PVA)).

3. Specimens should be kept between 2°C and 8°C and tested within 48 hours after collection. If testing cannot be performed within 48 hours, store samples at -20°C, or lower.

4. Minimize specimen freezing and thawing which may cause degradation/proteolysis of the antigen and result in loss of activity.

✓ Sample Processing

• For Manual Use

1. Set up one dilution tube for each specimen to be tested. 1.5 ml Eppendorf tubes are recommended for this purpose. Add 400 μl Sample Diluent to each tube. Label the tube.

2. Formed samples: Use a wooden applicator stick or a disposable teaspoon to transfer the fecal specimen to the tube. Transfer approximately 0.1 to 0.15 g of specimen (about the size of a small pea) to the Sample Diluent. Mix the collector in the Sample Diluent to remove as much sample as possible and squeeze the collector against the side of the tube to express any residual liquid. Liquid samples: transfer 150 μl of specimen to the tube. Make sure the liquid specimens are evenly suspended.

3. Thoroughly mix (vortex) the fecal specimen to ensure adequate sampling.

4. Let the tube stand for at least 10 minutes but not more than 30 minutes until large particulate matter is precipitated (decantation). Use upper liquid phase for testing.
• For Automation use
1. Set up one sample’s dilution tube for each specimen to be tested (use sample’s tubes compatible with the available automation equipment). Add 800 µl Sample Diluent to each sample’s tube. Label the tube.
2. Formed samples: Use a wooden collector or a disposable teaspoon to add the fecal specimen to the sample’s tube. Transfer approximately 0.2 to 0.3 g of specimen (about the size of 2 small peas) to the sample’s tube. Mix the collector in the Sample Diluent to remove as much sample as possible and squeeze the collector against the side of the tube to extract any residual liquid.
   Liquid samples: transfer 300 µl of specimen to the tube. Make sure the liquid specimens are evenly suspended.
3. Thoroughly mix (vortex) the fecal specimen to ensure adequate sampling.
4. Let the tube stand or at least 10 minutes until large particulate matter is precipitated (decantation).
   Ensure that the formed supernatant does not contain large particulate material. In case and required centrifuge the tubes at 1000 g for 30 sec.
5. Transfer the sample’s tube to the corresponding rack at the automation machine.

Assay Procedure

✓ For Manual Use
• Incubation of stool samples and controls
  1. Dispense 50 µl of ready-to-use conjugate into each well.
  2. Pipette 100 µl of Positive control and 100 µl of Negative Control (i.e., Sample Diluent) into separate wells of the test strip.
  3. Dispense 100 µl of diluted stool samples into separate wells of the test strip using the provided disposable pipettes (the lowest mark on the pipette).
  4. Cover the strips with a plate cover and incubate for 50 min at 37°C in a moisture chamber.
  5. Washing step: Discard the liquid content of the wells. Fill each well with Wash Buffer up to the end of the well (300 µl). Repeat this step 4 times to a total of FIVE times. Automatic washing machine can be used.
  6. Dry the strips and frame by gently tapping them over clean absorbent paper.

• Incubation with TMB Substrate
  1. Dispense 100 µl of TMB-Substrate into each well, cover the strips with a plate cover, and incubate at room temperature for 15 minutes.
  2. Stop the reaction by adding 100 µl of Stop Solution (1M H₂SO₄) into each well.

• Determination of Results
  Determine the absorbance at 450/620 nm and record the results. Determination should not exceed 10 minutes following stopping of the chromogenic reaction.

Note: Any air bubbles should be removed before reading. The bottom of the ELISA plate should be carefully wiped.
For Automation Use

- Incubation of conjugate with stool samples and controls

Dispense ready-to-use conjugate and samples into each well in consecutive sets of up to 4 strips at a time as follows:

1. Dispense 50 μl of ready-to-use conjugate into each well of up to 4 strips.
2. Pipette 100 μl of Positive control and 100 μl of Negative Control (i.e., Sample Diluent) into separate wells of the test strip (containing the ready-to-use conjugate).
3. Dispense 100 μl of diluted stool samples into separate wells of the test strip using the provided disposable pipettes (the lowest mark on the pipette).
4. Repeat dispensing of ready-to-use conjugate and samples to next 4 strip sets as described in steps 1 and 3.
5. Incubate the plate at 37°C for 30 minutes.
6. Perform 5X 300 μl wash cycles using the pre-diluted Wash Buffer.
7. Perform 2 aspirate cycles with aspirate sweep.

- Incubation with TMB Substrate

8. Dispense 100 μl of TMB-Substrate into each well, cover the strips with a plate cover, and incubate at room temperature for 15 minutes.
9. Stop the reaction by adding 100 μl of Stop Solution (1M H₂SO₄) into each well.

- Determination of Results

Determine the absorbance at 450/620 nm and record the results.

Please note that each automation machine has specific technical commands. Please implement this automation procedure for this kit on the operation protocol of your automation equipment.
Data Analysis

Calculation of Results

✔ Test Validation
For the test to be valid the following criteria must be met. If these criteria are not met the test should be considered invalid and should be repeated.

✔ Negative Control:
The absorbance value should be ≤ 0.07 at 450/620 nm.

✔ Positive Control:
The absorbance value should be ≥ 0.5 at 450/620 nm.

✔ Interpretation of Results
• Spectrophotometric Dual Wavelength at 450/620 nm
  Negative = OD < 0.08
  Positive = OD ≥ 0.08

Performance Characteristics

Study 1 and 2: Stool specimens were evaluated by Clostridium difficile GDH ELISA Kit. Study 1 was performed in house on a total of 90 samples (Table 1) and study 2 externally in an Israeli medical center on a total of 56 samples (Table 2). The results shown below were compared to results of an FDA approved commercial reference ELISA kit.

✔ Table 1

<table>
<thead>
<tr>
<th></th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Positive</td>
</tr>
<tr>
<td>Clostridium difficile</td>
<td>57</td>
</tr>
<tr>
<td>GDH ELISA Kit</td>
<td>3</td>
</tr>
<tr>
<td></td>
<td>60</td>
</tr>
</tbody>
</table>

Sensitivity: 95 %, Specificity: 100 %
Study 3: Stool specimens were evaluated by *Clostridium difficile* GDH ELISA Kit. The study was performed externally at a reference laboratory for Clostridium difficile, in Paris on a total of 298 samples (Table 3). The results shown below were compared to results of culture on selective medium as the gold standards.

<table>
<thead>
<tr>
<th></th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Positive</td>
</tr>
<tr>
<td><strong>Clostridium difficile GDH ELISA Kit</strong></td>
<td>31</td>
</tr>
<tr>
<td>Positive</td>
<td>1</td>
</tr>
<tr>
<td>Negative</td>
<td>32</td>
</tr>
</tbody>
</table>

Sensitivity: 96.9%, Specificity: 92.9%

Cross Reactivity and Interference by Mixed infections

The *Clostridium difficile* GDH ELISA Kit was evaluated using microbial culture isolates and stool samples*. No cross-reactivity was observed with any of the gastrointestinal pathogens and microbes listed below:

*Blastocystis, Campylobacter*, *Cryptosporidium parvum*, *Dientamoeba fragilis*, *Escherichia coli*, *Entamoeba histolytica*, *Enterococcus faecali, Enterococcus faesium, Enterococcus avium, Enterococcus aerogenes, Enterococcus cloacae, Enterococcus gallinarum, Enterococcus durans, Giardia lamblia*, *Helicobacter pylori*, *Klebsiella pneumonia Samonella enterica*, and *Shigella*.*
Resources

References


## Plate Layout

<table>
<thead>
<tr>
<th></th>
<th>A</th>
<th>B</th>
<th>C</th>
<th>D</th>
<th>E</th>
<th>F</th>
<th>G</th>
<th>H</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>3</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>4</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>5</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>6</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>7</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>8</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>9</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>10</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>11</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>12</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>