Trypanosoma cruzi (E coli, Recombinant) Antigen

Customer Service: Contact your local representative or find country specific contact information on www.abbottdiagnostics.com

Package insert instructions must be carefully followed. Reliability of assay results cannot be guaranteed if there are any deviations from the instructions in this package insert.

See REAGENTS section for a full explanation of symbols used in reagent component naming.
NAME AND INTENDED USE
ABBOTT ESA Chagas is an in vitro enzyme strip assay intended for the qualitative detection of antibodies to Trypanosoma cruzi (T cruzi) in human serum and plasma specimens. The assay is intended for use as an additional, more specific test on human serum or plasma specimens found to be repeatedly reactive using a licensed screening test for antibodies to T cruzi.

SUMMARY AND EXPLANATION OF THE TEST
Chagas disease or American Trypanosomiasis is caused by the parasite T cruzi. There are 3 morphologic forms in the life cycle of T cruzi: epimastigote (multiplying extracellular form in mammalian hosts); and trypanomastigote (multiplying intracellular form in mammalian blood and insect feces). The majority of T cruzi proteins are expressed in all 3 morphologic forms. ABBOTT ESA Chagas is based on recombinant proteins FP10, FP6, FP3, and TcF. In aggregate, these 4 hybrid recombinant proteins represent at least 14 distinct antigenic regions that broadly represent all 3 morphologic forms. Moreover, these recombinant proteins also contain epitopes recognized by antibodies present in persons with acute T cruzi infections as well as those with chronic Chagas disease.8

BIOLOGICAL PRINCIPLES OF THE PROCEDURE
ABBOTT ESA Chagas is a multi-step enzyme strip assay.

- Four individually prepared T cruzi recombinant antigens (FP10, FP6, FP3, and TcF), 3 onboard controls (2 onboard visual calibrators [human IgG], and 1 onboard sample addition control [anti-human IgG]), have been applied separately as discrete lines across strips that are composed of nitrocellulose membrane laminated onto a plastic support. These 4 T cruzi recombinant antigens are also used in ABBOTT PRISM Chagas in which they are coated onto the surface of microparticles. Licensed screening assays use the combined reactivity of their representative antigens to give a single composite signal. In ABBOTT ESA Chagas, the reactivity of each recombinant antigen is evaluated individually, resulting in a more specific test. The calibrators (H-CAL and L-CAL) are used to interpret results of the assay and indicate that conjugate was added to the strip. The control indicates that a sample was added to the strip.
- The strips are incubated with sample (either plasma, serum, or ABBOTT ESA Chagas Positive or Negative Control) and specimen diluent in the trough of the incubation tray. During incubation, T cruzi antibodies present in the sample bind to the antigen(s) on the strips.
- After first incubation is complete, the strips are washed with a 1x Wash Buffer. Then a goat anti-human:alkaline phosphatase conjugate is added to the strips and incubated. The conjugate binds antibody to T cruzi that is present.
- After the second incubation is complete, the strips are washed and the enzyme substrate (BCIP/NBT) is added and incubated.
- After incubation with the substrate, the strips are washed and dried. For each sample, color intensity at the location of each recombinant antigen is individually graded against the onboard calibrators to interpret the assay results (see Interpretation of Results).

REAGENTS
ABBOTT ESA Chagas Kit (REF 8L34-68)

- **STRIPS**
  - 1 Bottle (30 strips) T cruzi (Epitope, recombinant) Antigen Coated Strips: Each strip contains 1 goat anti-human IgG specimen control band, 2 human IgG calibrator bands, and 4 individual bands coated with T cruzi antigens. Minimum concentrations for the antigen bands: FP10: 2.5 µg/mL, FP6: 0.35 µg/mL, FP3: 0.2 µg/mL, and TcF: 37.5 µg/mL.

- **CONJUGATE**
  - 1 Bottle (38 mL) Anti-human (Goat):Alkaline Phosphatase Conjugate in TRIS buffer with protein stabilizers and detergent. Minimum concentration: 0.1 µg/mL. Preservative: 0.15% ProClin 950.

- **CONTROL**
  - 1 Bottle (0.25 mL) Negative Control is recalcified, human plasma. Preservative: 0.1% sodium azide and 0.15% ProClin 950.

- **SPECIMEN DILUENT**
  - 1 Bottle (38 mL) Specimen Diluent containing TRIS buffer with protein stabilizers and surfactants. Preservative: 0.15% ProClin 950.

- **CONCENTRATED WASH BUFFER**
  - 1 Bottle (38 mL) Concentrated Wash Buffer containing TRIS buffer and detergent. Preservative: 0.15% ProClin 950.

- **SUBSTRATE TABLETS**
  - 10 BCP/NBT Substrate Tablets (5-bromo-4-chloro-3-indoly phosphate/nitro blue tetrazolium).

- **INCUBATION TRAYS**
  - 10 Incubation Trays.

WARNINGS AND PRECAUTIONS

- **IVD**
  - For In Vitro Diagnostic Use
  - The performance characteristics of this product have not been established for the laboratory diagnosis of T cruzi infection.
  - Package insert instructions must be carefully followed. Reliability of assay results cannot be guaranteed if there are any deviations from the instructions in this package insert.

Safety Precautions

- CAUTION: This product contains human-sourced and/or potentially infectious components. Refer to the REAGENTS section of this package insert. No known test method can offer complete assurance that products derived from human sources or inactivated microorganisms will not transmit infection. Therefore, all human-sourced materials should be considered potentially infectious. It is recommended that these reagents and human specimens be handled in accordance with the OSHA Standard on Bloodborne Pathogens. Biosafety Level 2 or other appropriate biosafety practices should be used for materials that contain or are suspected of containing infectious agents. These precautions include, but are not limited to, the following:
  - Wear gloves when handling specimens or reagents.
  - Do not pipette by mouth.
  - Do not eat, drink, smoke, apply cosmetics, or handle contact lenses in areas where specimens and reagents are handled.
  - Clean and disinfect all spills of specimens or reagents using an appropriate disinfectant such as 0.1% sodium hypochlorite, or other suitable disinfectant.
  - Decontaminate and dispose of all specimens, reagents, and other potentially contaminated materials in accordance with local, state, and federal regulations.
  - The human plasma used in the negative control is nonreactive for antibodies to T cruzi, HBsAg, HIV-1 RNA or HIV-1 Ag, anti-HIV-1/HIV-2, and anti-HCV.
  - The human plasma used in the positive control contains antibodies to T cruzi, including mouse/human chimeric monoclonal antibody. The plasma is nonreactive for HBsAg, HIV-1 RNA or HIV-1 Ag, anti-HIV-1/ HIV-2, and anti-HCV.
  - The following warnings and precautions apply to the conjugate, controls, specimen diluent, and concentrated Wash Buffer.

Warning: Contains methylisothiazolinones. H317 May cause an allergic skin reaction.

Prevention:
P261 Avoid breathing mist/vapors/spray.
P272 Contaminated work clothing should not be allowed out of the workplace.
P280 Wear protective gloves/protective clothing/eye protection.

Response:
P305+P335 IF ON SKIN: Wash with plenty of water.
P333+P335 If skin irritation or rash occurs: Get medical advice/ attention.
P362+P364 If swallowed: Do not induce vomiting.
P361 Dispose of contents/container in accordance with local regulations.
P501 Dispose of product by methods recommended by the local authorities. P503 Take off contaminated clothing and wash it before reuse.
P504 Take off and wash immediately all contaminated clothing.
P505 Use suitable protective clothing.

- The negative and positive controls contain sodium azide. Contact with acids liberates very toxic gas. This material and its container must be disposed of in a safe way.
The following warnings and precautions apply to the substrate tablets.

**Warning:** Contains Trihydroxymethyl Aminomethane Magnesium Chloride hexahydrate Nitroblue Tetrazolium 5-Bromo-4-chloro-3-indolyl phosphate.

**H303** May be harmful if swallowed.

**H315** Causes skin irritation.

**H319** Causes serious eye irritation.

**H333** May be harmful if inhaled.

**H335** May cause respiratory irritation.

**Prevention:**
- P260 Avoid breathing dust.
- P264 Wash hands thoroughly after handling.
- P271 Use only outdoors or in a well ventilated area.
- P280 Wear protective gloves / protective clothing / eye protection.

**Response:**
- P305+P351+ P337+P313 If eye irritation persists: Get medical advice/attention.
- P302+P352 IF ON SKIN: Wash with plenty of water.
- P304+P340 IF INHALED: Remove victim to fresh air and keep at rest in a position comfortable for breathing.
- P312 Call a POISON CENTER or doctor/physician if you feel unwell.
- P362 Take off contaminated clothing and wash before reuse.

**Storage:**
- P403+P233 Store in a well-ventilated place. Keep container tightly closed.
- P401+P200 Store in a cool, dry place.
- P403+P233 Store in a cool, dry place.
- P403+P233 Store in a cool, dry place.

**Disposal:**
- P501 Dispose of contents/container in accordance with local regulations.

Safety Data Sheets are available at www.abbottdiagnostics.com or contact your local representative.

**Handling Precautions**
- Use caution to avoid microbial and chemical contamination of samples, reagents, and equipment.
- Use disposable pipette tips for all pipetting steps.
- Use a new pipette tip for each specimen.
- Do not interchange bottles or bottle caps between reagents.
- Do not use kits beyond the expiration date.
- Do not mix components and strips from kits with different lot numbers.
- Treat negative and positive controls as potentially infectious.
- Use only the ABBOTT ESA Chagas Positive and Negative Controls provided with the kit.
- Use accurately calibrated equipment.
- Distilled or deionized water (Clinical and Laboratory Standards Institute [CLSI] clinical laboratory reagent water or better) must be used for preparation of the 1x Wash Buffer and substrate. Store water, 1x Wash Buffer, and substrate in nonmetallic containers.
- Use forceps to hold the strip within the identification portion avoiding contact with the nitrocellulose membrane.
- Do not cut strips.
- Do not reuse strips or reaction troughs.
- Do not allow strips to dry out during the procedure, prior to completion of color development.
- To prevent fading, keep developed strips out of strong light (eg, direct sunlight).

**Storage Instructions**
- When stored and handled as directed, reagents are stable until the expiration date.
- Store ABBOTT ESA Chagas Kit at 2-8°C. Do not freeze.
- Do not interchange reagents or strips between kit lots.
- Do not cut strips.
- Do not use heat-inactivated specimens.
- Do not use specimens with abnormal chemical composition or gross lipemia.
- Performance has not been established using plasmapheresis or cadaveric specimens, umbilical cord blood, or body fluids such as urine, saliva, semen, amniotic fluid, cerebrospinal fluid, or pleural fluid.

**Indications of Instability or Deterioration of Reagents**
Changes in the physical appearance of the reagents supplied may indicate instability or deterioration of these materials. If changes in the physical appearance of the reagents are observed (eg, obvious changes in reagent color or cloudiness, which may be associated with microbial contamination), they should not be used.

**SPECIMEN COLLECTION AND PREPARATION FOR ANALYSIS**

**Specimen Types**
- Serum (including specimen collected in serum separator tubes), plasma collected in EDTA, potassium oxalate, sodium citrate, ACD-A, ACD-B, CP2D, CPD, or CPDA-1 anticoagulants, or plasma collected from segmented tubing may be used with ABBOTT ESA Chagas. Follow the manufacturer’s specimen collection instructions for serum and plasma collection tubes.

**CAUTION:** Do not use specimens collected in hepaticin.

**Specimen Conditions**
- This assay was designed and validated for use with individual human serum and plasma specimens. This assay has not been validated for use with pooled specimens.
- Serum from heparinized patients may be incompletely coagulated. Draw the specimen prior to heparin therapy or after heparin therapy is discontinued and activated partial thromboplastin time (aPTT) levels return within normal range.
- Do not use heat-inactivated specimens.
- Do not use specimens with visible microbial contamination or gross lipemia.
- Performance has not been established using plasmapheresis or cadaveric specimens, umbilical cord blood, or body fluids such as urine, saliva, semen, amniotic fluid, cerebrospinal fluid, or pleural fluid.

**Potential Interfering Substances**
- No qualitative performance differences were observed when a minimum of 26 nonreactive donor specimens and 27 reactive donor specimens, which were created by spiking with T cruzi antibody to low-level reactivity, were spiked with potentially interfering substances, creating samples with artificially elevated levels of bilirubin (≤ 20 mg/dL), hemoglobin in plasma (≤ 500 mg/dL), red blood cells (≤ 0.4% v/v), triglycerides (≤ 3,000 mg/dL), or protein (≤ 12 g/dL).
- In an additional study, 20 specimens from patients with elevated levels of endogenous hemoglobin in whole blood (16.2 to 18.1 g/dL), 19 specimens with elevated levels of endogenous triglycerides (1,029 to >10,450 mg/dL), 18 specimens with elevated levels of endogenous total protein (9.1 to 11.2 g/dL), and 20 specimens with elevated levels of endogenous bilirubin (5.1 to 11.2 mg/dL) were spiked with T cruzi antibody to target a low level of reactivity. All of these specimens prior to spiking were negative. All of the spiked specimens remained positive.

**Preparation for Analysis**

**FAILURE TO FOLLOW THE SPECIFIED CENTRIFUGATION PROCEDURE MAY GIVE ERRONEOUS OR INCONSISTENT TEST RESULTS.**

**Nonfrozen specimens** must be centrifuged such that g-minutes are between 30,000 and 75,000, and then may be stored at 2-8°C for up to 7 days. After 7 days, specimens need to be recentrifuged such that g-minutes are between 30,000 and 75,000.

**Previously frozen specimens** must be mixed gently and thoroughly after thawing and centrifuged such that g-minutes are between 30,000 and 75,000.

A refrigerated or nonrefrigerated centrifuge is acceptable for use. The acceptable time and force ranges that meet the criterion of Nonfrozen and Previously frozen specimens are described in the following table.

<table>
<thead>
<tr>
<th>Specimen Type</th>
<th>G-Minutes</th>
<th>Storage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nonfrozen</td>
<td>30,000-75,000</td>
<td>2-8°C</td>
</tr>
<tr>
<td>Previously frozen</td>
<td>30,000-75,000</td>
<td>2-8°C</td>
</tr>
</tbody>
</table>
Centrifugation Time

<table>
<thead>
<tr>
<th>Minutes</th>
<th>RCF (x g)</th>
<th>g-minutes</th>
</tr>
</thead>
<tbody>
<tr>
<td>10</td>
<td>3,000</td>
<td>30,000</td>
</tr>
<tr>
<td>15</td>
<td>2,000 - 3,000</td>
<td>30,000 - 45,000</td>
</tr>
<tr>
<td>20</td>
<td>1,500 - 3,000</td>
<td>30,000 - 60,000</td>
</tr>
<tr>
<td>25</td>
<td>1,300 - 3,000</td>
<td>32,500 - 75,000</td>
</tr>
</tbody>
</table>

Convert rpm to RCF as follows: RCF = 1.12 x \( \frac{rpm}{1000} \)^2

Converting RCF to rpm as follows: rpm = \( \frac{RCF}{1.12 \times \text{max}} \)

NOTE: If custom tube adapters (i.e., adapters not defined by the centrifuge manufacturer) are used, then the radius \( r_{\text{max}} \) should be manually measured in millimeters and the RCF calculated.

Storage and Shipping
- After collection, specimens may be stored at 30°C or colder for up to 7 days, 2-8°C for up to 14 days, or frozen at -20°C or colder for up to 2 months (inclusive of shipping time). Storage at a combination of 2-8°C and 30°C or colder may not exceed 14 days.
- Prior to freezing, the serum or plasma should be removed from the clot or red blood cells to avoid hemolysis.
- Specimens stored at -20°C or colder for greater than 2 months may be used for informational purposes (e.g., lookback testing, discardant sample testing, clinical and validation testing).
- When shipping specimens, package and label specimens in compliance with applicable regulations covering the transport of clinical specimens and infectious substances.
- Thirty nonreactive and 30 low-level reactive specimens showed no qualitative performance differences when subjected to 6 freeze/thaw cycles. However, some specimens that have undergone multiple freeze/thaw cycles, or have been stored frozen for prolonged periods, may give erroneous or inconsistent test results.

PROCEDURE

Materials Provided

- [**NEK** BL34-68] **ABBOTT ESA Chagas Kit**

Materials Required but not Provided

- Distilled or Deionized Water (CLSI clinical laboratory reagent water or better) \(^1\)
- 1 mL Pipette
- 20 \( \mu l \) Pipette
- Repeater Pipette (e.g., 25 mL Eppendorf)
- Microfuge or small centrifuge
- Bi-directional Orbital Rocker capable of maintaining 20-35 rpm
- Nonmetallic containers with caps
- Vacuum-powered aspirator with trap
- Forceps

Reagent Preparation

Before use, bring all reagents to room temperature (15-30°C).

Determine the number of specimens to be tested and the number of strips required. Each run requires the following number of strips:
- one strip for the positive control
- one strip for the negative control
- one strip for each test specimen

NOTE: Each tray holds 8 strips.

1x Wash Buffer

- The 1x Wash Buffer may be prepared before starting the **ABBOTT ESA Chagas Procedure** and must be used within 7 days.
- Gently invert the Concentrated Wash Buffer several times to ensure a homogeneous solution. Avoid foaming.
- Each strip requires 6 mL of 1x Wash Buffer. Make a 1:10 dilution of Concentrated Wash Buffer by adding 1 volume of Concentrated Wash Buffer to 9 volumes of distilled or deionized water in a nonmetallic container with cap and mix thoroughly by gently inverting to ensure a homogeneous solution. Avoid foaming.
- Use the table below as a guide for preparing the 1x Wash Buffer. When processing a number of strips not specified in the table, refer to the next highest maximum number of strips to be processed. For example, for 7 strips, use volumes for processing 8 strips.

<table>
<thead>
<tr>
<th>Maximum Number of Strips to be Processed</th>
<th>Volume of Concentrated Wash Buffer (mL)</th>
<th>Volume of Distilled or Deionized Water (mL)</th>
<th>Total Volume of Prepared 1x Wash Buffer (mL)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>3</td>
<td>2</td>
<td>18</td>
<td>20</td>
</tr>
<tr>
<td>6</td>
<td>4</td>
<td>36</td>
<td>40</td>
</tr>
<tr>
<td>8</td>
<td>5</td>
<td>45</td>
<td>50</td>
</tr>
<tr>
<td>10</td>
<td>7</td>
<td>63</td>
<td>70</td>
</tr>
<tr>
<td>15</td>
<td>10</td>
<td>90</td>
<td>100</td>
</tr>
<tr>
<td>20</td>
<td>12</td>
<td>117</td>
<td>130</td>
</tr>
<tr>
<td>25</td>
<td>16</td>
<td>144</td>
<td>160</td>
</tr>
<tr>
<td>30</td>
<td>19</td>
<td>171</td>
<td>190</td>
</tr>
</tbody>
</table>

* Provides sufficient quantity in slight excess of the required minimum to assure an uninterrupted test procedure.

Substrate

The substrate will be prepared during the **ABBOTT ESA Chagas Procedure** in Step 14.
- Use the substrate within 4 hours of preparation.
- Each strip requires 1 mL of substrate. Dissolve 1 substrate tablet in 20 mL of distilled or deionized water in a nonmetallic container with cap. Swirl container gently to obtain a homogeneous solution. Avoid foaming.
- Use the table below as a guide for preparing substrate.

<table>
<thead>
<tr>
<th>Number of Strips to be Processed</th>
<th>Number of Substrate Tablets</th>
<th>Volume of Distilled or Deionized Water (mL)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>3 - 19</td>
<td>1</td>
<td>20</td>
</tr>
<tr>
<td>20 - 30</td>
<td>2</td>
<td>40</td>
</tr>
</tbody>
</table>

* Provides sufficient quantity in slight excess of the required minimum to assure an uninterrupted test procedure.

**ABBOTT ESA Chagas Procedure**

**Initial Preparation**

1. Read **Handling Precautions** before performing the assay procedure.
2. Allow all test materials to reach room temperature (15-30°C) before performing the assay. Mix reagents thoroughly by gently inverting each component (except strips and substrate tablets) several times to ensure a homogeneous solution. Avoid foaming.
3. The entire assay procedure is performed at room temperature (15-30°C).
4. If not already prepared, prepare 1x Wash Buffer.
5. Using forceps, remove each strip from the container and place it with the filled rectangle facing down into each reaction trough of the incubation tray. Check to ensure that there is only 1 strip in each trough. Each run requires the following number of strips:
   • one strip for the positive control
   • one strip for the negative control
   • one strip for each test specimen
 NOTE: Each tray holds 8 strips.

6. Prepare a record to identify the numbers on the strips (ID) with the corresponding specimen identification numbers.

7. Place each incubation tray on the rocker.

Specimen Incubation

8. Add 1 mL of specimen diluent to each reaction trough containing a strip. Use forceps to gently submerge each strip in specimen diluent, touching the strip at the ID end.

9. Incubate each tray on the rocker at 20-35 rpm for 5-7 minutes to allow the strips to become saturated with specimen diluent. After incubation, remove the incubation tray(s) from the rocker.
 NOTE: The motion of the diluted specimen or controls over the strips, generated by the rocker, is important in achieving optimum performance of the assay. Periodically check to ensure that a rocking motion is maintained throughout the incubations. Improper functioning of the rocker, which may affect antibody binding, will invalidate the test results and require that the assay be repeated.

10. Deliver 20 µL of each specimen or each control to a single reaction trough containing strip and specimen diluent.

11. Cover the incubation tray(s). Incubate each tray on the rocker at 20-35 rpm for 2 hours ± 5 minutes.
 After incubation, remove the incubation tray(s) from the rocker. Remove the tray cover(s) and aspirate the liquid (at the strip ID end) from each reaction trough.

12. Wash the strips by adding 1 mL of 1x Wash Buffer to each reaction trough containing a strip. Return the incubation tray(s) to the rocker and incubate at 20-35 rpm for 4-5 minutes.

13. After incubation, remove the incubation tray(s) from the rocker and aspirate the liquid (at the strip ID end) from each reaction trough.
 Repeat this step two more times (3 washes total).

Conjugate Incubation

14. After washing the strips, add 1 mL of conjugate to each reaction trough containing a strip.

15. Cover the incubation tray(s). Incubate each tray on the rocker at 20-35 rpm for 1 hour ± 5 minutes.
 After incubation, remove the incubation tray(s) from the rocker. Remove the tray cover(s) and aspirate the liquid (at the strip ID end) from each reaction trough.

16. Wash the strips by adding 1 mL of 1x Wash Buffer to each reaction trough containing a strip. Return the incubation tray(s) to the rocker and incubate at 20-35 rpm for 4-5 minutes.

17. After incubation, remove the incubation tray(s) from the rocker and aspirate the liquid (at the strip ID end) from each reaction trough.
 Repeat this step two more times (3 washes total).

Substrate Incubation

18. After aspiration of 1x Wash Buffer, add 1 mL of substrate to each reaction trough containing a strip.

19. Using forceps, transfer strips to a paper towel with the filled rectangle facing up and aligned in a single direction. Gently blot dry with a paper towel. Then allow the strips to air dry for at least 30 minutes.

20. Interpret the results within 24 hours and 30 minutes of removal from the reaction trough. Record the results for each specimen and control as described in QUALITY CONTROL PROCEDURES and RESULTS.

QUALITY CONTROL PROCEDURES
The Abbott ESA Chagas Positive and Negative Controls must be included in each run of test specimens. A run may include any number of specimens, strips, or incubation trays. The controls are not run with each incubation tray.

Each strip includes onboard controls that must be evaluated before assessment of recombinant antigen bands. Human IgG low-level calibrator (L-CAL) and human IgG high-level calibrator (H-CAL) are used to interpret results of the assay. The anti-human IgG specimen addition control (SPM-CTL) is included to indicate that a specimen was added to the strip. If the negative and/or positive controls do not meet all of the criteria below, then the control result and associated test results are invalid. The samples must be retested.

Negative Control

Onboard Controls

• Must display clearly distinguishable onboard L-CAL and H-CAL bands.
• The L-CAL band (at the visual cutoff, 1+) must be clearly lighter (less intense) than the H-CAL band (at scoring of 3+).
• Must display a clearly distinguishable SPM-CTL band, with color intensity greater than the L-CAL (2+ or greater).

Antigen Bands

• Must not display a visible band at any of the four recombinant antigen locations.

The strips MUST NOT display uneven background color or colored deposits across any portion of seven possible visible bands. Some uneven background color or colored deposits between the bands or at the ends of the strip are acceptable.

Positive Control

Onboard Controls

• Must display clearly distinguishable onboard L-CAL and H-CAL bands.
• The L-CAL band (at the visual cutoff, 1+) must be clearly lighter (less intense) than the H-CAL band (at scoring of 3+).
• Must display a clearly distinguishable SPM-CTL band, with color intensity greater than the L-CAL (2+ or greater).

Antigen Bands

• Must display a clearly distinguishable band at the locations of recombinant antigens FP10, FP6, and TcF, with color intensities for each recombinant antigen greater than the L-CAL (2+ or greater).
• Must display a clearly distinguishable band at the location of recombinant antigen FP3 band with color intensity of +/- or greater.

The strips MUST NOT display uneven background color or colored deposits across any portion of seven possible visible bands. Some uneven background color or colored deposits between the bands or at the ends of the strip are acceptable.

RESULTS

Specimen Strip Validity Criteria

If a specimen strip does not meet all of the criteria below, then the associated test result is invalid and the specimen must be retested in a new run.

• Must display clearly distinguishable onboard L-CAL and H-CAL bands.
• The L-CAL band (at the visual cutoff, 1+) must be clearly lighter (less intense) than the H-CAL band (at scoring of 3+).
• Must display a clearly distinguishable SPM-CTL band, with color intensity greater than the L-CAL (2+ or greater).

The strips MUST NOT display uneven background color or colored deposits across any portion of seven possible visible bands. Some uneven background color or colored deposits between the bands or at the ends of the strip are acceptable.
Reading Results
For valid strips, reactivity of test specimens against individual \( T \) cruzi recombinant antigens (rAgs) is visually graded by comparing the color intensity of each of the four \( T \) cruzi antigens against the color intensity of the L-CAL (visual cutoff, intensity of 1+) and the H-CAL (intensity of 3+) on the strip incubated with the test specimen. The identity and location of the rAgs coated on the strips are shown below.

Interpretation criteria are summarized in the following tables. For valid strips, the following criteria should be used to interpret the result.

<table>
<thead>
<tr>
<th>REACTION BAND INTENSITY</th>
<th>VISUAL GRADE</th>
</tr>
</thead>
<tbody>
<tr>
<td>Absent</td>
<td>-</td>
</tr>
<tr>
<td>Faint or less than the L-CAL</td>
<td>+/-</td>
</tr>
<tr>
<td>Equal to the L-CAL</td>
<td>1+</td>
</tr>
<tr>
<td>Greater than the L-CAL but less than the H-CAL</td>
<td>2+</td>
</tr>
<tr>
<td>Equal to the H-CAL</td>
<td>3+</td>
</tr>
<tr>
<td>Greater than the H-CAL</td>
<td>4+</td>
</tr>
</tbody>
</table>

Occasionally, a strip may have a dark background. If the L-CAL and H-CAL onboard calibrator bands are distinguishable from the background (ie, darker than the background, with the H-CAL band darker than the L-CAL), the strip is interpretable and the intensity of the antigen bands should be compared to the onboard calibrators as described in this section.

Interpretation of Results
A NEGATIVE, INDETERMINATE, or POSITIVE interpretation for a specimen is based on the reaction pattern of the rAg bands present on the strip. For valid strips, the following criteria should be used to interpret the result.

Specimens with INDETERMINATE test results must be retested once and may need to be recentrifuged. Refer to the SPECIMEN COLLECTION AND PREPARATION FOR ANALYSIS section to determine if recentrifugation is needed.

<table>
<thead>
<tr>
<th>Initial Result</th>
<th>Retest Result</th>
<th>Interpretation</th>
</tr>
</thead>
<tbody>
<tr>
<td>POSITIVE</td>
<td>No retest required</td>
<td>Antibodies to ( T ) cruzi detected, indicative of a ( T ) cruzi infection</td>
</tr>
<tr>
<td>INDETERMINATE</td>
<td>Retest is POSITIVE</td>
<td>Antibodies to ( T ) cruzi detected, indicative of a ( T ) cruzi infection</td>
</tr>
<tr>
<td>Retest is INDETERMINATE</td>
<td>Antibodies to ( T ) cruzi may or may not be present. These individuals, especially those with risk factors*, may be retreated after 6 months using a freshly drawn specimen.</td>
<td></td>
</tr>
<tr>
<td>Retest is NEGATIVE</td>
<td>NEGATIVE: Antibodies to ( T ) cruzi not detected</td>
<td></td>
</tr>
</tbody>
</table>

NOTE: A01 (on the strip above) is an example of the printed strip ID.

Visual color intensities on each strip incubated with test specimen should be graded against the L-CAL and H-CAL bands on the same strip, not against strips incubated with the negative or the positive control. The intensity of reactivity against \( T \) cruzi rAgs is visually graded against the intensities of the onboard L-CAL and H-CAL as follows:

Interpretation criteria are summarized in the following tables.

Antigen Band Pattern

- No antigen bands visible or a SINGLE antigen band, having an intensity of 1+ or greater

- Two or more bands, all having a +/+ intensity

- Two or more bands, with at least 1 band having an intensity of 1+ or greater

- Two or more bands, all having a +/+ intensity

- Two or more bands, all having a +/+ intensity

SPECIFICATIONS

- A test result that is negative does not exclude the possibility of exposure or infection with \( T \) cruzi.

LIMITATIONS OF THE PROCEDURE

- This assay was designed and validated for use with individual human serum and plasma specimens. This assay has not been validated for use with pooled specimens.

- Do not use specimens collected in heparin.

- Serum from heparinized patients may be incompletely coagulated. Draw the specimen prior to heparin therapy or after heparin therapy is discontinued and activated partial thromboplastin time (aPTT) levels return within normal range.

- False-reactive test results can be expected with any test kit. False-reactive test results have been observed due to nonspecific interactions. Refer to the SPECIFIC PERFORMANCE CHARACTERISTICS section of this package insert for assay performance characteristics.

- Some specimens that have undergone multiple freeze/thaw cycles or have been stored frozen for prolonged periods may result in erroneous or inconsistent test results.

- All specimens must be centrifuged according to the Preparation for Analysis section of this package insert prior to running the assay.

- Performance has not been established using plasmapheresis or cadaveric specimens, umbilical cord blood, or body fluids such as urine, saliva, semen, amniotic fluid, cerebrospinal fluid, or pleural fluid. These specimens should not be tested using ABBOTT ESA Chagas.

- Additional testing for Leishmania should be considered for individuals with indeterminate results with ABBOTT ESA Chagas who have identifiable risk factors for leishmaniasis.

- Do not use heat-inactivated specimens.

- Do not use specimens with obvious microbial contamination or gross lipemia.

- Do not use serum or plasma specimens with obvious gross hemolysis (dark red to black). No qualitative performance differences were observed when specimens were spiked with 500 mg/dL of hemoglobin.

- The ABBOTT ESA Chagas Procedure and RESULTS must be closely followed when testing serum or plasma specimens for the presence of antibodies to \( T \) cruzi.

- Avoid microbial contamination of reagents by carefully following handling precautions within this package insert.

- A test result that is negative does not exclude the possibility of exposure to or infection with \( T \) cruzi.

* Identifiable risk factors include: born in a Chagas endemic area or parent born in an endemic area, etc.

PREPARATION FOR ANALYSIS

- No retest required

- Retest is NEGATIVE

- Retest is INDETERMINATE

- Retest is POSITIVE

- No test kit available

- A freshly drawn specimen

- Refer to the SPECIFIC PERFORMANCE CHARACTERISTICS section of this package insert for assay performance characteristics.
**SPECIFIC PERFORMANCE CHARACTERISTICS**

**Clinical Reproducibility**

Reproducibility was determined at the clinical testing sites with ABBOTT ESA Chagas by testing a 6-member panel. Panel Members 1 and 2 were T cruzi antibody-positive specimens. Panel Member 3 was a T cruzi antibody-negative specimen. Each panel member was tested once per day over 3 days with each of 3 reagent lots at 4 clinical sites, with 1 technician at each site.

There were 36 strips tested for each of the 6 panel members with 4 antigen bands per strip for a total of 864 antigen band readings. 99.19% (857/864) of the antigen band readings were within one level of intensity of the band reading reported by the majority of strip readers. 0.81% (7/864) of the antigen band readings were within one level of intensity higher than the band reading reported by the majority of strip readers. 0.81% (7/864) of the antigen band readings were within one level of intensity higher than the band reading reported by the majority of strip readers.

Table I shows the percent agreement for the strip interpretation for each of the panel members. Eight (8) out of 36 strips for the indeterminate Panel Member 3 were interpreted as negative.

This study shows acceptable reproducibility performance for ABBOTT ESA Chagas.

<table>
<thead>
<tr>
<th>Panel Member</th>
<th>Expected Reactivity</th>
<th>ABBOTT ESA Chagas</th>
<th>% Agreement</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 Positive</td>
<td>36 0 0</td>
<td>36 0</td>
<td>100.00</td>
</tr>
<tr>
<td>2 Negative</td>
<td>36 0 0</td>
<td>36 0</td>
<td>100.00</td>
</tr>
<tr>
<td>3 Indeterminate</td>
<td>36 0 28</td>
<td>8 77.78</td>
<td>100.00</td>
</tr>
<tr>
<td>4 Positive</td>
<td>36 0 0</td>
<td>0 0</td>
<td>100.00</td>
</tr>
<tr>
<td>5 Positive</td>
<td>36 0 0</td>
<td>0 0</td>
<td>100.00</td>
</tr>
<tr>
<td>6 Positive</td>
<td>36 0 0</td>
<td>0 0</td>
<td>100.00</td>
</tr>
</tbody>
</table>

POG = Positive, IND = Indeterminate, NEG = Negative

**Clinical Specificity in US Blood Donors**

A total of 330 serum and plasma specimens from United States (US) blood donors were tested with ABBOTT ESA Chagas (Table II). The specimens were presumed negative for antibodies to T cruzi based on a T cruzi antibody licensed enzyme-linked immunosorbent assay (T cruzi antibody licensed ELISA). In this study, 327 out of 330 specimens (99.1%, with a 95% confidence interval of 97.4% to 99.8%) were negative by ABBOTT ESA Chagas, 3 (0.9%) were indeterminate, and none were positive. Of the 3 specimens with indeterminate ABBOTT ESA Chagas results, 2 were negative with a laboratory-developed T cruzi radioimmuno precipitation assay (T cruzi RIPA), not licensed by the FDA, and the other specimen was not tested by T cruzi RIPA. There were no false positives with ABBOTT ESA Chagas.

In an ABBOTT ESA Chagas post-marketing study, an additional 300 serum and plasma specimens from United States (US) blood donors were tested with ABBOTT ESA Chagas (Table II). The specimens were presumed negative for antibodies to T cruzi based on ABBOTT PRISM Chagas. In this study, 297 out of 300 specimens (99.0%, with a 95% confidence interval of 97.1% to 99.8%) were negative by ABBOTT ESA Chagas, 3 (1.0%) were indeterminate, and none were positive. Of the 3 specimens with indeterminate ABBOTT ESA Chagas results, all 3 were negative with T cruzi RIPA. There were no false positives with ABBOTT ESA Chagas.

**Specificity with Specimens from Individuals with Medical Conditions or Containing Potentially Interfering Substances**

Specificity of ABBOTT ESA Chagas was evaluated internally at Abbott by testing 618 frozen serum and plasma specimens collected from individuals with medical conditions unrelated to T cruzi infection or containing potentially interfering substances (Table III). Specimens that were positive or indeterminate with ABBOTT ESA Chagas were tested further with T cruzi RIPA (shown in brackets).

**Specificity with Specimens from Individuals with Medical Conditions Unrelated to T cruzi Infection**

<table>
<thead>
<tr>
<th>Category</th>
<th>Total</th>
<th>POS</th>
<th>IND</th>
<th>NEG</th>
</tr>
</thead>
<tbody>
<tr>
<td>Leishmaniasis</td>
<td>59</td>
<td>1 [0-0-0-0]</td>
<td>24 [2-0-2-0]</td>
<td>33 [0-0-0-33]</td>
</tr>
<tr>
<td>Malaria Positive</td>
<td>32</td>
<td>0 [0-0-0-0]</td>
<td>3 [0-0-0-3]</td>
<td>29 [0-0-0-29]</td>
</tr>
<tr>
<td>Syphilis Serologic Positive</td>
<td>16</td>
<td>0 [0-0-0-0]</td>
<td>1 [0-0-0-1]</td>
<td>15 [0-0-0-15]</td>
</tr>
</tbody>
</table>

**Other Medical Conditions Unrelated to T cruzi Infection and Specimens Containing Potentially Interfering Substances**

<table>
<thead>
<tr>
<th>Substances</th>
<th>Total</th>
<th>POS</th>
<th>IND</th>
<th>NEG</th>
</tr>
</thead>
<tbody>
<tr>
<td>Malaria Positive</td>
<td>512</td>
<td>0 [0-0-0-0]</td>
<td>17 [3-1-3-0]</td>
<td>495 [0-0-0-495]</td>
</tr>
</tbody>
</table>

**Table I ABBOTT ESA Chagas Reproducibility**

**Table II Specificity of ABBOTT ESA Chagas with Specimens from US Blood Donors**

**Table III Specificity of ABBOTT ESA Chagas with Specimens from Individuals with Medical Conditions Unrelated to T cruzi Infection**
Supplemental Testing of US Blood Donor Specimens Repeatedly Reactive by ABBOTT PRISM Chagas and/or the T cruzi Antibody Licensed ELISA

A total of 329 US blood donor specimens were tested with ABBOTT ESA Chagas and T cruzi Antigen ELISA. Of these specimens, 202 were preselected donor specimens repeatedly reactive by T cruzi antibody licensed ELISA, 64 repeatedly reactive by T cruzi antibody licensed ELISA and/or ABBOTT PRISM Chagas were prospectively identified by testing 41,760 fresh donor specimens and 63 were PRISM Chagas repeatedly reactive identified during the ABBOTT ESA Chagas post-marketing study.

A comparison of ABBOTT ESA Chagas results and T cruzi RIPA results for the 329 US blood donors that were repeatedly reactive on a T cruzi antibody licensed ELISA and/or ABBOTT PRISM Chagas is shown in Table IV. Of the 329 repeatedly reactive specimens, 151 were positive on both ABBOTT ESA Chagas and T cruzi RIPA, 5 were positive on ABBOTT ESA Chagas and indeterminate on T cruzi RIPA and 24 were positive on ABBOTT ESA Chagas and negative on T cruzi RIPA.

This study shows a high level of concordance of positives (151/153 or 98.7%) on ABBOTT ESA Chagas with specimens positive by T cruzi RIPA, demonstrating consistency with T cruzi RIPA for samples repeatedly reactive on ABBOTT PRISM Chagas and/or the T cruzi antibody licensed ELISA.

The most probable T cruzi antibody status for specimens with discordant results between ABBOTT ESA Chagas and T cruzi RIPA in these studies was interpreted based on the available evidence for those specimens, recognizing that the laboratory-developed assay, T cruzi RIPA, is not an absolute standard for determination of T cruzi antibody status. Specimens that were repeatedly reactive on the two licensed screening assays and positive with 3 or more antigen bands with ABBOTT ESA Chagas were interpreted as true positive. Specimens that were repeatedly reactive on one screening test and positive with 3 or less antigen bands with ABBOTT ESA Chagas were interpreted as an inconclusive status. Risk factors for T cruzi infection such as immigration from a Chagas-endemic country and intensity of the bands on ABBOTT ESA Chagas were also considered in the interpretation of the status as true positive and as inconclusive. Specimens that were negative on T cruzi RIPA and indeterminate with ABBOTT ESA Chagas with 2 or fewer reactive bands were interpreted as true negative.

A negative result on a follow-up sample for both ABBOTT ESA Chagas and T cruzi RIPA were used to reinterpret the status of the donor as true negative.

### Table IV

<table>
<thead>
<tr>
<th>T cruzi RIPA</th>
<th>Total</th>
<th>T cruzi Antibody Status</th>
</tr>
</thead>
<tbody>
<tr>
<td>ABBOTT ESA Chagas</td>
<td>POS</td>
<td>IND</td>
</tr>
<tr>
<td>POS</td>
<td>180</td>
<td>151</td>
</tr>
<tr>
<td>IND</td>
<td>14</td>
<td>1</td>
</tr>
<tr>
<td>NEG</td>
<td>136</td>
<td>1 TN</td>
</tr>
<tr>
<td>Total</td>
<td>329</td>
<td>153</td>
</tr>
</tbody>
</table>

*PO = Positive, IND = Indeterminate, NEG = Negative, Tp = True Positive, TN = True Negative*

- All 5 of these specimens were repeatedly reactive with both ABBOTT PRISM Chagas and a T cruzi antibody licensed ELISA, 4 specimens were positive with 3 or more antigen bands with ABBOTT ESA Chagas and were from donors from Chagas endemic countries, and 1 specimen was positive with 2 antigen bands with ABBOTT ESA Chagas.
- 19 specimens were repeatedly reactive with both ABBOTT PRISM Chagas and a T cruzi antibody licensed ELISA and positive for 3 or more antigen bands with ABBOTT ESA Chagas.
- One specimen was repeatedly reactive with both ABBOTT PRISM Chagas and a T cruzi antibody licensed ELISA and positive for 2 antigen bands with ABBOTT ESA Chagas.
- One specimen was repeatedly reactive with both ABBOTT PRISM Chagas and a T cruzi antibody licensed ELISA and positive for 2 antigen bands with ABBOTT ESA Chagas.
- One specimen was repeatedly reactive with both ABBOTT PRISM Chagas and a T cruzi antibody licensed ELISA and positive for 2 antigen bands with ABBOTT ESA Chagas.
- One specimen was nonreactive with one of the licensed screening assays and a follow-up specimen was nonreactive with both licensed screening assays and indeterminate with ABBOTT ESA Chagas.

A total of 284 US blood donor specimens that were repeatedly reactive by ABBOTT PRISM Chagas were tested with ABBOTT ESA Chagas and T cruzi RIPA. Of the 284 blood donor specimens, 58 were prospectively identified by testing 41,760 fresh donor specimens, 163 were identified by testing 202 preselected donor specimens that were repeatedly reactive by a T cruzi antibody licensed ELISA and 63 were PRISM Chagas repeatedly reactive identified during the ABBOTT ESA Chagas post-marketing study.

A comparison of ABBOTT ESA Chagas results and T cruzi RIPA results for the 284 ABBOTT PRISM Chagas repeatedly reactive US blood donor specimens is shown in Table V. Of the 284 repeatedly reactive specimens, 153 were positive by T cruzi RIPA, of which 151 were ABBOTT ESA Chagas positive, one specimen was ABBOTT ESA Chagas negative and one specimen was ABBOTT ESA Chagas indeterminate. Follow-up testing on a new specimen from the donor that was ABBOTT ESA Chagas positive was nonreactive by both screening assays, and negative by both ABBOTT ESA Chagas and T cruzi RIPA, indicating that this donor was most likely not infected with T cruzi, and the negative result with ABBOTT ESA Chagas for the initial specimen was correct. There was no follow-up for the ABBOTT ESA Chagas indeterminate specimen.

The combined studies showed a high level of concordance of positive results (151/153 or 98.7%) from ABBOTT ESA Chagas with T cruzi RIPA positive results, demonstrating consistency with T cruzi RIPA for samples repeatedly reactive with ABBOTT PRISM Chagas.

### Table V

<table>
<thead>
<tr>
<th>T cruzi RIPA</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>ABBOTT ESA Chagas</td>
<td>POS</td>
</tr>
<tr>
<td>POS</td>
<td>151</td>
</tr>
<tr>
<td>IND</td>
<td>1*</td>
</tr>
<tr>
<td>NEG</td>
<td>0</td>
</tr>
<tr>
<td>Total</td>
<td>153</td>
</tr>
</tbody>
</table>

*PO = Positive, IND = Indeterminate, NEG = Negative*

- The follow-up testing results for a new specimen from the same donor were nonreactive by ABBOTT PRISM Chagas (S/CO = 0.95, 0.91, 0.82), nonreactive by T cruzi antibody licensed ELISA (S/CO = 0.094), negative by ABBOTT ESA Chagas (FP10 -, FP6 +/-, FP3 - and TcF -), and negative by T cruzi RIPA.
- One specimen was nonreactive with T cruzi antibody licensed ELISA. There was no follow-up specimen provided.

Of the 41,760 blood donors screened with ABBOTT PRISM Chagas, 58 were repeatedly reactive of which 9 were positive with ABBOTT ESA Chagas, 2 specimens were repeatedly reactive with both screening assays and positive with 3 antigen bands with ABBOTT ESA Chagas. The follow-up specimen for the first donor was positive with 3 antigen bands with ABBOTT ESA Chagas, T cruzi RIPA indeterminate, ABBOTT PRISM Chagas nonreactive (0.97 S/CO), nonreactive on ELISA (0.924 S/CO), and indicated several risk factors for T cruzi infection. No follow-up specimen was available for the second donor. The third specimen was repeatedly reactive with ABBOTT PRISM Chagas and positive with 2 antigen bands with ABBOTT ESA Chagas. A follow-up specimen tested indeterminate with ABBOTT ESA Chagas, negative by T cruzi RIPA, and nonreactive by the two licensed screening assays.

This study shows concordance (6 out of 6) for ABBOTT ESA Chagas positive results with T cruzi RIPA positive results for prospectively acquired US blood donor specimens found repeatedly reactive by ABBOTT PRISM Chagas.
SUPPLEMENTAL TESTING OF ABBOTT PRISM CHAGAS REPEATEDLY REACTIVE SPECIMENS

**Table I**

<table>
<thead>
<tr>
<th>Category</th>
<th>Number of Specimens Tested</th>
<th>ABBOTT PRISM Chagas Repeatedly Reactive/Number of Specimens Tested</th>
<th>ABBOTT ESA Chagas Positive/Number of Specimens Tested</th>
</tr>
</thead>
<tbody>
<tr>
<td>US Blood Donors</td>
<td>41,760</td>
<td>58/41,760 (0.14%)</td>
<td>9/58 (15.52%)</td>
</tr>
</tbody>
</table>

Of the 63 specimens that were identified during the ABBOTT ESA Chagas post-marketing study as PRISM Chagas repeatedly reactive, 7 were positive with ABBOTT ESA Chagas. Of these 7 specimens, 6 were positive with both ABBOTT ESA Chagas and T cruzi RIPA and 1 was ABBOTT ESA Chagas positive with 2 antigen bands and T cruzi RIPA negative. The specimen that was positive with ABBOTT ESA Chagas and T cruzi RIPA negative was repeatedly reactive with ABBOTT PRISM Chagas (1.26, 1.29, and 1.19 S/CO) and nonreactive on ELISA (0.361 S/CO). A follow-up specimen tested positive with 2 antigen bands on ABBOTT ESA Chagas, negative by T cruzi RIPA, and nonreactive by the two licensed screening assays.

The post-marketing study shows concordance (6 out of 6) for ABBOTT ESA Chagas positive results with T cruzi RIPA positive results for prospectively acquired US blood donor specimens found repeatedly reactive by ABBOTT PRISM Chagas.

The combined studies show concordance of (12 out of 12) for ABBOTT ESA Chagas positive results with T cruzi RIPA positive results for prospectively acquired US blood donor specimens found repeatedly reactive by ABBOTT PRISM Chagas.

**Supplemental Testing of US Blood Donor Specimens Repeatedly Reactive by a T cruzi Antibody Licensed ELISA**

A total of 221 US blood donor specimens that were repeatedly reactive by a T cruzi antibody licensed ELISA were tested with ABBOTT ESA Chagas and with T cruzi RIPA. Of the 221 blood donor specimens, 13 were from testing 16,292 fresh donor specimens, 202 were from testing preselected donor specimens repeatedly reactive by a T cruzi antibody licensed ELISA and 6 were identified during the ABBOTT ESA Chagas post-marketing study.

A comparison of ABBOTT ESA Chagas results with T cruzi RIPA results for the 221 T cruzi antibody licensed ELISA repeatedly reactive US blood donors is shown in Table VII. Of the 221 specimens, 149 were positive by both T cruzi RIPA and ABBOTT ESA Chagas. This study shows a high level of concordance of positive results (149/149 or 100.0%) on ABBOTT ESA Chagas with specimens positive by T cruzi RIPA, demonstrating consistency with T cruzi RIPA for samples repeatedly reactive on the T cruzi antibody licensed ELISA.

**Table II**

<table>
<thead>
<tr>
<th>Category</th>
<th>Number of Specimens Tested</th>
<th>ABBOTT PRISM Chagas Repeatedly Reactive/Number of Specimens Tested</th>
<th>ABBOTT ESA Chagas Positive/Number of Specimens Tested</th>
</tr>
</thead>
<tbody>
<tr>
<td>US Blood Donors</td>
<td>16,292</td>
<td>13/16,292 (0.08%)</td>
<td>7/13 (53.85%)</td>
</tr>
</tbody>
</table>

Of the 6 specimens that were identified during the ABBOTT ESA Chagas post-marketing study as T cruzi antibody licensed ELISA repeatedly reactive, 5 were positive with both ABBOTT ESA Chagas and T cruzi RIPA and 1 was negative with both ABBOTT ESA Chagas and T cruzi RIPA. A follow-up specimen tested negative with both ABBOTT ESA Chagas and T cruzi RIPA, and repeatedly reactive by the two licensed screening assays.

The post-marketing study shows concordance (5 out of 5) for ABBOTT ESA Chagas positive results with T cruzi RIPA positive results for prospectively acquired US blood donor specimens found repeatedly reactive by T cruzi antibody licensed ELISA.

**Clinical Sensitivity in Parasitologically Positive Non-US Specimens**

A total of 110 serum specimens from individuals known to be positive for the T cruzi parasite were tested with ABBOTT PRISM Chagas and ABBOTT ESA Chagas (Table IX). Of the 110 specimens, 65 were from individuals that tested positive by identification of the parasite with xenodiagnosis. The remaining 45 specimens were from individuals known to be positive for the T cruzi parasite by historical identification of the parasite with xenodiagnosis or hemoculture. The specimens were obtained from the Chagas-endemic countries of Argentina, Bolivia, Brazil, and Peru. All 110 specimens were repeatedly reactive on ABBOTT PRISM Chagas and were positive for T cruzi antibodies with ABBOTT ESA Chagas demonstrating 100% sensitivity (Table IX).

**Clinical Sensitivity in Serologically Positive Non-US Specimens**

A total of 85 serum specimens from individuals reactive for T cruzi antibodies based on 2 different serologic tests for antibodies to T cruzi (ie, ELISA, immunofluorescence assay [IFA], or indirect hemagglutination assay [IHA]), were obtained from Argentina and were tested with ABBOTT PRISM Chagas and with ABBOTT ESA Chagas. All 85 specimens were repeatedly reactive on ABBOTT PRISM Chagas and were positive for T cruzi antibodies with T cruzi RIPA and with ABBOTT ESA Chagas demonstrating 100% sensitivity (Table X).

**Prospective Studies in High Risk Populations**

A total of 524 serum specimens from individuals residing in Chagas-endemic areas were tested with ABBOTT ESA Chagas, ABBOTT PRISM Chagas, and T cruzi antibody licensed ELISA. Specimens were obtained from Argentina, Brazil, Guatemala, Panama, and Peru. Specimens that were positive and indeterminate with ABBOTT ESA Chagas and/or repeatedly reactive with either ABBOTT PRISM Chagas or a T cruzi antibody licensed ELISA, if available, were tested further with T cruzi/RIPA. A comparison of ABBOTT ESA Chagas results and T cruzi RIPA results for the 524 high risk specimens is shown in Table XI. This study shows a high level of concordance of positives (130/132 or 98.5%) on ABBOTT ESA Chagas.
Chagas with specimens positive by T. cruzi/RIPA, demonstrating consistency with T. cruzi/RIPA for samples repeatedly reactive on ABBOTT PRISM Chagas and/or the T. cruzi antibody licensed ELISA.

The most probable T. cruzi antibody status for specimens in these studies was interpreted based on the available evidence for those specimens, recognizing that the laboratory-developed assay, T. cruzi RIPA, is not an absolute standard for determination of T. cruzi antibody status. Specimens that were repeatedly reactive on the two licensed screening assays and positive with 3 or more antigen bands with ABBOTT ESA Chagas were interpreted as true positive. Specimens that were repeatedly reactive on one screening test and positive with 3 or less antigen bands with ABBOTT ESA Chagas were interpreted as an inconclusive status. Risk factors for T. cruzi infection such as from a Chagas-endemic country and intensity of the bands on ABBOTT ESA Chagas were also considered in the interpretation of the status as true positive and as inconclusive.

### Table XI

<table>
<thead>
<tr>
<th>T. cruzi Antibody Status</th>
<th>ABBOTT ESA Chagas</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Positive (POS)</td>
<td>147</td>
<td>3 TP*</td>
</tr>
<tr>
<td>Indeterminate (IND)</td>
<td>21</td>
<td>1 INC</td>
</tr>
<tr>
<td>Negative (NEG)</td>
<td>356</td>
<td>6 TNE</td>
</tr>
<tr>
<td>Total</td>
<td>524</td>
<td>381</td>
</tr>
</tbody>
</table>

Note: T. cruzi RIPA results are in the bracket: [POS-IND-NEG]

* TP = True Positive, TN = True Negative
  * POS = Positive, IND = Indeterminate, NEG = Negative
  * Out of 21 specimens that tested indeterminate with ABBOTT ESA Chagas, 7 specimens were not available for testing by T. cruzi RIPA.
  * T. cruzi RIPA testing was not performed for 354 nonreactive specimens.

### BIBLIOGRAPHY


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