Anti PR3 (c-ANCA) ELISA Assay Kit
Quantitative determination of IgG class antibodies against proteinase 3 (PR3) in human serum or plasma.

INTENDED USE
Eagle Biosciences Anti PR3 (c-ANCA) ELISA Assay Kit is an indirect solid phase enzyme immunoassay (ELISA) for the quantitative measurement of IgG class autoantibodies against proteinase 3 (PR3) in human serum or plasma. The assay is intended for research use only and is not intended for diagnostic procedures.

1. CLINICAL SIGNIFICANCE
Anti-neutrophilic-cytoplasm antibodies (ANCA) represents a group of autoantibodies directed towards the cytoplasmatic components of the neutrophil granulocytes and monocytes. The classical methods for the determination of ANCA are the immunofluorecent methods. With these indirect immunofluorescence techniques two main patterns are recognized, a cytoplasmatic (c-ANCA) and a perinuclear (p-ANCA) type.

2. PRINCIPLE
Anti PR3 (c-ANCA) ELISA Assay Kit is a test is based on the binding of the antibodies in the sample with human neutrophil proteinase 3 coated into the microplates. In the first step the antibodies in calibrators, controls or prediluted patient samples bind to the inner surface of the wells. After a 60 minutes incubation the microplate is washed with wash buffer for removing non-reactive serum components. In the second step an anti-human-IgG horseradish peroxidase conjugated solution recognizes the IgG class antibodies bound to the immobilized antigens. After a 30 minutes incubation any excess enzyme conjugate which is not specifically bound is washed away with the wash buffer. Then a chromogenic substrate solution containing TMB is dispensed into the wells. After 15 minutes incubation the color development is stopped by adding the Stop Solution. The solution color change into yellow. The amount of color is directly proportional to the concentration of IgG antibodies present in the original sample. The concentration of IgG antibodies in the sample is calculated through a calibration curve.

3. REAGENTS, MATERIALS AND INSTRUMENTATION

3.1. Reagents and materials supplied in the kit
1. Anti-PR3 Calibrators (5 vials, 1.2 mL each)
   - Phosphate buffer 0,1M, NaN₃ < 0,1%, human serum
   - CAL0
   - CAL1
   - CAL2
   - CAL3
   - CAL4
   
2. Controls (2 vials, 1.2 mL each)
   - Phosphate buffer 0,1M, NaN₃ < 0,1%, human serum
   - Negative control
   - Positive control
   
3. Sample Diluent (1 vial, 100 mL)
   - Phosphate buffer 0,1 M NaNa₃ < 0,1%
   
4. Conjugate (1 vial, 15 mL)
   - Anti h-IgG conjugated with horseradish peroxidase (HRP), BSA 0,1%, Proclin < 0,0015%
   
5. Coated Microplate
   (1 breakable microplate coated with human neutrophil proteinase 3)
   
6. TMB Substrate (1 vial, 15 mL)
   - 3,3',5,5'-tetramethylbenzidine 0.26 g/L, hydrogen peroxide 0.05%
   
7. Stop Solution (1 vial, 15 mL)
   - Sulphuric acid 0.15M
   
8. 10X Conc. Wash Solution (1 vial, 50 mL)
   - Phosphate buffer 0,2M, proclin < 0,0015%

9. IVD
See external label

10. LOT
See external label

11. 2°C ± 8°C

12. 0°C ± 3°C

13. 2°C ± 8°C

14. 7. Stop Solution

15. 6. TMB Substrate

16. 5. Coated Microplate

17. 4. Conjugate

18. 3. Sample Diluent

19. 2. Controls

20. 1. Anti-PR3 Calibrators
3.2. Reagents necessary not supplied
Distilled water.

3.3. Auxiliary materials and instrumentation
Automatic dispenser.
Microplate reader (450 nm).

4. WARNINGS
- This Anti PR3 (c-ANCA) ELISA Assay Kit is intended for research use by professional persons only. Not for internal or external use in Humans or Animals.
- Use appropriate personal protective equipment while working with the reagents provided.
- Follow Good Laboratory Practice (GLP) for handling blood products.
- All human source material used in the preparation of the reagents has been tested and found negative for antibody to HIV 1&2, HbsAg, and HCV. No test method however can offer complete assurance that HIV, HBV, HCV or other infectious agents are absent. Therefore, Calibrators and Controls should be handled in the same manner as potentially infectious material.
- Material of animal origin used in the preparation of the kit has been obtained from animals certified as healthy and the bovine protein has been obtained from countries not infected by BSE, but these materials should be handled as potentially infectious.
- Some Anti PR3 (c-ANCA) ELISA Assay Kit reagents contain small amounts of Sodium Azide (NaN₃) or Proclin 300® as preservatives. Avoid the contact with skin or mucosa.
- Sodium Azide may be toxic if ingested or absorbed through the skin or eyes; moreover it may react with lead or copper plumbing to form potentially explosive metal azides. If you use a sink to remove the reagents, allow scroll through large amounts of water to prevent azide build-up.
- The TMB Substrate contains an irritant, which may be harmful if inhaled, ingested or absorbed through the skin. To prevent injury, avoid inhalation, ingestion or contact with skin and eyes.
- The Stop Solution consists of a diluted sulphuric acid solution. Sulphuric acid is poisonous and corrosive and can be toxic if ingested. To prevent chemical burns, avoid contact with skin and eyes.
- Avoid the exposure of reagent TMB/H₂O₂ to directed sunlight, metals or oxidants. Do not freeze the solution.

5. PRECAUTIONS
- Please adhere strictly to the sequence of pipetting steps provided in this protocol. The performance data represented here were obtained using specific reagents listed in this Instruction For Use.
- All Anti PR3 (c-ANCA) ELISA Assay Kit reagents should be stored refrigerated at 2-8°C in their original container. Any exceptions are clearly indicated. The reagents are stable until the expiry date when stored and handled as indicated.
- Allow all Anti PR3 (c-ANCA) ELISA Assay Kit components and specimens to reach room temperature (22-28°C) and mix well prior to use.
- Do not interchange Anti PR3 (c-ANCA) ELISA Assay Kit components from different lots. The expiry date printed on box and vials labels must be observed. Do not use any kit component beyond their expiry date.
- **WARNING:** the conjugate reagent is designed to ensure maximum dose sensitivity and may be contaminated by external agents if not used properly: therefore, it is recommended to use disposable consumables (tips, bottles, trays, etc.). For divided doses, take the exact amount of conjugate needed and do not re-introduce any waste product into the original bottle. In addition, for doses dispensed with the aid of automatic and semi-automatic devices, before using the conjugate, it is advisable to clean the fluid handling system, ensuring that the procedures of washing, deproteinization and decontamination are effective in avoiding contamination of the conjugate; this procedure is highly recommended when the kit is processed using analyzers which are not equipped with disposable tips.
- For this purpose, Diametra supplies a separate decontamination reagent for cleaning needles.
- If you use automated equipment, the user has the responsibility to make sure that the kit has been appropriately tested.
- The incomplete or inaccurate liquid removal from the wells could influence the assay precision and/or increase the background.
- It is important that the time of reaction in each well is held constant for reproducible results. Pipetting of samples should not extend beyond ten minutes to avoid assay drift. If more than 10 minutes are needed, follow the same order of dispensation. If more than one plate is used, it is recommended to repeat the dose response curve in each plate.
- Addition of the TMB Substrate solution initiates a kinetic reaction, which is terminated by the addition of the Stop Solution. Therefore, the TMB Substrate and the Stop Solution should be added in the same sequence to eliminate any time deviation during the reaction.
- Observe the guidelines for performing quality control in medical laboratories by assaying controls and/or pooled sera.
- Maximum precision is required for reconstitution and dispensation of the reagents.
- Samples microbiologically contaminated, highly lipemic or haemolysed should not be used in the assay.
- Plate readers measure vertically. Do not touch the bottom of the wells.

6. PROCEDURE

6.1. Preparation of the Calibrators (C₀…C₄)
Since no international reference preparation for Anti-proteinase 3 antibodies is available, the assay system is calibrated in relative arbitrary units. The Calibrators are ready to use and have the following concentration:

<table>
<thead>
<tr>
<th>AU/mL</th>
<th>C₀</th>
<th>C₁</th>
<th>C₂</th>
<th>C₃</th>
<th>C₄</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0</td>
<td>10</td>
<td>20</td>
<td>40</td>
<td>160</td>
</tr>
</tbody>
</table>

Once opened, the Calibrators are stable 6 months at 2-8°C.

6.2. Preparation of the Sample
Either human serum or plasma samples can be used for the test execution. Test samples should be clear. Contamination by lipemia is best avoided, but does not interfere with this assay. Specimens may be refrigerated at 2-8°C for up to five days or stored at -20°C up to six months. Avoid repetitive freezing and thawing of serum or plasma samples. This may result in variable loss of autoantibody activity. Testing of heat-inactivated serum or plasma samples is not recommended.

All serum or plasma samples must be prediluted 1:100 with sample diluent; for example 10 µL of sample should be diluted with 990 µL of sample diluent. The Controls are ready to use.
6.3. Preparation of the Wash Solution

Dilute the content of each vial of the "10X Conc. Wash Solution" with distilled water to a final volume of 500 mL prior to use. For smaller volumes respect the 1:10 dilution ratio. The diluted wash solution is stable for 30 days at 2-8°C.

In concentrated wash solution is possible to observe the presence of crystals; in this case mix at room temperature until the complete dissolution of crystals; for greater accuracy, dilute the whole bottle of concentrated wash solution to 500 mL, taking care to transfer completely the crystals, then mix until crystals are completely dissolved.

6.4. Procedure

- Allow all reagents to reach room temperature (22-28°C) for at least 30 minutes.
- Unused coated microwell strips should be released securely in the foil pouch containing desiccant and stored at 2-8°C.
- To avoid potential microbial and/or chemical contamination, unused reagents should never be transferred into the original vials.
- As it is necessary to perform the determination in duplicate in order to improve accuracy of the test results, prepare two wells for each point of the calibration curve (C0-C4), two for each Control, two for each sample, one for Blank.

<table>
<thead>
<tr>
<th>Reagent</th>
<th>Calibrator</th>
<th>Sample/ Controls</th>
<th>Blank</th>
</tr>
</thead>
<tbody>
<tr>
<td>Calibrator</td>
<td>C0-C4</td>
<td>100 µL</td>
<td></td>
</tr>
<tr>
<td>Controls</td>
<td></td>
<td>100 µL</td>
<td></td>
</tr>
<tr>
<td>Diluted</td>
<td>Sample</td>
<td>100 µL</td>
<td></td>
</tr>
</tbody>
</table>

Incubate 60 minutes at room temperature (22-28°C). Remove the content from each well, wash the wells 3 times with 300 µL of diluted wash solution.

Conjugate 100 µL 100 µL

Incubate 30 minutes at room temperature (22-28°C). Remove the contents from each well, wash the wells 3 times with 300 µL of diluted wash solution.

TMB Substrate 100 µL 100 µL 100 µL

Incubate 15 minutes in the dark at room temperature (22-28°C).

Stop Solution 100 µL 100 µL 100 µL

Shake the microplate gently. Read the absorbance (E) at 450 nm against Blank within 5 minutes.

7. QUALITY CONTROL

- The PR3 IgG Positive and the Negative Control should be run with every batch of samples to ensure that all reagents and procedures perform properly.
- Because Positive and the Negative Control are prediluted, they do not control for procedural methods associated with dilution of specimens.
- Additional suitable control sera may be prepared by aliquoting pooled human serum specimens and storing at < -20°C.
- In order for the test results to be considered valid, all of the criteria listed below must be met. If any of these are not met, the test should be considered invalid and the assay repeated:

8. RESULTS

For the Anti PR3 (c-ANCA) test a 4-Parameter-Fit with lin-log coordinates for optical density and concentration is the data reduction method of choice. Smoothed-Spline Approximation and log-log coordinates are also suitable. We recommend using a Lin-Log curve. First calculate the averaged optical densities for each calibrator well. Use lin-log graph paper and plot the averaged optical density of each calibrator versus the concentration. Draw the best fitting curve approximating the path of all calibrator points. The calibrator points may also be connected with straight line segments. The concentration of unknowns may then be estimated from the calibration curve by interpolation.

9. REFERENCE VALUES

In a normal range study with serum samples from healthy blood donors the following ranges have been established with the Anti PR3 (c-ANCA) test:

<table>
<thead>
<tr>
<th>Cut-off</th>
<th>Anti-PR3 (c-ANCA)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>20 AU/mL</td>
</tr>
</tbody>
</table>

Please pay attention to the fact that the determination of a range of expected values for a "normal" population in a given method is dependent on many factors, such as specificity and sensitivity of the method used and type of population under investigation. Therefore each laboratory should consider the range given by the Manufacturer as a general indication and produce their own range of expected values based on the indigenous population where the laboratory works. Positive results should be verified concerning the entire clinical status of the patient. Also every decision for therapy should be taken individually. It is recommended that each laboratory establishes its own normal and pathological ranges of serum anti PR3 antibodies.

10. LIMITATIONS OF PROCEDURE

The presence of immune complexes or other immunoglobulin aggregates in the patient sample may cause an increased level of non-specific binding and produce false positives in this assay.
11. PERFORMANCE AND CHARACTERISTICS

11.1. Precision and reproducibility
Precision and reproducibility are evaluated by eight reply of two positive samples by two different runs with two different lots. Dispensing and washing operations were performed manually by an operator. The results in terms of Calibrator deviation and coefficient of variation were below:

<table>
<thead>
<tr>
<th>Sample</th>
<th>1</th>
<th>2</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>SD</td>
<td>CV%</td>
</tr>
<tr>
<td>Intra-test</td>
<td>0.16</td>
<td>4.2</td>
</tr>
<tr>
<td>Inter-test</td>
<td>0.35</td>
<td>7.5</td>
</tr>
</tbody>
</table>

11.2. Specificity
Comparison test against a commercial reference kit, performed on 32 sera (3 of them positive sera and 29 negative sera) showed a 100% specificity.

11.3. Sensitivity
Comparison test against a commercial reference kit, performed on 32 sera (3 of them positive sera and 29 negative sera) showed a 100% sensibility.

11.4. Detection Limit
The lowest concentration of anti PR3 antibodies that can be distinguished from the Calibrator 0 is about 0.13 AU/mL with a confidence limit of 98%.

12. WASTE MANAGEMENT
Reagents must be disposed off in accordance with local regulations.

BIBLIOGRAPHY

ERROR POSSIBLE CAUSES / SUGGESTIONS

No colorimetric reaction
- no conjugate pipetted reaction after addition
- contamination of conjugates and/or of substrate
- errors in performing the assay procedure (e.g. accidental pipetting of reagents in a wrong sequence or from the wrong vial, etc.)

Too low reaction (too low ODs)
- incorrect conjugate (e.g. not from original kit)
- incubation time too short, incubation temperature too low

Too high reaction (too high ODs)
- incorrect conjugate (e.g. not from original kit)
- incubation time too long, incubation temperature too high
- water quality for wash buffer insufficient (low grade of deionization)
- insufficient washing (conjugates not properly removed)

Unexplainable outliers
- contamination of pipettes, tips or containers
  insufficient washing (conjugates not properly removed) too high within-run
- reagents and/or strips not pre-warmed to CV% Room Temperature prior to use
- plate washer is not washing correctly (suggestion: clean washer head)
  too high between-run - incubation conditions not constant (time, CV % temperature)
- controls and samples not dispensed at the same time (with the same intervals) (check pipetting order)
- person-related variation