INTRODUCTION
SAA, a protein of ~12 kDa, is a positive acute phase reactant that circulates in blood mostly associated with high density lipoprotein (HDL). In pigs, basal serum levels are approximately 5 µg/ml. Levels can reach 500 µg/ml during infection.2,3

PRINCIPLE OF THE ASSAY
The assay uses two different peptide-specific pig SAA antibodies; one for solid phase immobilization and the other, conjugated to horseradish peroxidase (HRP), for detection. Serum samples are first denatured by heating for 1 hour at 60°C. The denaturing step dissociates SAA from interfering factors. Subsequently, the denatured samples are diluted. Standards and diluted samples are incubated, in the microtiter wells, together with HRP conjugate for one hour. This results in SAA molecules being sandwiched between immobilization and detection antibodies. The wells are then washed to remove unbound HRP-conjugate. TMB is added and incubated for 20 minutes. If SAA is present a blue color develops. Color development is stopped by addition of Stop solution, changing the color to yellow, and absorbance is measured at 450 nm. The concentration of SAA is proportional to absorbance and is derived from a standard curve.

MATERIALS AND COMPONENTS
Materials provided with the kit:
- SAA antibody coated 96-well plate (12 x 8-well strips)
- HRP Conjugate, 11 ml
- SAA stock (lyophilized)
- 20x Wash solution; TBS50-20, 50 ml
- Diluent; YD50-1, 50 ml
- TMB, TMB11-1, 11 ml
- Stop solution, SS11-1, 11 ml

Materials required but not provided:
- Pipettors and tips
- Distilled or deionized water
- Polypropylene or glass tubes
- Vortex mixer
- Water bath
- Absorbent paper or paper towels
- Plate incubator/shaker
- Plate washer
- Plate reader capable of measuring absorbance at 450 nm.
- Curve fitting software

STORAGE
The SAA stock should be stored at or below -20°C for optimum stability. The remainder of the kit should be stored at 2-8°C and the microtiter plate should be kept in a sealed bag with desiccant. Kits will remain stable for six months from the date of purchase.

GENERAL INSTRUCTIONS
1. All reagents should be allowed to reach room temperature before use.
2. Reliable and reproducible results will be obtained when the assay is carried out with a complete understanding of the instructions and with adherence to good laboratory practice.
3. The wash procedure is critical. Insufficient washing will result in poor precision and falsely elevated absorbance readings.
4. Laboratory temperature will influence absorbance readings. Our ELISA kits are calibrated using shaking incubators set at 150 rpm and 25°C. Performance of the assay at lower temperatures will result in lower absorbance values.

WASH SOLUTION PREPARATION
The wash solution is provided as a 20x stock. Prior to use, dilute the contents of the bottle (50 ml) with 950 ml of distilled or deionized water.

STANDARD PREPARATION
1. Reconstitute the SAA stock as described on the vial label. Mix gently several times before use. The stock does not require heat treatment
2. Label 7 polypropylene tubes as 100, 50, 25, 12.5, 6.25, 3.13 and 1.56 ng/ml.
3. Into the tube labeled 100 ng/ml, pipette the volume of diluted SAA stock detailed on the SAA stock vial label. Then add the indicated volume of SAA and mix gently. This provides the 100 ng/ml standard.
4. Dispense 250 µl of diluted SAA standard into the tubes labeled 50, 25, 12.5, 6.25, 3.13 and 1.56 ng/ml.
5. Pipette 250 µl of the 100 ng/ml SAA standard into the tube labeled 50 ng/ml and mix. This provides the 50 ng/ml standard.
6. Similarly prepare the remaining standards by two-fold serial dilution.

SAMPLE PREPARATION
Denaturation
1. Dispense 100 µl of each serum sample into a polypropylene microcentrifuge tube and tightly seal.
2. Incubate the samples at 60°C in a water bath for one hour.

Dilution
1. Dilute the denatured samples at least 400-fold with the diluent provided with the kit. Optimum dilutions must be empirically determined.

ASSAY PROCEDURE
1. Secure the desired number of 8-well strips in the holder. Unused strips should be stored in the re-sealed bag with desiccant at 2-8°C for future use.
2. Dispense 100 µl of standards and samples into the wells (we recommend that standards and samples be run in duplicate).
3. Add 100 µl of HRP-conjugate into each well.
4. Incubate on a plate shaker at 150 rpm and 25°C for one hour.
5. Empty and wash the microtiter wells 5x with 1x wash solution using a plate washer (400 µl/well).
6. Strike the wells sharply onto absorbent paper or paper towels to remove all residual droplets.
7. Dispense 100 µl of TMB into each well.
8. Incubate on an orbital micro-plate shaker at 150 rpm at 25°C for 20 minutes.
9. After 20-minutes, stop the reaction by adding 100 µl of Stop solution to each well.
10. Gently mix. It is important to make sure that all the blue color changes to yellow.
11. Read absorbance at 450 nm with a plate reader within 5 minutes.

**CALCULATION OF RESULTS**
1. Using curve fitting software, construct a standard curve by plotting absorbance values of the standards versus log10 of the concentration.
2. Fit the standard curve to a four-parameter logistic regression (4PL) equation (x axis = log10 concentration) and determine the concentration of the samples from the standard curve (remember to derive the antilog).
3. Multiply the derived concentration by the dilution factor to determine the actual concentration in the serum or plasma sample.
4. If the A450 values of samples fall outside the standard curve, samples should be diluted appropriately and re-tested.

**TYPICAL STANDARD CURVE**
A typical standard curve with absorbance at 450 nm on the Y-axis against SAA concentrations on the X-axis is shown below. This curve is for illustration only.

<table>
<thead>
<tr>
<th>SAA (ng/ml)</th>
<th>A450</th>
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<tbody>
<tr>
<td>100</td>
<td>2.451</td>
</tr>
<tr>
<td>50</td>
<td>1.867</td>
</tr>
<tr>
<td>25</td>
<td>1.224</td>
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<tr>
<td>12.5</td>
<td>0.877</td>
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<tr>
<td>6.25</td>
<td>0.576</td>
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<td>3.125</td>
<td>0.416</td>
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<tr>
<td>1.563</td>
<td>0.305</td>
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**ASSAY CHARACTERISTICS**
Typical results obtained with three normal Yorkshire pig serum samples are shown in the following table. Samples were heated at 60°C for one hour then diluted as indicated and concentrations (ng/ml) derived. Serum concentrations (SC, µg/ml) were calculated by multiplying the concentrations of the diluted samples by their respective dilution factors. Average values, standard deviation (SD) and coefficient of variation (CV) were then calculated.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Dilution</th>
<th>A450</th>
<th>ng/ml</th>
<th>SC (µg/ml)</th>
<th>Average</th>
<th>SD (µg/ml)</th>
<th>CV</th>
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<tbody>
<tr>
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<td>1.29</td>
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**REFERENCES**

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For technical assistance please email us at techsupport@lifediagnostics.com