Complement C3c FS*

Diagnostic reagent for quantitative in vitro determination of complement component C3c in serum or plasma on photometric systems

Order Information
Cat. No. Kit size
1 1802 99 10 930 R1 4 x 20 mL + R2 1 x 8 mL
1 1802 99 10 935 R1 2 x 20 mL + R2 1 x 8 mL
5 9200 99 10 037 3 x 1 mL TruCal Protein high
5 9200 99 10 038 5 x 1 mL TruCal Protein:
              Calibrator set with 5 different levels

Summary [1,2]
The complement system represents a group of at least 20 plasma proteins and several receptor proteins that interact in a regulated proteolytic cascade in order to destroy invading bacteria and prevent deposition of immunocomplexes. The activation results in decreased concentrations of C3 and/or C4 due to consumption of the intact proteins. The complement cascade can be activated by two different pathways. The classical pathway is activated by immunocomplexes or antibodies bound to bacteria or virus. The cascade starts with the binding of the C1q part of C1 to the Fc-part of the antibodies and it activates C3 by proteolysis of C4. The alternative pathway is activated independent of antibodies by microorganisms, Polysaccharides, autolysis of C3 or aggregated immunoglobulins. The alternative pathway does not need C4 protein. Because C3 is common to both pathways, lowered concentrations are indicating a general complement activation. Lowered C3 values are found in inflammatory and infectious diseases especially in glomerulonephritis and SLE (Systemic Lupus erythematoses). Depending on the activated pathway C4 values can be lowered or stay normal. Isolated low values of C4 can occur in hereditary and acquired angioneuritic oedema. Hereditary deficiency states of both complement factors have been reported. C3 as well as C4 react as acute phase proteins. This increase due to an inflammatory process may mask a moderately increased complement consumption.

Method
Immunoturbidimetric test

Principle
Determination of the concentration of C3c by photometric measurement of antigen-antibody-reaction of antibodies to human C3c with C3c present in the sample.

Reagents
Components and Concentrations
R1: TRIS pH 7.5 100 mmol/L
      NaCl 320 mmol/L
R2: TRIS pH 8.0 100 mmol/L
      NaCl 300 mmol/L
Anti-human C3c antibody (goat) < 1%

Storage Instructions and Reagent Stability
The reagents are stable up to the end of the indicated month of expiry, if stored at 2 – 8°C and contamination is avoided. Do not freeze the reagents and protect reagents from light!

Warnings and Precautions
2. The reagents contain sodium azide (0.95 g/L) as preservative. Do not swallow! Avoid contact with skin and mucous membranes!
3. Reagent 2 contains animal material. Handle the product as potentially infectious according to universal precautions and good clinical laboratory practices.
4. In very rare cases, samples of patients with gammapathy might give falsified results [6].
5. Please refer to the safety data sheets and take the necessary precautions for the use of laboratory reagents. For diagnostic purposes, the results should always be assessed with the patient’s medical history, clinical examinations and other findings.
6. For professional use only!

Waste Management
Please refer to local legal requirements.

Reagent Preparation
The reagents are ready to use.

Materials required but not provided
NaCl solution 9 g/L
General laboratory equipment

Specimen [3]
Serum, heparin plasma or EDTA plasma

During storage of serum, the C3 and C4 proteins slowly degrade into C3c resp. C4c fragments (fragmentation is inhibited by EDTA). These fragments still contain the reactive epitopes and may even display higher signals than the intact protein. Depending on the conditions of this aging process, fresh serum samples may show up to 30% lower C3 values than samples stored at 4 °C for 8 days. The fragmentation of C4 is much slower than for C3 and only 15% lower values can be observed under similar storage conditions. Discard contaminated specimens.

Assay Procedure for Analyzers

Application sheets for automated systems are available on request.

| Wavelength  | 340 nm |
| Optical path | 1 cm |
| Temperature  | 37°C |
| Measurement  | Against reagent blank |

<table>
<thead>
<tr>
<th>Blank</th>
<th>Sample or calibrator</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sample or calibrator</td>
<td>-</td>
</tr>
<tr>
<td>Dist. water</td>
<td>5 μL</td>
</tr>
<tr>
<td>Reagent 1</td>
<td>350 μL</td>
</tr>
<tr>
<td>Mix, incubate for 3 – 5 min., read absorbance (A1), then add:</td>
<td></td>
</tr>
<tr>
<td>Reagent 2</td>
<td>70 μL</td>
</tr>
<tr>
<td>Mix, incubate for 5 min., read absorbance (A2).</td>
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ΔA = (A2 – A1) sample or calibrator

* fluid stable
Calculation
The concentration of C3c in unknown samples is derived from a calibration curve using an appropriate mathematical model such as logit/Log or spline. The calibration curve is obtained with 5 calibrators at different levels and NaCl solution (9 g/L) for determination of the zero value.
Stability of calibration: 4 weeks

Calibrators and Controls
For the calibration of automated photometric systems, DiaSys TruCal Protein calibrator set or the TruCal Protein high calibrator is recommended. The assigned values of the calibrators have been made traceable to the ERM®-DA470k/IFCC Reference Material. DiaSys TruLab Protein control should be assayed for internal quality control. Each laboratory should establish corrective action in case of deviations in control recovery.

Performance Characteristics

Measuring Range
The test has been developed to determine concentrations of C3c within a measuring range from 1 – 500 mg/dL (0.01 – 5.0 g/L) – depending on the highest calibrator. When values exceed this range samples should be diluted 1 + 1 with NaCl solution (9 g/L) and the result multiplied by 2.

Prozone Limit
No prozone effect was observed up to an C3c value of 1000 mg/dL (10.0 g/L).

Specificity/Interferences
Due to its antibodies, DiaSys Complement C3c is a specific immunoassay for human C3c. No interference was observed by conjugated and unconjugated bilirubin up to 60 mg/dL, hemoglobin up to 1000 mg/dL, lipemia up to 2000 mg/dL triglycerides and RF up to 1200 IU/mL. Interference by monoclonal gammopathies was not observed up to 6400 mg/dL IgA. 4100 mg/dL IgM and 6400 mg/dL IgG. For further information on interfering substances refer to Young DS [5].

Sensitivity/Limit of Detection
The lower limit of detection is 1 mg/dL (0.01 g/L).

Imprecision

<table>
<thead>
<tr>
<th>Intra-assay precision</th>
<th>Mean [mg/dL]</th>
<th>SD [mg/dL]</th>
<th>CV [%]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sample 1</td>
<td>143</td>
<td>3.00</td>
<td>2.10</td>
</tr>
<tr>
<td>Sample 2</td>
<td>188</td>
<td>5.84</td>
<td>3.10</td>
</tr>
<tr>
<td>Sample 3</td>
<td>220</td>
<td>7.73</td>
<td>3.51</td>
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<tbody>
<tr>
<td>Sample 1</td>
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<td>5.70</td>
<td>4.00</td>
</tr>
<tr>
<td>Sample 2</td>
<td>186</td>
<td>6.58</td>
<td>3.54</td>
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<tr>
<td>Sample 3</td>
<td>219</td>
<td>6.93</td>
<td>3.16</td>
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</table>

Total precision according to protocol EP-5 of the NCCLS (National Committee of Clinical Laboratory Standards):

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</thead>
<tbody>
<tr>
<td>Sample 1</td>
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<td>4.23</td>
<td>2.97</td>
</tr>
<tr>
<td>Sample 2</td>
<td>185</td>
<td>9.18</td>
<td>4.95</td>
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Method Comparison
A comparison of DiaSys Complement C3c FS (y) to an immunoturbidimetric test (x) using 107 samples gave following results:
\[
y = 0.83 x + 6.72 \text{ mg/dL}; r = 0.982
\]
A comparison of DiaSys Complement C3c FS (y) to a nephelometric test (x) using 95 samples gave following results:
\[
y = 1.11 x - 7.02 \text{ mg/dL}; r = 0.961
\]

Reference Range [4]
90 – 180 mg/dL (0.9 – 1.8 g/L)
In case of fresh samples lower reference ranges are expected for C3c. Each laboratory should establish own reference ranges in order to reflect its specific working conditions.

Literature

Manufacturer
DiaSys Diagnostic Systems GmbH
Alte Strasse 9
65558 Holzheim, Germany