Introduction

Many disorders of the central nervous system (CNS) are accompanied by increased protein concentration in the cerebrospinal fluid (CSF), which can be attributable to an altered permeability of the blood-brain barrier, to intrathecal synthesis of immunoglobulins, or to the combination of both (1). Examination of CSF total protein and specific proteins is used chiefly to detect increased permeability of the blood-brain barrier to plasma proteins or to detect increased intrathecal production of immunoglobulins. The degree of permeability of the blood-brain barrier can be evaluated by immunochemical measurements of albumin in CSF and in serum specimens obtained at the same time. Albumin is a particularly suitable protein because it is neither synthesized nor metabolized intrathecally. Demonstration of increased intrathecal immunoglobulin synthesis resulting from infiltration of IgG-producing lymphocytes inside brain and spinal cord is a specific indication for the presence of inflammation in the CNS. Albumin can cross the blood-brain barrier, but in contrast to IgG, cannot be produced in the CNS itself (2).

Several disorders of the CNS such as bacterial meningitis, multiple sclerosis and other CNS inflammatory diseases, are associated with an increase in protein concentration in CSF (3, 4). In this context, the examination of the CSF includes quantitative measurements of immunoglobulin and albumin concentrations in comparison with the serum. The purpose of this work was to establish the degree of blood brain barrier damage in inflammatory and noninflam-
inflammatory diseases, intrathecal synthesis of IgG and to use the data in differential diagnosis between the two types of CNS diseases.

Materials and Methods

The investigations were performed on patients with inflammatory and noninflammatory disease of CNS. Patients with inflammatory disease (n = 20) had clinically confirmed diagnosis of tuberculosis meningitis. In the group with noninflammatory disease (n = 20) were patients with diagnosis of poliradicu-

loneuritis, vascular disorders and brain carcinoma, while tuberculosis meningitis was excluded. Patients who had normal concentrations of total proteins, albumin and globulins in both serum and CSF were used as the control.

CSF and blood were obtained by lumbar puncture (L5-L6) and venipuncture, respectively. Blood samples were drawn without anticoagulant and centrifuged (3000 g for 10 min at 4 °C), and serum was harvested. All samples were stored at –20 °C before being processed. Albumin and IgG were measured in serum and unconcentrated CSF by immunoturbidimetry using Turbitimer (Dade-Behring) with calibrators and internal controls provided by Dade-Behring and according to manufacturer’s recommendations.

As a mirror of blood-CSF barrier status, the CSF/serum albumin ratio was calculated; its threshold of positivity varied between 6 × 10⁻³ and 8 × 10⁻³ with the age of patients. To detect intrathecal synthesis of IgG (ISI), we calculated the IgG/albumin index (5) and the Schuller index (6). The IgG/albumin index was calculated by the following formula:

\[ \text{IgG/Albumin index} = \frac{\text{IgG}_{\text{CSF}}/\text{IgG}_{\text{serum}}}{\text{albumin}_{\text{CSF}}/\text{albumin}_{\text{serum}}} \]

and was positive above 0.65. The Schuller index was defined by the following formula:

\[ \text{ISI (mg/L)} = \text{IgG}_{\text{CSF}} - (30 + [(\text{albumin}_{\text{CSF}} - 240)/60] \times \text{IgG}_{\text{serum}}) \]

Intrathecal synthesis of IgG is suspected when the result is > 0 mg/L.

Results and Discussion

The data we have obtained for CSF albumin and immunoglobulin G are presented in Table I.

The CSF albumin concentration was significantly increased both in patients with inflammatory and noninflammatory disease, which was already confirmed in the literature (7). Significant difference was found between each patient group and the control one (p < 0.001), but not between the patient groups themself (p > 0.05).

The concentrations of IgG in CSF were increased in both groups of patients, and significant differences was found in comparison to the control (p < 0.001), which is in agreement with literature data (8). The differences in IgG concentration were also found between patients with inflammatory and noninflammatory CNS disease (p < 0.05).

For the evaluation of the blood-brain barrier damage, we have calculated the albumin index. The results are presented in Table II.

Compared to the control group there was a significant increment in patients both with noninflammatory (p < 0.001) and inflammatory disease (p < 0.001). No significant differences were found for albumin index between the two patient groups (p > 0.05).

The preferred method for the measurement of blood-CSF barrier dysfunction is the analysis of albumin quotient and its evaluation with regard to the agerelated reference range (9). The analysis of total protein is still frequently used for evaluation of barrier dysfunction because it does offer some clinical value in spite of its broad reference range, i.e., lower sensitivity, and specificity than albumin quotient. Due to systematic inflammatory diseases with high total protein concentrations in blood, high total concentrations in CSF are subsreantlrty observed. This could lead to false positive interpretation but could be avoided by using the albumin quotient (10).

The immunoglobulin index was calculated in order to prove possible intrathecal synthesis of IgG. In patients with inflammatory CNS disease, the index was increased, which was not the case with noninflammatory diseases (Table III).

Table I   Concentrations of CSF albumin and immunoglobulin G in the control group (I) and patients with inflammatory (II) and noninflammatory CSN disease

<table>
<thead>
<tr>
<th>N</th>
<th>I (x ± SD)</th>
<th>II (x ± SD)</th>
<th>III (x ± SD)</th>
<th>p (I – II)</th>
<th>p (I – III)</th>
<th>p (II – III)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Albumin (mg/L)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>20</td>
<td>206.6 ± 45.2</td>
<td>1314 ± 114.7</td>
<td>1478 ± 142.4</td>
<td>&lt; 0.001</td>
<td>&lt; 0.001</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td>IgG (mg/L)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>20</td>
<td>26.09 ± 3.12</td>
<td>334 ± 30.3</td>
<td>152 ± 12.5</td>
<td>&lt; 0.001</td>
<td>&lt; 0.001</td>
<td>&lt;0.05</td>
</tr>
</tbody>
</table>
This finding confirms the existence of IgG synthesis as a response to the infectious agent (11). By comparison of IgG index between the investigated groups, statistically significant differences were found \( p < 0.001 \).

The determination of CSF/serum quotients is regarded as a suitable evaluation and interpretation method for the detection of intrathecal IgG, IgA or IgM synthesis. As reviewed recently, the frequently used IgG Index or IgG Synthesis rate give up to 90% false-positive values for high albumin quotients (blood-CSF barrier dysfunction) compared to the gold standard, which is the detection of oligoclonal bands on isoelectric focusing (IEF). Regarding quantitative analyses of intrathecal immunoglobulin synthesis, there is an agreement to refer to hyperbolic discrimination line, expressed in either a numerical or graphical format (12, 13). This is often facilitated by use of PC-based software for evaluation of CSF data profiles, and thus knowledge-based interpretation programs support routine CSF analysis (12, 14).

<table>
<thead>
<tr>
<th>Table II</th>
<th>Blood-CSF barrier permeability abnormalities assessed by the albumin ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>Groups</td>
<td>CSF/albumin ratio</td>
</tr>
<tr>
<td>Control (I)</td>
<td>4.75</td>
</tr>
<tr>
<td>Inflammatory diseases (II)</td>
<td>36.40</td>
</tr>
<tr>
<td>Noninflammatory diseases (III)</td>
<td>35.10</td>
</tr>
<tr>
<td>Inflammatory/ noninflammatory diseases</td>
<td>0.48</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Table III</th>
<th>Intrathecal IgG synthesis assessed by the IgG/albumin index and the Schuller index</th>
</tr>
</thead>
<tbody>
<tr>
<td>Groups (( n = 20 ))</td>
<td>IgG/albumin index &gt; 0.65, n (%)</td>
</tr>
<tr>
<td>Control (I)</td>
<td>0.48</td>
</tr>
<tr>
<td>Inflammatory diseases (II)</td>
<td>0.77 (12)</td>
</tr>
<tr>
<td>Noninflammatory diseases (III)</td>
<td>0.39 (12)</td>
</tr>
<tr>
<td>Inflammatory/ noninflammatory diseases</td>
<td>1.974 (4)</td>
</tr>
</tbody>
</table>
DIJAGNOSTIČKI ZNAČAJ ODREĐIVANJA PROTEINA U CEREBROSPINALNOJ TEČNOSTI KOD DIFERENCIJALNE DIJAGNOSTIKE OBOLJENJA CNS-a

Dragana Vukosavljević1, Ljiljana Bumbaširević2, Aleksandra Stefanović3, Marina Stojanov3

1Institut za medicinsku biohemiju, Klinički centar Srbije, Beograd, Srbija i Crna Gora
2Institut za neurologiju, Klinički centar Srbije, Beograd, Srbija i Crna Gora
3Institut za medicinsku biohemiju, Farmaceutski fakultet Univerziteta u Beogradu, Beograd, Srbija i Crna Gora

Kratak sadržaj: Diferencijalna dijagnoza inflamatornih i neinflamatornih oboljenja CNS-a, obuhvata kvantitativno određivanje koncentracije imunoglobulina i albumina, kao i njihovo poređenje sa serumom. Koncentracije albumina i imunoglobulina G određene su u cerebrospinalnoj tečnosti (CST) u serumu pacijenata sa inflamatornim i neinflamatornim oboljenjima CNS-a. Ovi parametri određeni su i u kontrolnoj grupi, sačinjenoj od pacijenata kod kojih su ispitivani parametri bili u granicama referentnih vrednosti. Dobijeni rezultati statistički su obrađeni neparametarskim Mann-Whitney testom. Izračunate vrednosti CSF/albumin pokazale su statistički značajne razlike između obe grupe pacijenata i kontrolne grupe (p < 0,001). Ovo dokazuje da je kod obe grupe pacijenata krvno moždano barjera oštećena. Za procenu intratekalne sinteze IgG izračunati su Ig/albumin indeks i Schuller-ov index. Poređenjem vrednosti za pacijente sa nezaplaljenim oboljenjima i kontrolne grupe nisu nađene značajne razlike (p > 0,05). Suprotno, statistički značajne razlike uočene su između grupe sa inflamatornim oboljenjima i kontrolne grupe (p < 0,05), što ukazuje na postojanje intratekalne sinteze IgG-a. Vrednosti izračunatih indeksa za zapaljenske procese bile su značajno više u odnosu na nezaplaljena oboljenja (p < 0,05), što se može iskoristiti za diferencijalnu dijagnozu ovih oboljenja.

Ključne reči: cerebrospinalna tečnost, albumin, imunoglobulin G, odnos CSF/albumin, IgG/albumin indeks

References


Received: December 9, 2004
Accepted: December 16, 2004