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Cerebrospinal fluid to serum glucose ratio in non-hypoglycorrachic neurological conditions

Objective. To explore the relevance of cerebrospinal fluid to serum glucose ratio in non-hypoglycorrachic conditions.

Design. Retrospective observational study.

Setting. Neurology ward, university teaching hospital, Hong Kong.

Patients. Adult patients with conditions unrelated to hypoglycorrachia who underwent lumbar puncture.

Main outcome measures. Cerebrospinal fluid and simultaneous serum glucose concentrations, and their ratio to each other.

Results. Between September 1998 and August 2003, 170 cerebrospinal fluid and serum glucose samples were collected from 138 patients. Mean cerebrospinal fluid to serum glucose ratio was 0.61 (standard deviation, 0.142; range, 0.21-1.00). With the exception of cerebrospinal fluid protein level, laboratory parameters were similar among different diseases. The glucose ratio was lower than 0.6 in 43% and lower than 0.5 in 19% of samples. Cases with a low glucose ratio appeared to have higher serum glucose concentrations (significant among groups with different glucose ratios, P<0.001). The mean glucose ratio (0.65) was also significantly higher in patients with serum glucose concentration of lower than 7.8 mmol/L compared with those with serum glucose concentration between 7.8 and 11.1 mmol/L (mean, 0.46), or higher than 11.1 mmol/L (mean, 0.46) [P<0.001]. There was a strong negative correlation between the glucose ratio and serum glucose concentration (r= –0.704, P<0.001).

Conclusion. A lowered cerebrospinal fluid to serum glucose ratio is often seen in the absence of an appropriate disorder, especially when simultaneous serum glucose concentration is elevated. This may be explained by the saturation kinetics of glucose transportation in hyperglycaemia, and the time lag for cerebrospinal fluid and glucose to equilibrate when the blood level fluctuates.
normal.1-3,5-7  Less than this level indicates pathological hypoglycorrhachia. This misleading effect can be corrected by estimating the CSF to blood glucose ratio, which derives a fairly constant value. The widely accepted normal ratio is between 0.6 and 0.8, although 0.5 has also been considered the lower limit of normal.1-3,5-7 Less than this level indicates pathological hypoglycorrhachia.

The concept of a ‘normal’ ratio had been challenged as an over-simplification of the relationship between CSF and serum glucose at different concentrations.8,9 A study of 79 men with no CSF abnormalities and the published data from about 100 normal subjects revealed that the CSF to blood glucose ratio was not a valid measure of their true relationship.9 In addition, previous reports often focused on healthy subjects or excluded those with abnormal CSF findings. This study aimed to explore the relevance of CSF to serum glucose ratio in patients with neurological disorders.

Methods

The admission and follow-up records of patients who had CSF collected under the care of the neurology ward between September 1998 and August 2003 were retrospectively reviewed. Only LP samples were studied. Patients diagnosed with conditions associated with CSF hypoglycorrhachia and patients whose CSF white cell count exceeded 20 x 10^6 /L were excluded. Simultaneous serum glucose was defined as the glucose concentration in blood taken within 1 hour of LP (before or after). Patients in whom a simultaneous serum glucose level was unavailable were also excluded. Blood samples were transported to the laboratory in standard fluoride/oxalate tubes. Glucose concentrations were measured by the hexokinase method on a Hitachi 747 analyser (Boehringer Mannheim, Mannheim, Germany). Fasting before LP was not mandatory. None of the patients had received concentrated intravenous glucose solution within 4 hours of LP, but infusion of glucose solution of 5% or lower was permitted as this would not affect the equilibrium between CSF and glucose levels.8 Diagnosis, CSF findings (protein and glucose concentrations, white cell count), simultaneous serum glucose concentrations, and CSF to serum glucose ratios were recorded for each set of samples.

All statistical analyses were performed using the Statistical Package for the Social Sciences (Windows version 12.0; SPSS Inc, Chicago [IL], US). For group comparisons, depending on homogeneity of variance, independent-samples t test, one-way analysis of variance, or Kruskal-Wallis test (H-test) was applied. The relationship between CSF to serum glucose ratio and serum glucose concentration was tested with Spearman’s rank order correlation coefficient.

Results

One hundred and seventy sets of samples from 138 patients were studied (22 patients underwent more than one LP). The mean age of patients was 51.4 years (range, 19-86 years). The final diagnoses are shown in Table 1. Follow-up records for at least 1 year after LP were available in 135 patients. The remaining three patients had died—a 70-year-old man with peripheral neuropathy died of a ruptured aortic aneurysm; another patient diagnosed to have transverse myelitis died of pneumonia; the last patient had Lambert-Eaton syn-
drome and carcinoma of the lung. No brain metastasis was found on magnetic resonance imaging at the time of LP. Cytological examinations of CSF were also negative.

The overall mean CSF to serum glucose ratio was 0.61 (standard deviation, 0.142; range, 0.21-1.00), with a mean CSF glucose concentration of 3.89 mmol/L (range, 2.1-11.8 mmol/L) and simultaneous serum glucose concentration of 6.78 mmol/L (range, 3.4-27.6 mmol/L). Mean CSF protein level was 0.85 g/L (range, 0.13-4.74 g/L) and white cell count was 2.29 x 10^6/L (range, 0-18 x 10^6/L). The CSF glucose concentration was lower than 2.5 mmol/L in two samples: one patient with transverse myelitis had an initial CSF glucose concentration of 2.1 mmol/L but was normal on repeated LP. No infection or malignancy was identified. Another patient had associated hypoglycaemia whose CSF glucose concentration was 2.3 mmol/L and serum level was 3.4 mmol/L.

There were no significant differences among the nine disease groups in CSF to serum glucose ratio, CSF glucose concentration, serum glucose concentration, or CSF white cell count. The CSF protein level appeared higher in patients with demyelinating or inflammatory conditions (H-test, P<0.05).

Table 2 shows the number of samples with different CSF to serum glucose ratios. The ratio was lower than 0.6 in 43% and lower than 0.5 in 19% of samples. Serum glucose concentration was significantly different among the five groups (H-test, P<0.001): the concentration was highest for samples with a ratio of lower than 0.4 and it decreased as the ratio increased. Mean CSF glucose for each group ranged from 3.55 to 4.51 mmol/L. Although the difference was statistically significant (H-test, P<0.05), a trend similar to that of serum glucose concentration was not found. The means of age (H-test, P=0.117), CSF protein level, and white cell count were not significantly different among the five groups.

Thirty sets of samples were from patients with diabetes mellitus: two controlled by diet alone, 16 on oral hypoglycaemic agents, and 12 on insulin. Their mean serum glucose concentration was 11.03 mmol/L (range, 5.8-27.6 mmol/L), compared with 5.88 mmol/L (range, 3.4-11.8 mmol/L) in non-diabetic patients (t test, P<0.001). The differences in CSF to serum glucose ratio and CSF glucose concentration were significant between the two groups (t test, P<0.05 and <0.001, respectively). The mean glucose ratio in samples with simultaneous serum glucose concentration of lower than 7.8 mmol/L was 0.65 (range, 0.44-1.00; n=132). In samples with serum glucose concentration of 7.8 to 11.1 mmol/L, the mean ratio was 0.46 (range, 0.21-0.74; n=25). In samples with serum glucose concentration of higher than 11.1 mmol/L, the mean ratio was also 0.46 (range, 0.32-0.75; n=13). The difference among the three mean ratios was significant (H-test, P<0.001). Respectively to the three groups of samples, the glucose ratio were lower than 0.6 in 31%, 84%,
The CSF to serum glucose ratio was plotted against the simultaneous serum glucose concentration (Fig). The ratio did not remain constant but showed a strong negative correlation ($r = -0.704, P<0.001$), which remained significant after excluding the outliers ($r = -0.665, P<0.001$).

Discussion

The CSF to serum glucose ratio is conventionally used to adjust CSF glucose concentration for blood glucose concentration at different glycaemic levels. Previous studies focused on patients without CSF abnormalities or with specific disorders.8,9 This study included patients with a spectrum of commonly encountered neurological conditions. Many patients had a ratio below the normal range (ie <0.6 or <0.5). Their diagnoses and records were reviewed carefully, and none of them had any condition associated with hypoglycorrhachia. A significant negative correlation was also found between serum glucose concentration and the CSF to serum glucose ratio.

In meningitis, CSF glucose is depressed by pathogens as well as anaerobic glycolysis in adjacent neural tissue.1 Glucose is also consumed by neutrophils although to a lesser extent; in order to minimise such effect in this study, case inclusion was arbitrarily cut-off at a CSF white cell count of 20 x 10⁶ /L. Although this did not exclude all samples with pleocytosis, patients with low glucose ratios did not have more leukocytes than those with higher ratios.

Abnormal glycolysis generates lactate.3 Measurement of CSF lactate level, though not performed routinely in this study, may distinguish a false low CSF to serum glucose ratio from pathological glucose consumption. Nevertheless, elevated lactate level is not specific for meningitis; lactate also accumulates in the CSF in the presence of many other neurological and systemic diseases, including stroke, epilepsy, multiple sclerosis, cervical spondylosis, pneumonia, heart failure, uraemia, and hepatic failure.4

Concentration of glucose in CSF is mainly regulated by the choroid plexus through facilitated transportation that exhibits saturable kinetics.10-13 Animal data reveal that the rate of glucose extraction increases steadily with rising blood glucose concentration.11,12 In cats, when blood glucose concentration is beyond 11.1 mmol/L, the extraction rate will gradually plateau until it is saturated.12 This dissociated increment was illustrated in the present study by lowering of CSF to serum glucose ratio at hyperglycaemia, although the scatterplot (Fig) failed to match a logistic regression curve expected from the saturation kinetics of glucose transportation. Conversely, a reduced glucose ratio was also present in some normoglycaemic patients. When the serum glucose level fluctuates rapidly, there is a delay of 1 to 2 hours before a steady state can be re-established in the CSF because of its slow turnover rate.8 In this situation, a low glucose ratio may reflect inadequate CSF responsiveness to an upsurge in blood glucose from procedure-related stress.14

A previous group devised a linear regression normogram for ascertaining hypoglycorrhachia at various glycaemic levels (though the model could not address the deviation from linearity at high serum concentrations when glucose transportation saturates), and calculated that a CSF to serum glucose ratio of 0.6 or 0.5 could no longer be assumed when blood

Table 2. Number and cumulative percentage of samples with different cerebrospinal fluid (CSF) to serum glucose ratios, and comparisons of serum and CSF glucose concentrations, CSF protein level, and white cell count among groups with different ratios

<table>
<thead>
<tr>
<th>CSF to serum glucose ratio</th>
<th>No. of samples, n=170</th>
<th>Cumulative percent</th>
<th>Mean values</th>
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<tbody>
<tr>
<td></td>
<td></td>
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<td>Serum glucose* (mmol/L)</td>
</tr>
<tr>
<td>&lt;0.40</td>
<td>11</td>
<td>7</td>
<td>11.06</td>
</tr>
<tr>
<td>0.40-0.49</td>
<td>22</td>
<td>19</td>
<td>10.16</td>
</tr>
<tr>
<td>0.50-0.59</td>
<td>40</td>
<td>43</td>
<td>6.77</td>
</tr>
<tr>
<td>0.60-0.69</td>
<td>52</td>
<td>74</td>
<td>5.78</td>
</tr>
<tr>
<td>≥0.70</td>
<td>45</td>
<td>100</td>
<td>5.27</td>
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* Significantly different (Kruskal-Wallis H-test, P<0.001)
† Significantly different (Kruskal-Wallis H-test, P<0.05)
‡ Not significant (One-way ANOVA, F=1.086; P=0.365)
§ Not significant (Kruskal-Wallis H-test, P=0.187)
glucose concentration exceeded 6.9 or 10.6 mmol/L, respectively. In the present study, glucose ratios were significantly lower, and more cases had ratios below 0.6 or 0.5 when random glucose was between 7.8 and 11.1 mmol/L or higher than 11.1 mmol/L, respectively. The latter is diagnostic of diabetes mellitus and the former indicates impaired glucose tolerance that is associated with stress-induced hyperglycaemia.15,16

One potential source of error in this study was the difficulty in controlling the time between blood sampling and determination of serum glucose concentration. Because of transportation and laboratory handling, some delay in separation of serum from red cells was inevitable. It is also well-known that glycolysis prior to separation cannot be completely inhibited by the standard fluoride/oxalate preservatives. Simultaneous serum glucose levels in the specimens may thus have been underestimated (ie CSF to serum glucose ratio would be overestimated). Nevertheless, if the samples were tested within a few hours, a major reduction in serum glucose concentration should not be expected. More importantly, these data reflect the actual situation in clinical practice and patient management.

**Conclusion**

A reduced CSF to serum glucose ratio was often seen in the absence of an appropriate neurological disorder. This is contrary to the conventional belief. The ratio may be unreliable even with modest elevations of simultaneous serum glucose level during LP. An isolated, clinically unexplained lowering of CSF to serum glucose ratio should therefore be interpreted with caution and not be overstated.

**Acknowledgement**

We thank Ms Eliza Chan for her helpful advice in the statistical methods.

**References**

4. Pryce JD, Gant PW, Saul KJ. Normal concentrations of lactate, glucose, and protein in cerebrospinal fluid, and the diagnostic