Effect of Sodium Valproate on Renal Cell Brush-Border Enzymes in Rats

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Summary: The effect of sodium valproate (200 mg/kg body weight) on renal cells was investigated during a 10-day trial, by determining the catalytic activities of alanine aminopeptidase, γ-glutamyltransferase and alkaline phosphatase in the membrane cell brush-border of the renal proximal tubules.

Four days after the administration of sodium valproate, a significant increase was observed in the volume of urine (9.2 ± 4.2 ml/18-h volume; control group, 3.06 ± 1.8 ml/18-h volume), and in the catalytic activities of γ-glutamyltransferase (2.69 times that measured in the control group) and alkaline phosphatase (3.02 times that measured in the control group) in urine. After prolonged treatment the urine became alkaline (pH = 10.5 on day 10), thereby excluding alanine aminopeptidase as a useful indicator of renal cell changes following the administration of sodium valproate. The activities of alkaline phosphatase and γ-glutamyltransferase in isolated vesicles of renal brush-border membrane cells were significantly increased (p < 0.05) and decreased, respectively.

On the basis of the results obtained, we believe that the determination of the catalytic activities of γ-glutamyltransferase and alkaline phosphatase in urine might prove useful for the follow-up of the renal cell state during therapy with sodium valproate.

Introduction

Sodium valproate (sodium dipropyl acetate) is an antiepileptic which has a number of side effects, such as gastrointestinal discomforts, pancreatitis, body weight gain, hyperglycaemia and hyperglycinuria, changes of coagulation, etc. (1). In quite a number of adult patients and especially in children, the administration of sodium valproate is also associated with the development of acute (2, 3) or chronic hyperammonaemia (4, 5). The genesis of hyperammonaemia has not yet been fully explained, but two mechanisms have attracted particular attention: decreased ammonium detoxification via urea synthesis in the liver (6), and increased glutamine metabolism and increased ammonium synthesis in the kidney (7—9). Cases of serious renal damage, such as interstitial nephritis (10) and proximal renal tubule dysfunction (11) have also been reported following sodium valproate therapy.

Considering the numerous metabolic effects of sodium valproate (12), especially on the metabolism of fatty acids, it may also conceivably disturb the integrity or function of the renal cell membrane. On the other hand, increased catalytic activities of some renal enzymes in urine (otherwise localized on the cell membrane of the renal proximal tubules) are known to be among the earliest indicators of kidney cell pathologic alterations (13).

Therefore, the effect of sodium valproate on rat kidney cells was followed by monitoring the catalytic activities of alanine aminopeptidase¹), alkaline phosphatase¹) and γ-glutamyltransferase¹) in urine and isolated brush-border membrane vesicles of the renal proximal tubules.

¹) Enzymes:
Alanine aminopeptidase, EC 3.4.11.2
Alkaline phosphatase, EC 3.1.3.1
γ-Glutamyltransferase, EC 2.3.2.2
The study was performed on male Fisher rats (n = 29) aged 1 month, mean body weight 100 g. The animals had free access to chow and water throughout the study. The animals were divided into four groups: the control group (n = 6) received 1 ml saline (0.154 mol/1 NaCl) daily per os. The experimental group (n = 23) received 200 mg/kg body weight sodium valproate daily per os in the same volume. Experimental animals were sacrificed by severing the cranial spine on day 1 (n = 8), day 4 (n = 7) and day 10 (n = 8).

Prior to sacrifice, the 18-h urine was collected in metabolic cages for each group of animals. Urine volume and pH were readily determined by test strips. Urine was tested for the presence of protein, glucose, haemoglobin and ketone bodies, and centrifuged at 1500 min⁻¹ for 10 min in a Tehnica LC 320 laboratory centrifuge. Dialysis (PM exclusion 12000-14000), PolyLabo, Strasbourg, France) was performed against water for 3 h to remove possible enzyme inhibitors. Catalytic activities of the enzymes were then readily determined in the dialysed urine samples.

After sacrificing animals under ether narcosis on day 1, 4 and 10, blood was withdrawn from the umbilical vein. Blood serum was separated by centrifugation at 3000 min⁻¹ for 10 min and serum glucose, haemoglobin and ketone bodies, and centrifuged at 1500 min⁻¹ for 10 min in a Tehnica LC 320 laboratory centrifuge. Dialysis (PM exclusion 12000-14000), PolyLabo, Strasbourg, France) was performed against water for 3 h to remove possible enzyme inhibitors. Catalytic activities of the enzymes were then readily determined in the dialysed urine samples.

Immediately after sacrifice, the kidneys were excised, washed in cold saline, dried with filter paper and freed from the connective tissue and capsules.

Renal cortex sections prepared with a microtome were minced, placed in 15 ml buffer (Tris-HCl, 12 mmol/1, pH = 7.4; EGTA, 5 mmol/l; mannitol, 300 mmol/l), then homogenized with an Ultra Turrax homogenizer, type 18-10 (Janke & Kunkel, GK). Brush border membrane vesicles were isolated by precipitation with Mg ions, according to Biber et al. (14), using a Beckman L5-65 ultracentrifuge. Protein concentration and catalytic activities of alanine aminopeptidase, alkaline phosphatase and γ-glutamyltransferase were determined in isolated brush-border membrane vesicles. Mean values of the factor of enzymatic activity enrichment, following brush border membrane vesicle isolation, were f = 11.0, f = 8.0 and f = 10.0 for alanine aminopeptidase, alkaline phosphatase and γ-glutamyltransferase, respectively.

The catalytic activity of alanine aminopeptidase was determined by the procedure of Haschen (15) on a Pye Unicam SP-6-550 UV/ VIS spectrophotometer, that of alkaline phosphatase by the procedure of McComb & Bowers (16), and that of γ-glutamyltransferase by the procedure of Szász et al. (17) on a Centrifichem 400 autoanalyser. Serum urea concentration was determined by the method of Sampson et al. (18) on a Centrifichem 400 autoanalyser.

The protein concentration of isolated brush-border membrane vesicle preparations was determined by the procedure of Bradford et al. (19) on a Pye Unicam SP-6-550 UV/VIS spectrophotometer.

Table 1. Urine pH and volume changes in experimental rats

<table>
<thead>
<tr>
<th>Day after first application of valproate</th>
<th>Urine pH</th>
<th>18-h urine volume (ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control group</td>
<td></td>
<td></td>
</tr>
<tr>
<td>(n = 6)</td>
<td>6.0</td>
<td>3.06 ± 1.8</td>
</tr>
<tr>
<td>(n = 8)</td>
<td>5.0</td>
<td>2.10 ± 1.1</td>
</tr>
<tr>
<td>(n = 7)</td>
<td>5.0</td>
<td>9.20 ± 4.2*</td>
</tr>
<tr>
<td>(n = 8)</td>
<td>10.5</td>
<td>1.58 ± 0.5</td>
</tr>
</tbody>
</table>

* p < 0.05
Fig. 1 Effect of sodium valproate on the catalytic activity of alanine aminopeptidase in rat urine and kidney brush border membrane vesicles. Changes in enzyme catalytic activity are presented as relative activities in comparison to control group values. Control group activities in urine (○—○) and kidney brush border membrane vesicles (●—●) were: 0.036 ± 0.01 U/18 h volume and 299.5 ± 55 U/mg protein, respectively; *p < 0.05

Fig. 2 Effect of sodium valproate on the catalytic activity of γ-glutamyltransferase in rat urine an kidney brush border membrane vesicles. Changes in enzyme catalytic activity are presented as relative activities in comparison to control group values. Control group activities in urine (■—■) and kidney brush border membrane vesicles (○—○) were: 0.685 ± 0.13 U/18 h volume and 12955 ± 2142 U/mg protein, respectively; *p < 0.05

Fig. 3 Effect of sodium valproate on the catalytic activity of alkaline phosphatase in rat urine an kidney brush border membrane vesicles. Changes in enzyme catalytic activity are presented as relative activities in comparison to control group values. Control group activities in urine (Π—α) and kidney brush border membrane vesicles (O—O) were: 0.10 ± 0.06 U/18 h volume and 2485 ± 178 U/mg protein, respectively; *p < 0.05; **p < 0.01

The activity of γ-glutamyltransferase in isolated renal brush-border membrane vesicles was decreased about 0.30-, 0.40- and 0.18-fold on days 1, 4 and 10, respectively, whereas the activity of alkaline phosphatase was significantly increased (from 1.45-fold on days 1 and 4 to 2.18-fold on day 10 of the trial; p < 0.05).

The investigated enzymes are known as surface enzymes or integral constituents of the brush-border membrane cells of the kidney proximal tubules. Their urinary excretion is considered to be a useful early indicator of kidney alterations (13).

The time-dependent and significantly elevated catalytic activities of the three enzymes in the urine of experimental animals pointed to an adverse action of the drug on the kidney cells. The catalytic activity of alanine aminopeptidase was significantly increased as early as
the first day after sodium valproate administration, which is consistent with the finding that the enzyme is readily released from the surface of the cell membrane (23). A relatively early urinary excretion of alanine aminopeptidase was also observed by Holmes et al. (24) in their study of the coordinated release of urinary lysosomal and brush-border enzymes following renovascular surgery. The authors conclude that cell components, including enzymes, are affected by transitory pathophysiological changes in the kidney. We postulate that the decreased enzyme activity in urine after 10 days of treatment was probably due to a significant change in urine pH. By urine dialysis, we eliminated the inhibitors of alanine aminopeptidase and other enzymes. However, it is known from the literature (25) that enzymatic activity can be decreased by prolonged urine retention in the bladder, as well as by urine collection over a prolonged period of time, as in our experiment (18 h). The use of sodium valproate is associated with an increased formation of ammonium ions in the body. Thus, the above data, and the fact that ammonium ions are known to inhibit alanine aminopeptidase (26), appears to invalidate this enzyme as a diagnostic indicator of adverse changes in kidney cells during the administration of this drug.

Although the incorporation of γ-glutamyltransferase into the cell membrane lipid matrix is, according to the literature, weaker than that of alkaline phosphatase (23), marked enzymuria, together with notable polyuria, was observed concurrently for both enzymes on day 4 to the drug administration. These results are consistent with the observations of Jung et al. (27), who found that an increased water intake, i.e. increased diuresis in normal subjects, resulted in an increased excretion of soluble forms of the brush-border membrane enzymes, whereas the excretion of particulate forms was independent of diuresis. Therefore, in our future studies with sodium valproate, it might prove useful to determine both forms of γ-glutamyltransferase and alkaline phosphatase, or to express the enzyme catalytic activities in relation to creatinine concentration, which is one of the recommended ways of expressing urinary enzyme catalytic activities (25). The observed normalization of urine enzyme activities after 10 days may reflect the adaptation of renal cells to the presence of the drug. These changes in the enzyme catalytic activities are consistent with data obtained in similar studies of renal cell impairment (28).

Most of these studies show an intermittent increase in urinary enzyme activities, depending on the study duration or the concentration of the potentially toxic substance.

Whereas the activity of γ-glutamyltransferase in isolated renal brush-border membrane vesicles decreased as expected, the marked increase in the activity of alkaline phosphatase was probably due to the enhanced synthesis of enzyme multiple forms in the renal tissue (intestinal-like and hepatic types) and subsequent excretion in the urine. Thus, Pfeiderer et al. (29) found an increased amount of the intestinal-like forms of alkaline phosphatase in the urine from subjects receiving potentially toxic drugs.

In our experiments with rats, 200 mg/kg body weight sodium valproate per day appeared to cause transient changes in renal cell membranes, resulting, over a period of 10 days, in increased enzymuria of the specific enzymes, polyuria, and changes in urine pH. Proteinuria, microhaematuria and histological alterations due to the use of sodium valproate have also been demonstrated in humans (11). Thus, the data presented appear to suggest that the state of renal cells should be monitored during therapy with sodium valproate. This is especially important if it is given over a long period, during which not only transitory but also more severe damage to the renal tubular cells can be expected to occur. As urinary enzymes are very early indicators of adverse renal cell changes, the present results indicate that the determination of urinary catalytic activities of γ-glutamyltransferase and alkaline phosphatase may be useful for monitoring patients for adverse effects of sodium valproate treatment.

References


