SHORT COMMUNICATIONS
A POSSIBLE ROLE OF THE KIDNEY AND URINARY BLADDER IN UREA CONSERVATION OF *BUFO VIRIDIS* UNDER HIGH SALT ACCLIMATION

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The toad *Bufo viridis* can be acclimated to high salt solutions up to 800 mosmol kg\(^{-1}\) of NaCl (Tercafs & Schoffeniels, 1962; Katz, 1973). Despite extensive studies on various aspects of adaptation to high salt in this species (Gordon, 1962; Katz, 1979; Katz, Garcia-Romeu, Masoni & Isaia, 1981) only little attention has been paid to the osmoregulatory roles of the kidneys and urinary bladder during high salt acclimation. In *Rana cancrivora*, a species which can be acclimated to salinities even higher than *B. viridis* (and which shares the property of urea accumulation) Schmidt-Nielsen & Lee (1962), found a lower urine flow under high salt and consequently reduced urea secretion, while Gordon & Tucker (1968) observed urea retention at first, but increasing loss at salinities higher than 400 mosmol kg\(^{-1}\). Ferreira & Jesus (1973), presented evidence in salt adapted *B. bufo* suggesting increased osmolarity (mainly sodium) in the urinary bladder. We therefore attempted to assess the turnover of water and urea in *B. viridis* with special emphasis on the contribution of the renal and the urinary bladder.

Toads were collected near Jerusalem and after 2–3 weeks in the laboratory, they were acclimated to NaCl solutions, as previously described (Katz, 1973). Acclimated toads spent at least 12 days in either tap water (approx. 15 mosmol kg\(^{-1}\)), 230 or 500 mosmol kg\(^{-1}\) of NaCl. The experiments were carried out from May throughout July at 22 ± 2 °C. Both sexes were used and the toads were not fed. Water uptake was determined on individually caged animals (Katz, 1974), urine was collected through the cloaca using a glass tube, and the volume was estimated from the difference in weight before and after catheterization. Surface area was calculated according to the empirical equation A (cm\(^2\)) = W\(^{6.3}\) × 6.3 (W = weight in grams). Blood was taken from the heart into heparinized tubes. Osmolality was determined from the freezing point depression with a Knauer, Berlin semimicro osmometer, chloride was titrated with Radiometer CMT 10 chloridometer and urea was assayed colorimetrically at 570 nm using Sigma reagents (Bull, No. 640) and t-test was used for statistical analyses.

Fig. 1 compares the water uptake and urine production which were determined on the same animals. Urine production was estimated by catheterization through the cloaca during 2-h periods. It shows that the toads are at steady-state and the overall water turnover can be calculated from either measurements. Since the osmotic gradient across the skin of the '230' group has decreased significantly, while the water

Key words: *Bufo viridis*, urine composition, urinary chloride, urea loss.
uptake was not different from the tap water group, it follows that its osmotic permeability has increased considerably.

Table 1 summarizes the measurements of the urine and blood composition under each experimental condition. Note that the urine was catheterized either repeatedly at short intervals (2 h) which we think should represent nearly ureteral urine, or at long intervals (8–10 days) which represents the bladder urine. A difference was found in the osmolality, urea and chloride concentrations between the ureteral and bladder urine, under the salt acclimation conditions; this is most significant in the '500' group. Quite surprisingly we found the urine volume in the undisturbed '500' toads at the long intervals, to be quite similar to the other two groups. To verify this observation we followed the weight of caged undisturbed toads under the three acclimation conditions for a number of days. The weight of the '500' acclimated toads was quite steady, nearly unchanged (Fig. 2), reflecting low water uptake as compared with the 'tap water' and '230' acclimated groups, indicating that these toads do not void urine, or do so only rarely. It seems then, that the urine which is produced at a low rate under this condition (Fig. 1), is accumulated slowly and stored in the urinary bladder. To assess the urea turnover, we analysed the solution of a container in which we kept four to five toads under each acclimation condition overnight. The results (Table 2) show that urea loss is greatly diminished under conditions of acclimation to high salt, and is far below the calculated loss on the basis of urinary urea concentration and urine production.

The net result of these experiments points to an increasing contribution of the kidney and urinary bladder to osmoregulation under high salt conditions. Similar results indicating osmolar and sodium excretion in the urinary bladder were obtained by Ferreira & Jesus (1973) in B. bufo, and by Middler, Kleeman & Edwards (1968) in B. marinus. Chew, Elliott & Wong (1972) found a decreased ability to accumulate urea in the plasma under saline conditions in cystectomized Rana cancrivora. These authors suggested that urea permeability changes which are induced by a hormone (oxytocin), contribute to the immediate increase of urea concentration in the plasma. An increased urea permeability was found in the isolated urinary bladder of B. marinus after acclimation to 0·6% NaCl (J. Handler, personal communication).
Table 1. Urine and plasma composition in toads acclimated to NaCl solutions

<table>
<thead>
<tr>
<th>Conditions of acclimation</th>
<th>Tap water (15 mosmol l(^{-1}))</th>
<th>230 mosmol l(^{-1})</th>
<th>500 mosmol l(^{-1})</th>
</tr>
</thead>
<tbody>
<tr>
<td>Osmolality (mosmol kg(^{-1}))</td>
<td>Osmolality (mosmol kg(^{-1}))</td>
<td>Urea (mmol l(^{-1}))</td>
<td>Cl (mmol l(^{-1}))</td>
</tr>
<tr>
<td>Short intervals (2 h)</td>
<td>81 ± 6</td>
<td>24±5 ± 3·0</td>
<td>28·2 ± 4·1</td>
</tr>
<tr>
<td>Long intervals (8–10 days)</td>
<td>48 ± 3</td>
<td>12·4 ± 1·7</td>
<td>18·0 ± 1·0</td>
</tr>
<tr>
<td>Difference</td>
<td>-33</td>
<td>-12·1</td>
<td>-10·2</td>
</tr>
<tr>
<td>P</td>
<td>&lt;0·1</td>
<td>&lt;0·001</td>
<td>&lt;0·005</td>
</tr>
<tr>
<td>Urine volume (ml)</td>
<td></td>
<td>3·12 ± 1·15</td>
<td></td>
</tr>
<tr>
<td>(long intervals)*</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Plasma</td>
<td>326</td>
<td>42·0</td>
<td>119·0</td>
</tr>
</tbody>
</table>

Mean ± s.e. of six toads under each condition.

* Three determinations at 6-day intervals.
Table 2. The effect of salt acclimation on urea loss from Bufo viridis

<table>
<thead>
<tr>
<th>Conditions of acclimation</th>
<th>Osmolality (mosmol kg⁻¹)</th>
<th>Urea loss (mmol 100 g⁻¹ 24 h⁻¹)</th>
<th>Urea in urine (mmol l⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tap water</td>
<td></td>
<td>1.52 ± 0.38</td>
<td>21 ± 8</td>
</tr>
<tr>
<td>500 (NaCl)</td>
<td></td>
<td>0.31 ± 0.11</td>
<td>172 ± 31</td>
</tr>
<tr>
<td>P</td>
<td></td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

Four toads were kept in a bath with 500 ml solution for 12–20 h. Mean ± s.e. of five separate observations.

is also our experience that repeated catheterization of the toads (B. viridis) interferes strongly with their ability to acclimate to high NaCl solutions.

However, the first stage in the modification of the urine has already taken place in the kidneys. U/P ratio for osmolar, chloride and urea excretion increases with the salinity of acclimation in a selective way, indicating the efficiency and specificity of the kidneys in handling the major solutes under these conditions.

In B. marinus (Maffly, Hays, Lamdin & Leaf, 1960) and B. spinulosus (Monge,
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(3) the concentration of urinary urea was rather constant in forced hydrated and in dehydrated animals, from which they concluded that the urinary bladder is important in the maintenance of body fluid volume but has no control over the concentrations of the 'internal milieu'.

We conclude from our experiments that both the kidneys and the urinary bladder of *Bufo viridis* play an important role in the control of solutes of the plasma, and which is critical to the ability of the toads to acclimate to high salinity. It would therefore be interesting to carry out a careful study of the regulation of kidney function and urinary bladder permeability under water and high salt acclimation in order to understand the cellular mechanisms involved.

REFERENCES


