BRIEF COMMUNICATION

TRANSAMINASES IN THE EPIDIDYMAL FLUID OF THE RAM*

EUGENIA ALUMOT AND H. SCHINDLER

National and University Institute of Agriculture, Rehovot, Israel

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In a study of the composition of the epididymal fluid of rams, attention was given to the fact that nitrogen compounds constitute a great part of the dry matter and that so far seventeen free amino acids have been detected in the fluid (Ch. Neumark, unpublished data). It was therefore deemed of interest to study the enzymes of their metabolism. Two principal transaminases were investigated: glutamic–aspartic transaminase (GOT) and glutamic–alanine transaminase (GPT). The presence of these enzymes has been established in human, bovine and rabbit seminal plasma (Flipse, 1960; Povoa & Villela, 1960; Gregoire, Rakoff & Ward, 1961), but no report on their presence in the epididymal fluid was found in the literature.

Table 1

<table>
<thead>
<tr>
<th>Sample No.</th>
<th>Epididymal fluid</th>
<th>Seminal plasma</th>
<th>Blood serum</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>GOT</td>
<td>GPT</td>
<td>GOT</td>
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</tr>
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<td>20</td>
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<tr>
<td>5</td>
<td>—</td>
<td>—</td>
<td>26</td>
</tr>
</tbody>
</table>

Epididymal fluid obtained from fistulated rams (A. Tadmor & H. Schindler, unpublished information) served for the enzyme determination. For comparison, the enzymatic activity was also investigated in the seminal plasma and blood serum of rams.

The procedure of Reitman & Frankel (1957) for the determination of transaminases in blood serum was adapted to the epididymal fluid as follows: samples of epididymal semen were centrifuged with equal volumes of isotonic (0.15 M) phosphate buffer at pH 7.3; the supernatant was diluted with 4 vol. of distilled water, 0.2 ml of which (corresponding to 0.02 ml of the original

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sample) were taken for the enzyme determinations. The average dry matter content of the epididymal fluid was 8.0% and nitrogen content was 0.7%.

The enzyme activity data are presented in Table 1. The results are reported in micromoles of pyruvate formed/1 ml original fluid/hr. (In the case of G0T the oxaloacetate formed was determined as pyruvate, after decarboxylation.)

It can be seen from Table 1 that the activity of G0T is about ten times greater in the epididymal fluid and seminal plasma than in the blood serum. On the other hand, almost no GPT activity was detected in the epididymal fluid and seminal plasma, with the exception of one sample. GPT was tested also in an undiluted sample of epididymal fluid with the same negative results.

As far as the results can be compared, owing to the diversity of units of activity, the data reported in this study for G0T are of the same order of magnitude as the values found by other authors. Gregoire et al. (1961) found that in normal human seminal plasma G0T activity was about twenty times greater than in blood serum, Povoa & Villela (1960) found it ten times greater, and Flipse (1960) found that the GPT activity in bovine seminal plasma was very weak as compared to the G0T activity, which was forty times greater.

The fact that G0T transaminase in the epididymal fluid exerts an activity exceeding by many times that found in the blood and that, on the other hand, GPT is found only in traces in all but one case, despite its presence in the blood, points to the existence of a transaminase mechanism in the epididymal fluid, independent of that in the blood. It appears that the epididymal fluid is one of the sources of G0T in the seminal plasma.

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REFERENCES