An enzyme-linked immunosorbent-assay test for hepatitis B surface antigen

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SUMMARY  A new commercial test for hepatitis B surface antigen based on an enzyme-linked immunosorbent-assay system was evaluated and compared with other tests already available. It was of comparable sensitivity to the radioimmunoassay test but required less expensive equipment (in terms of both outlay and upkeep) and did not use radioisotopes; the reagents were more stable.

The third phase tests for hepatitis B surface antigen (HBsAg)—passive haemagglutination (pHA) and radioimmunoassay (RIA)—are now well established. Although regarded as the most sensitive and most specific, the RIA test has its disadvantages; it is expensive, the reagents have a short shelf-life, and it requires specialised equipment and radiological protection.

Enzyme-linked immunosorbent assays were devised by Engvall and Perlmann (1971) and their wide application has been indicated by Voller et al. (1976). An assay of this type was developed by Wolters et al. (1976) for HBsAg detection. The Virus Reference Laboratory was able to assess its performance as part of an organised multicentred study (Kacaki et al., 1977b) before the reagents were available commercially.

Material and methods

TECHNIQUES
A commercial ELISA test system (Hepanostica) developed by Organon Scientific Development Group was used as described by Wolters et al. (1976). It was available in microtitre plates and the results were read visually. A positive reaction was visible as a yellow colour which was easily read by eye against a diffuse light source placed under the microtitre plate. Each plate included two HBsAg positive controls (a weak and a strong) and one HBsAg negative control. A specimen was regarded as HBsAg positive when the colour produced was stronger than that of the negative control.

At first the specificity test recommended for con-
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Table 1  Comparison of Hepanosticon and ELISA tests

<table>
<thead>
<tr>
<th></th>
<th>Total</th>
<th>HA− ELISA−</th>
<th>HA+ ELISA−</th>
<th>HA− ELISA+</th>
<th>HA+ ELISA+</th>
</tr>
</thead>
<tbody>
<tr>
<td>'High risk'</td>
<td>276</td>
<td>223 (81%)</td>
<td>16 (5.7%)</td>
<td>35 (12.6%)</td>
<td>2 (0.7%)</td>
</tr>
<tr>
<td>Routine</td>
<td>497</td>
<td>471 (94.8%)</td>
<td>22 (4.4%)</td>
<td>4 (0.8%)</td>
<td>0</td>
</tr>
<tr>
<td>Total</td>
<td>773</td>
<td>694 (89.7%)</td>
<td>38 (5%)</td>
<td>39 (5%)</td>
<td>2 (0.25%)</td>
</tr>
</tbody>
</table>

Table 2  Comparative titration of prototype HBs antigens by different methods

<table>
<thead>
<tr>
<th>Method</th>
<th>Antigen specimen</th>
<th>Subtype ad</th>
<th>Subtype ay</th>
</tr>
</thead>
<tbody>
<tr>
<td>IEOP</td>
<td>1/16</td>
<td>1/128</td>
<td></td>
</tr>
<tr>
<td>HA (Hepanosticon)</td>
<td>1/400</td>
<td>1/1600</td>
<td></td>
</tr>
<tr>
<td>HA (Hepatest)</td>
<td>1/1600</td>
<td>1/1280</td>
<td></td>
</tr>
<tr>
<td>HA (Auscell)</td>
<td>1/1600</td>
<td>1/3200</td>
<td></td>
</tr>
<tr>
<td>RIA (Austria II)</td>
<td>1/32000</td>
<td>1/160000</td>
<td></td>
</tr>
<tr>
<td>ELISA</td>
<td>1/51200</td>
<td>1/160000</td>
<td></td>
</tr>
</tbody>
</table>

ratios above 1 but only one was above 2:1—the level regarded as significant when serum is tested.

Discussion

This new enzyme-linked immunosorbent-assay test is comparable in sensitivity and specificity with RIA. Unlike RIA it does not require radiological protection and expensive equipment (although an automatic washer/aspirator is most useful). An automatic dispenser originally supplied proved to be unsatisfactory but multichannel pipettes facilitating the dispensing of conjugate, substrate, and sulphuric acid are now available. Thus the system can be automated and should prove suitable for testing large numbers of specimens. Each plate can be used for testing about 100 specimens but, if necessary, unused wells can be used in a second run provided they are sealed during the first run and used within one week. The reagents have a shelf-life of about one year (compared with the three weeks of Austria II). As with Austria II a relatively long incubation period is required, although the overnight period at room temperature can be reduced to 2 hours at 37°C. As the time taken to read the results is very much shorter than with Austria II, results can be made available on the day of testing. However, both Austria II and ELISA in their present format are not suitable for testing the single urgent specimen.

References


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doi: 10.1136/jcp.30.8.714

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