Western blotting is useful in the salivary diagnosis of Helicobacter pylori infection

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Abstract

Background—The salivary diagnosis of Helicobacter pylori infection offers attractive possibilities for the epidemiological study of infection in children. Salivary enzyme linked immunosorbent assay (ELISA) is less reliable than serum ELISA, owing to variable transudation of immunoglobulin. In addition, children are more difficult to study because of lower specific serum antibody concentrations to H pylori. The performance of salivary western blotting in comparison with serum western blotting and serum ELISA was investigated in school children.

Subjects and methods—Paired serum and saliva specimens were obtained from 665 school children aged 9–11 in 10 British towns. All saliva and serum specimens were first analysed by ELISA; subsequently, western blotting of both specimens was performed on 31 and 34 specimens, respectively, to establish the criteria for positivity for western blotting. The remaining 121 specimens were then tested blindly and saliva was compared with the serum.

Results—The sensitivity and specificity of salivary ELISA in the 665 specimens was 32 of 50 (64%) and 530 of 691 (78%), respectively, when compared with serum ELISA. The western blotting validation was performed on 28 subjects with positive serum and positive salivary ELISA, 28 saliva positives with negative serum, 16 saliva negatives with positive serum, and 50 doubly negative subjects. Compared with serum western blots, the sensitivity and specificity of salivary western blots was 35 of 47 (61%) and 68 of 75 (91%), respectively. Using serum ELISA as the gold standard, the sensitivity and specificity were 32 of 44 (73%) and 72 of 78 (92%), respectively, the specificity being significantly higher than salivary ELISA (p < 0.001).

Conclusion—Salivary western blotting for IgG is useful in the diagnosis of H pylori infection and is superior to ELISA. It also permits the identification of pathogenic strains.

Keywords: Helicobacter pylori; western blotting; enzyme linked immunosorbent assay; salivary specimens

Helicobacter pylori is a chronic bacterial infection of the human stomach that is associated with peptic ulcer and gastric cancer, as well as with possible extra-intestinal manifestations. The diagnosis of H pylori can be made on biopsies obtained at endoscopy, but these are unsuitable for epidemiological studies, which have relied on serology and the urea breath test. Salivary antibody testing is a potentially useful alternative for epidemiological studies, especially in children, because it is non-invasive and no sample preparation is required.

The specific antibody response to H pylori can be detected using a variety of methods, but ELISA and immunoblotting (western blotting) for specific IgG are the most reliable. ELISA is more convenient than western blotting, but specific anti-H pylori IgG values tend to be lower in children than adults, and most assays have reported a slightly lower sensitivity when compared with adults.

Although IgA is the predominant immunoglobulin present within saliva, poor sensitivity and specificity are obtained in comparison with IgG when assayed by ELISA. IgG gains access to saliva by transcapillary leakage, but values are approximately 1/1000 of those in serum. The low concentrations, together with variable transcapillary leakage, are the main reasons why salivary IgG determination is less reliable than serum determination. Immunoblotting is a sensitive and qualitative method for determining the presence of antibodies, and hence would be expected to overcome some of these difficulties. It also has the advantage of being able to detect antibodies to CagA and VacA, proteins that are markers of H pylori virulence.

Therefore, we investigated whether salivary western blotting was superior to salivary ELISA for the detection of IgG antibodies to H pylori compared with serum ELISA and serum western blotting. We also determined whether salivary western blotting had the ability to detect antibodies to VacA and CagA, as determined by serum western blotting.

Subjects and methods

Our report is based on the 8–11 year old examination of the “ten towns” children study. The general design of our study has been described elsewhere. Paired saliva and serum samples were available for 665 children from the last five towns screened; three with particularly low adult cardiovascular mortality (Leatherhead, Bath, and Tunbridge Wells) and two with particularly high cardiovascular mortality (Rhonda, Rochdale). ELISA antibody determinations were performed on both serum and saliva with the Launch Premier ELISA kit (Launch Premier, Longfield, Kent, UK). This has been shown previously to be 100% sensitive and 96% specific for serum in adults. Serum was tested according to the manufacturer’s instructions. Optimum performance for salivary ELISA was obtained when undiluted saliva was incubated with IgG conjugate.
Figure 1 Frequency distribution of optical densities in 665 subjects who had salivary enzyme linked immunoabsorbent assay (ELISA) performed and who had matching serum specimens. A cut off point of 0.41 optical density units was used.

Figure 2 Frequency distribution of serum enzyme linked immunoabsorbent assay (ELISA) values (n = 129).

(A2290; Sigma, Poole, Dorset, UK) at a concentration of 1/1500.

The specimens selected for the validation study had an over representation of false positive saliva by ELISA to allow a better evaluation of specificity. All the seropositive specimens from the original 665 paired samples tested on ELISA with remaining saliva were included. In total, 122 paired saliva and serum samples were studied by western blotting. They comprised 28 subjects who were ELISA seropositive and salivary IgG positive, 16 subjects who were seropositive and saliva IgG negative, 28 subjects who were seronegative but saliva IgG positive (OD > 0.41), and 50 subjects who were seronegative and saliva IgG negative. A further seven samples were unreadable owing to gross contamination or insufficient saliva.

Criteria for a positive serum western blot were first established on 18 positive and 16 negative sera measured by ELISA, and for a positive salivary western blot on saliva specimens from 13 seropositive and 18 seronegative subjects by serum ELISA. When the criteria had been established the remainder of the samples were tested blindly.

Western blotting was performed as follows: 100 µg/ml of a crude lysate of a CagA VacA positive strain of H pylori was run on a 10–15% gradient gel and then blotted on to nitrocellulose for one hour at 20 V, 300 mamps. The H pylori were grown on selective plates at 37°C in a controlled atmosphere (5% oxygen, 10% carbon dioxide) for five days and then washed and stored in phosphate buffered saline (PBS) at −70°C. The nitrocellulose was blocked overnight at 4°C in blocking solution (15 g Marvel milk in 200 ml PBS). The nitrocellulose was incubated with 5 µl serum and 1 ml saliva and control serum samples for one hour and at the dilutions stated below in TTBS (300 ml PBS/Tween, 9 g Marvel milk). The blots were washed in PBS/Tween (one litre PBS/4 ml Tween 20) three times for 10 minutes. The IgG conjugate (HRP conjugated goat antihuman IgG; catalogue number A2290; Sigma) was diluted at concentrations stated below in TTBS and incubated again for one hour. Washing was then performed four times for 15 minutes in PBS/Tween and the signal developed using the Amersham ECL (Amersham, Little Chalfont, Buckinghamshire, UK) detection system.

Conjugate, serum, and saliva concentrations were optimised using paired samples. The optimum dilution of saliva was 1/4 to minimise sample use and 1/500 for the IgG conjugate. The optimal concentrations for serum were 1/1500 and IgG conjugate concentration of 1/6000.

Some saliva specimens contained an excess of debris on visual inspection and were centrifuged for five minutes at 5000 × g before testing.

**STATISTICAL METHODS**

The sensitivity of saliva ELISA and saliva western blot test were compared by focusing on “true” positives as defined by serum ELISA and serum western blot, respectively. The sensitivity of the two tests was then compared using McNemar’s test for paired samples with continuity correction. The specificity of the two saliva tests was similarly compared by focusing on the “true” negatives defined by the gold standard.

We calculated κ statistics for comparing the agreement for saliva and serum western blot using EPI INFO version 6.

**Results**

**PERFORMANCE OF SALIVARY ELISA VERSUS SERUM ELISA**

Of the 665 sera and saliva samples tested by ELISA, 50 (8%) were seropositive (fig 1). The sensitivity and specificity of salivary ELISA against serum ELISA as gold standard was 32 of...
50 (64%) and 530 of 619 (87%), respectively, at a cut off point of 0.41 OD (optical density) units for saliva and 0.1 OD units for serum (fig 2), as suggested by the manufacturer. The cut off point of 0.41 maximised specificity without compromising sensitivity too far (fig 3).

ESTABLISHING CRITERIA FOR POSITIVE AND NEGATIVE SALIVARY WESTERN BLOT

The criteria for positivity on serum/saliva western blotting were either the recognition of the CagA (120 kDa) band alone, which has previously been shown to be specific for the presence of H pylori infection 3 or a minimum of six bands from any other region of the blot, because several bands between 30 and 60 kDa were non-specific, as has been reported previously. 3–10 Using these criteria, 17 of 18 (94%) positive sera were western blot seropositive and 16 of 16 (100%) negative sera were western blot seronegative (fig 4).

The same criteria were applied to the saliva western blot samples and 11 of 13 (84%) of the saliva specimens from seropositive subjects were western blot positive, and 18 of 18 (100%) of the saliva specimens from seronegative subjects were western blot negative (fig 5).

PERFORMANCE OF SALIVARY WESTERN BLOT COMPARED WITH SERUM WESTERN BLOT

A total of 16 samples had to be centrifuged before western blotting and of these seven still yielded uninterpretable blots and were excluded from the analysis. Using the serum western blot as the gold standard, the sensitivity and specificity of the salivary western blot was 38 of 47 (81%) and 68 of 75 (91%), respectively (table 1). The sensitivity and specificity of saliva ELISA on these same specimens was 29 of 47 (62%) and 48 of 75 (64%), respectively. Saliva western blotting was significantly more sensitive (p = 0.039) and significantly more specific (p = 0.00005) than saliva ELISA.

The presence of bands to the cagA and vacA proteins on the salivary and serum western blots were compared, with good overall agreement between the two methods (κ = 0.75, p < 0.000001 for CagA; κ = 0.66, p < 0.000001 for VacA; table 2).

PERFORMANCE OF SALIVAR WESTERN BLOT AGAINST SERUM ELISA

The sensitivity and specificity of saliva western blot compared with serum ELISA was 32 of 44 (73%) and 65 of 78 (83%), respectively (table 3). The sensitivity of salivary western blotting was not significantly greater than salivary ELISA (p = 0.39), whereas specificity was significantly greater (p = 0.0071).

COMPARISON OF SERUM WESTERN BLOT AGAINST SERUM ELISA

There was good overall agreement between serum western blot and serum ELISA (κ = 0.84, p < 0.000001; table 4). Of the six ELISA negative, but western blot positive,
Western blotting to diagnose Helicobacter pylori infection

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for the detection of Helicobacter pylori infection. Serum western blotting has a significantly better specificity than ELISA. Our study demonstrates that western blotting can detect organisms with which the patient is infected.

Table 2 Presence of cagA and vacA bands on serum and salivary western blots

<table>
<thead>
<tr>
<th></th>
<th>CagA serum positive</th>
<th>CagA serum negative</th>
<th>VacA serum positive</th>
<th>VacA serum negative</th>
</tr>
</thead>
<tbody>
<tr>
<td>Saliva positive</td>
<td>20</td>
<td>4</td>
<td>12</td>
<td>4</td>
</tr>
<tr>
<td>Saliva negative</td>
<td>6</td>
<td>92</td>
<td>6</td>
<td>100</td>
</tr>
</tbody>
</table>

$\kappa = 0.75$ $\kappa = 0.66$

Table 3 Performance of salivary western blot (WB) compared with serum enzyme linked immunosorbent assay (ELISA) by salivary ELISA status

<table>
<thead>
<tr>
<th>Salivary WB</th>
<th>Serum ELISA positive</th>
<th>Serum ELISA negative</th>
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<tr>
<td>Saliva Positive</td>
<td>25</td>
<td>3</td>
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<tr>
<td>Saliva Negative</td>
<td>3</td>
<td>21</td>
</tr>
<tr>
<td>Serum Positive</td>
<td>7</td>
<td>6</td>
</tr>
<tr>
<td>Serum Negative</td>
<td>9</td>
<td>44</td>
</tr>
</tbody>
</table>

$\kappa = 0.75$

Table 4 Agreement between serum enzyme linked immunosorbent assay (ELISA) and serum western blot (WB)

<table>
<thead>
<tr>
<th></th>
<th>Serum WB positive</th>
<th>Serum WB negative</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum ELISA positive</td>
<td>41</td>
<td>3</td>
<td>44</td>
</tr>
<tr>
<td>Serum ELISA negative</td>
<td>6</td>
<td>72</td>
<td>78</td>
</tr>
<tr>
<td>Total</td>
<td>47</td>
<td>75</td>
<td>122</td>
</tr>
</tbody>
</table>

Discussion

Our study demonstrates that western blotting has a significantly better specificity than ELISA for the detection of H. pylori antibodies in the saliva of children and is comparable with serum ELISA. Therefore, salivary western blotting might be useful in the study of children, where high specificity is of particular importance, owing to the low prevalence of infection. In comparison with serum western blotting, salivary western blotting was more sensitive than salivary ELISA. It was also superior when both were compared with serum ELISA, although this failed to reach significance.

Our study might have underestimated the specificity of salivary western blotting significantly, because a stratified sample was used, in which saliva positive serum negative specimens on ELISA were over represented. Although western blotting of saliva was very specific, it had a slightly lower sensitivity than serum ELISA, using serum western blot as the gold standard. However, specificity is more important than sensitivity in studying low prevalence populations, such as children in the developed world.

The qualitative nature of the western blot does appear to overcome the problems of variable transudation of serum immunoglobulin into saliva, which might affect ELISA optical density. Western blotting has the additional advantage of providing more information from the saliva samples, such as seropositivity to CagA or VacA, which can be used to type the organisms with which the patient is infected. There was good agreement between bands detected on serum and saliva western blotting, supporting the use of saliva for this purpose.

Saliva has some disadvantages compared with serum. In a number of specimens there was excessive debris, which required centrifugation. Unfortunately, there were still seven samples that yielded uninterpretable blots, and these were not included in the results. Another potential disadvantage is the sample volume required, although only 1 ml was used, which is not excessive.

Other problems that might impair the sensitivity of saliva western blotting are the freeze–thaw cycle of samples and the circumstances of sample collection. In our study, the saliva was three years old and had been subjected to a previous freeze–thaw cycle. The means of collection might also be important: theoretically, unstimulated saliva is better than stimulated saliva.

Saliva has several advantages over other methods of determining H. pylori status. It is non-invasive, unlike serum testing, and it is much cheaper and more convenient than C13 urea breath testing. New tests to detect H. pylori antigen in faeces hold promise, but faecal specimens are more difficult to obtain, particularly in children, and harder to work with in the laboratory. Preliminary reports of evaluation in children have given promising results.

Although western blotting is a reliable method and overcomes the inconvenience of a capture assay, such as ELISA, it is time consuming, but could and is being automated. However, because of the ease of collection and non-invasiveness, saliva remains attractive for the diagnosis of H. pylori infection, and salivary western blotting is therefore a potentially useful method for epidemiology and paediatric diagnosis.

Rationalised virological electron microscope specimen testing policy

I read the two letters from McCaughey and Curry and their respective colleagues with astonishment. Their debate on a policy for electron microscopy (EM) use reminded me of those mediaeval ones about the number of angels that could dance on the head of a pin. Interesting debate, pity it missed the point. They failed, I think, to address three important aspects:

(1) The nature of virus diagnosis. No one can pretend that virus laboratories can investigate every individual "viral" illness in the community, but they have an obligation to monitor what is prevalent in it. Although attractive to a cash strapped service, selecting specimens on the basis that some are more likely to yield positives than others misses the point of diagnosis—more than one virus can cause many "virus like" syndromes and the true prevalence does not parallel exactly the presence of symptoms. To discard specimens taken from those who are recovering because the yield may be low, or where the cause is apparently obvious, strikes me as arrogant. If someone has taken the trouble to send a specimen, it seems reasonable to look at it, if only in the hope that the sender might be encouraged to send others in the future, especially if it turns out to be positive. It is a constant battle to get worthwhile virological specimens sent to the laboratory—choking them off is daft. Moreover, I would further confirm the Irish view that solid stools may often yield positives and also that making the diagnosis by holding the specimen up to the light (metaphorically speaking) is very misleading. Trends in infection can only be given some credence if the specimen base remains more or less constant. Arbitrary and variable selection of what will be examined, and what will not be included in this base, Elaborating a "rational policy" is always a recipe for cutting down what is done (why else have one?), usually in pursuit of saving money, and is rarely based on sound science.

(2) The question of money. EM is unique among virological laboratory techniques in that the major costs of using it diagnostically (equipment, staff) are incurred in setting it up in the first place. Running costs, in comparison, are trivial, but savings are thought to be possible by allowing the operator to work only part time. Used full time, the technique becomes less expensive for each specimen the more it is used, but this use must be sensible, and humane to the operator. Microscopy, and EM in particular, can never really be a part time occupation—the operator needs to keep in constant practice and to be committed to it. The cream of his/her work should not be skimmed off by using other techniques (enzyme immunoassays, etc.)—to detect rotavirus or adenovirus—for example. Like everyone else, electron microscopists thrive on getting positive results and these reward immensely what can be a lonely working existence. Moreover, the laboratory gets several simultaneous tests for its money, results are produced more quickly than with other tests and, in most cases, with better economy.

(3) The need to retain EM as a non-centralised resource. With changes in travel and climate, "new" viruses can appear anywhere at any time. Part of characterising a putative new virus, whether truly novel or merely transferred to a new habitat, is to know what it looks like—other properties correlate surprisingly well with structure—and having only a national, or even a regional, facility for this purpose is usually a recipe for serious delay. Specimens from the periphery do not carry the same urgency as the facility's own work, especially if the latter has to meet internal performance targets. Passing perhaps the most interesting part to another laboratory is hardly likely to encourage the local virus hunters either.

Clamping EM into a "rational policy", instead of encouraging its use wherever possible, will ultimately cause it to die a slow death from a downward spiral of discourage ment. Put in rugby terms, use it or lose it. EM is the only truly "catch all" technique in diagnostic virology. If lost, it will be replaced (partially, even at best) with a battery of tests which, collectively, will cost more, take longer to do, and will frequently miss the dual or triple gut infections that are common in children.

The authors reply

McAuliffe’s perception of the continuing usefulness of diagnostic electron microscopy (EM) in virus laboratories is very different from ours. We believe that diagnostic EM is becoming largely redundant, electronic microscopy simplifies the training and skill mixes required to provide a comprehensive service compared with that required for a service delivered via diverse traditional methods.

The term “catch all” was first applied to diagnostic EM in the 1970s and is now inappropriate. EM is insensitive and non-specific. Our nested multiplex PCR assay can distinguish between HSV-1, HSV-2, and VZV in a slide from a vesicular skin lesion. EM will often miss the diagnosis, and even when positive will only indicate the presence of a virus from the Herpesviridae family.

In 30 years of diagnostic EM we have not discovered any new viruses. In the first year of routinely using nested PCR on all skin specimens we have discovered a deletion mutant of HSV type 1. Looking for new viruses via routine diagnosis is not a function of a diagnostic virology laboratory. Producing results that affect patient management and infection control in an efficient, timely, and economic fashion is a major function.

If Professor McAuliffe considers the rewarding aspect of diagnostic virology to be the positive result, then he should welcome the availability of more sensitive methodology. There is a responsibility for diagnostic laboratories continuously to improve their performance, rather than to continue with assays whose appeal is aesthetic rather than functional.

The authors reply

Professor McAuliffe’s comments are very welcome but his letter opens up the debate about electron microscopy (EM) to a much greater extent than that originally intended by either the rationalised EM policy paper or the recent correspondence relating to it. We strongly agree that EM should not be allowed to wither and die, but maintaining the relative abundance of EM units that existed into the late 1980s was untenable for two main reasons. First, the development of new investigative and diagnostic methodologies (initially
enzyme immunoassays and subsequently the polymerase chain reaction) and secondly financial pressures that have affected, not only the PHLS, but also hospital pathology departments and universities. Both these factors have meant that the use of EM in many centres has been critically reviewed. Many universities have centralised their EM facilities into units either specialising in materials science or biomedical science. Within pathology, many laboratories have given up locally provided EM services because of cost considerations and buy only the EM services that they require from established units, which is more cost effective than maintaining a local EM facility. Within the PHLS, a strategy for EM was formulated, which resulted in the formation of a strategic network of EM units in England and Wales being retained. This rationalised EM service provided significant improvements in virus surveillance and showed—for example, the true importance of small round structured viruses (SRSVs) as a cause of outbreaks of gastroenteritis in the UK.

A fundamental limitation of EM is that every specimen needs individual examination by a skilled microscopist. This aspect of EM cannot be automated and restricts the number of specimens that can be examined by a microscopist within a working day. Only through the provision of that warrant individual attention should be examined, and this is at odds with Professor Madeley's view that if a specimen is submitted it should be examined. Newer diagnostic methods are very sensitive, much cheaper for each test, can handle greater numbers, and often require less skilled staff to perform them. It is because of such developments that EM would no longer be considered as a front line test—for example, group A rotaviruses.

Professor Madeley also makes the point that EM is relatively cheap after the initial capital investment. However, this is over simplistic because microscopes and associated ancillary equipment all have finite lives. Even if an electron microscope were in operation for 20+ years, few organisations would have the money to replace such an instrument on anything other than an exchange or one to one basis. Even the remaining microscope units, the cost of annual maintenance contracts for such instruments is high, and when taken with the limited number of specimens that can be examined and the costs of a skilled electron microscopist, this means that a virological examination costs tens of pounds to perform. We would agree that EM is a "catch all" method, but specimen numbers and turn round times must be considered if an efficient diagnostic service is to be provided to the customer, who has to pay for such investigations.

What we have tried to achieve with our specimen testing policy is to provide a range of diagnostic techniques to our customers, which necessitates testing only the most appropriate specimens by EM; this is underpinned by epidemiological evidence of the relative benefit of discriminating between samples as opposed to the "shotgun" effect of unconsidered and unstructured sampling.

EM does have a future, but this probably lies with units that have a broad range of ultrastructural interpretative skills, carry out several preparative methods (such as negative staining and thin sectioning), and have a high throughput of specimens, thus maximising the use of these expensive facilities. Units able to undertake a variety of work, with skilled interpretation, will prosper. However, an important problem in the medium term is how to pass on established EM skills to a new generation of electron microscopists. Few organisations that have retained EM facilities have grappled with this problem of succession planning.

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Aberrant CD10 expression by NHL

The report by Millar and colleagues' of aberrant expression of CD10, the common acute lymphoblastic leukaemia (ALL) antigen, by the cells of a marginal zone non-Hodgkin's lymphoma (NHL) is important because it reminds histopathologists not to rely too heavily on the results of single surface marker tests. "Aberrant" expression of CD antigens by malignant lympho-haemopoietic cells is a phenomenon with which haematologists have long been familiar; expression of "myeloid" antigens CD13 and CD33 by the blast cells of common ALL and the "B lymphoid" antigen CD19 by those of acute myeloid leukaemia with the (8;21) translocation are two classic examples. But is not the range of CD10 positive NHL cases already rather broader than they suggest? We have known for years that up to 20% or so of lymphoblastic lymphomas have an immunophenotype already rather broader than they suggest? We have known for years that up to 20% or so of lymphoblastic lymphomas have an immunophenotype in distinguishable from common ALL and that a similar proportion of cell lymphoblastic lymphomas, albeit not acknowledged by the REAL classification, may also express CD10. In addition, the REAL classification also points out that some cases of mantle cell and diffuse large B cell lymphoma express CD10.

The range of monoclonal antibodies that work superbly on sections or by flow cytometry or both, will surely reveal more "aberrant" results, and may eventually modify our views about what "aberrant" means.

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that goes into the required details when discussing the diseases that are specific for the upper aerodigestive tract. The growth in knowledge and proliferation of diagnostic entities associated with the upper aerodigestive tract are reflected in the fact that the current fascicle counts 455 pages, whereas the head and neck fascicle from the previous series had 343. Another difference to the previous edition is the use of colour illustrations, which greatly enhances the quality of the book.

Of course, there are some issues on which there may be disagreement with the authors. The use of the term transitional carcinoma as a synonym for undifferentiated carcinoma is confusing because in the WHO classification this definition is used as a synonym for ciliated cell carcinoma, the tumour originally described by Ringertz in his seminal monograph. Moreover, equating psammomatoid ossifying fibroma with juvenile ossifying fibroma is unjustified; these lesions differ in histology as well as in predestination site and age. Finally, the discussion of the bony lesions occurring in the jaw is meagre but I suppose they will be the subject of a separate fascicle, as was the case in the previous series.

I can heartily recommend this book for anyone who has diagnostic responsibilities for specimens from the upper aerodigestive tract.

PIETER SLOOTWEG

CD-ROM review


I am a lover of well written textbooks and, I suspect, like most histopathologists a relatively late adopter of new technologies such as the use of the digital image. I therefore faced the review of this CD-ROM atlas of surgical pathology with some trepidation—the reader should take this into account. This CD-ROM installs, or uses, Quick Time, Adobe Acrobat, and Folio Bound Views to present and navigate between what is stated to be over 4000 images taken from 1500 different lesion types. The images are almost all haematoxylin and eosin stained sections. The quality of photomicrography is good—illumination is even, the staining is not obviously faded, and there are no colour casts. The colour intensity is rather strong and occasionally slightly muddy. The resolution of the images is not obviously stated, but is such that a full screen image looks good, without obvious pixellation, but when the zoom in facility is used, the pixellation becomes obvious when the magnification has been increased by 50% or more. In practice, this meant that the images were best viewed at full screen and not zoomed at all.

An animated tutorial takes the viewer through the possible ways of using the atlas. I found watching the mouse arrow move around the screen by itself and press buttons slightly tedious, and aborted this programme in favour of getting started and pressing all the buttons myself.

There appear to be two main ways of using this atlas. The first is entitled “pattern recognition”. In this, one selects the desired topo-

graphic site from a reasonably comprehensive list, presented around an image of a ghastly blue coloured human body, and then screens are presented primarily by histological pattern, with possible diagnoses as the subheadings. It immediately becomes apparent that to navigate this, one has to know the name of the histological picture being looked up—if you do not know what it is called, you do not know where to start looking. Thus—for example, if you were a trainee presented with a case of crescentic glomerulonephritis but did not know what it was called, you would have to leaf through all of the pictures of glomerular disease to find out what it was called. However, once there, information about the conditions is succinct and worthy of study. The juxtaposition of competing differential diagnoses makes navigating this section rewarding and attractive.

The second main mode of using the atlas is the “self assessment quiz mode”. In this, one again selects the desired topographical site and then brief histories are presented, together with a collection of thumbnail sketch histological images. These can then be interrogated further by enlargement before arriving at a mental diagnosis. There does not appear to be anywhere to record one’s diagnosis to the computer, and to have it marked. Instead, the diagnosis is revealed after clicking on a button. This at least avoids the frustration of computer marking of free text answers, where permissibility near misses get (infuriatingly) marked wrong. I found it interesting that it was reasonably easy to predict which cases were benign and which were malignant from the number of images presented. Malignant conditions can be justified within a few images. Benign ones usually take a lot of images at different magnifications to provide evidence of lack of malignancy. There is obviously a “learning curve” with the manipulation of digital images, and I found the exercise very difficult compared with direct microscopy of slides. The main problem appeared to be the slight over emphasis on high power images, the inability to select one's own field, and the lack of resolution compared with real microscopy.

On the positive side, however, was the requirement to make a commitment to a diagnosis on the basis of a given field. This is obviously something that could be used in the future for testing professional interpretation in a very standardised way.

All of this raises the question “who would derive most value from the program?”. I don’t think there is a lot of new information here for existing consultants. However, I can imagine the images being useful in teaching (I should note that I attempted to cut and copy some of the images out of the program, without knowing whether I was allowed to or not, but in any event, the program would not allow me to do this).Trainees might prefer this to a paper atlas because a large number of reasonably high quality images are contained here, and the indexing is fine for someone used to navigating electronic indexes. However, they would need to be at a level where they know the names of basic histological patterns. I would therefore envisage that this would be invaluable to SpRs as an adjunct to viewing slide collections and participating in bench work.

I would recommend that departments setting up an electronic learning resource for their trainees add this to the collection of available CD-ROM programs and think that practical use would be seen at least by those trainees who enjoy operating this type of technology.

TIM STEPHENSON

www.jclinpath.com

Letters, Book reviews, CD-ROM review, Notices, Correction

Notices

Muscle Disorders: Pathophysiology Applications and Techniques in Veterinary Pathology

New Millennium Bugs

Cytopathology Update: Making Cervical Cytopathology Work

27 September 2000, 5 October 2000, 18 October 2000, and 7 December 2000, respectively

Four one day symposia held at the Royal College of Pathologists, 2 Carlton House Terrace, London, SW1Y 5AF, UK. The symposia are open to members of the college, to trainee pathologists, and to workers in other disciplines with an interest in the subject. The programme is approved for CPD.

Further details: Scientific Meetings Officer, Royal College of Pathologists, 2 Carlton House Terrace, London SW1Y 5AF, UK; tel: +44 020 7451 6740/6739; email: www.repath.org

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6–8 November 2000

Royal Brompton Hospital, Imperial School of Medicine, and National Heart and Lung Institute

This “hands on” course approaches in detail the problems that face the diagnostic pathologist when dealing with cardiovascular pathology. Congenital heart disease will be highlighted. The approach to a cardiac necropsy and sudden death will be emphasised. Cardiac specimens will be made available for analysis and practical demonstrations as well as video demonstrations. A slide seminar is also included. The course is aimed at trainees studying for the MRCPATH and also senior pathologists who wish to update their knowledge of cardiac disease, both congenital and adult. Course fee £200.00 or £200.00 for juniors in training.

Details from: Short Course Office, National Heart and Lung Institute, Dovehouse Street, London SW3 6LY; tel: +44(0)20 73518172; fax +44(0)20 73518246; email: shortcourse.NHLI@IC.AC.UK

XIII International Congress for Quantitative Diagnostic Pathology

October 2000

Adelaide, Australia

The web site for this meeting will be available at the following address from 12th May 2000: http://som. flinders.edu.au/fusa/AnatPath/congress/ main1.htm
Correction


The authors would like to apologise for an error that occurred in this paper. It was stated throughout the paper that 665 children were tested but in fact the number was 669. In addition, in the first sentence of the results section of the abstract 530 of 691 (87%) should be 530 of 619 (86%). The authors stress that these errors do not change the message of the paper.