Bilirubin is a specific marker for the diagnosis of acute appendicitis

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Received September 27, 2020; Accepted January 19, 2021

DOI: 10.3892/etm.2021.10490

Abstract. Total serum bilirubin and other biochemical parameters have been associated with acute appendicitis, mainly in complicated cases. The present study aimed to evaluate the role of biochemical parameters in the diagnosis of acute appendicitis, and to further investigate the role of bilirubin as a diagnostic marker irrespective of the severity of the pathology. All recorded cases of appendicectomies in a 1-year period in a single institution were reviewed. The median values of white cell count, C‑reactive protein and total serum bilirubin on admission were associated with final histology, and their respective rates of abnormal and normal values were compared between patients who were proven to have negative histology and patients who were proven to have acute appendicitis. A total of 300 patients were studied. Median total serum bilirubin, white cell count and C‑reactive protein on admission were significantly associated with acute appendicitis (P<0.001). Respective rates of normal and abnormal values were significantly associated with final histology (P<0.001). Total serum bilirubin demonstrated higher specificity (0.88) but lower sensitivity (0.26) and diagnostic accuracy (0.40) for acute appendicitis. In conclusion, total serum bilirubin on admission should be considered in the diagnostic workup to confirm rather than exclude appendicitis, without focusing on subgroups of specific severity of the disease. White cell count and C‑reactive protein may also contribute to the diagnostic work‑up, although with limited accuracy.

Introduction

Acute appendicitis (AA) is currently the commonest acute surgical pathology, with a lifetime risk of 7-9% (1,2). Although various techniques have been advocated with the aim to reduce negative appendectomy rates in patients with acute abdominal pain, these remain between 15 and 50% (3) and seem higher in young female, elderly and paediatric patients (4). A safe, cheap, rapid, widely available and accurate diagnostic marker for AA would be useful, however currently there is no single diagnostic tool that can lead to a definitive diagnosis of AA (5,6). In the past, different authors have studied the diagnostic value of various laboratory tests, but results are contradictory (4). White Cell Count (WCC) as well as C‑reactive protein (CRP) are two of the markers that can contribute to the diagnosis of AA, with the advantages of being relatively inexpensive and carrying no risks, contrary to other methods such as computerized tomography with exposure to radiation or laparoscopy with potential surgical complications. Nevertheless, none of them is diagnostic or specific. Previous retrospective studies have demonstrated hyperbilirubinaemia to be a useful predictor of complicated acute appendicitis and appendiceal perforation (7,8) although contradiction exists in literature (2,9,10), but recently others have suggested an association between Total Serum Bilirubin (TSB) and simple appendicitis (3,6,11).

The present study's aim is to evaluate the usefulness of biochemical parameters (WCC, CRP, TSB) including a Receiver Operating Characteristic (ROC) curve analysis in the diagnosis of AA and to evaluate further specifically the role of TSB as preoperative diagnostic marker irrespective of the severity of the disease.

Materials and methods

Patients and parameters. All electronically recorded consecutive cases of appendicectomies in a 1-year period in a single centre were reviewed. Demographic data of all patients have been recorded. All cases in which symptoms were attributed to ovarian pathologies or other problems identified during surgery and those in which appendicectomy was not the primary procedure have been excluded from the study. In the cases in which the study of the surgical specimen did not confirm acute appendicitis or other pathology the histology was considered as negative. In cases where the study of the specimen confirmed acute appendicitis or any other pathology explaining the clinical findings the histology was considered as positive.

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Key words: bilirubin, white cell count, C‑reactive protein, acute appendicitis, histology
Median TSB value, as well as median WCC and CRP on day of admission were statistically studied in relation to the final histology result. A further comparison has been performed between patients shown to have negative histology results and patients demonstrated to have acute appendicitis regarding the percentage of abnormal and normal values of WCC, CRP and TBS on admission in each group. Any values of WCC higher than 11x10⁹ wbc/l, CRP higher than 5.5 mg/l and TSB higher than 21 µmol/l were considered as abnormal, while any values of WCC, CRP and TSB below 11x10⁹/l, 5.5 mg/l and 21 µmol/l respectively were considered as normal. CRP given as <5 mg/l was accounted as 4 mg/l.

As all cases in the current study derive from a single centre, the diagnostic work up and clinical management across the patients, such as history taking, clinical examination and blood tests are homogeneous. Based on the judgment of the responsible clinician, as well as on other factors such as patient's gender, age, availability of techniques, timing etc., the imaging method, if any, has varied among the patients (ultrasound scan, computerized tomography, magnetic resonance imaging or none). Conduction of this work is in full compliance with local Ethical Regulations and Anonymization standards. Approval from local ethical committee was not required as this was not an interventional study, involving only retrospective analysis of clinical data associated with diagnostic and therapeutic techniques performed without any deviation from institute's local guidelines. Data were analysed retrospectively thus informed consent from the patients prior to their inclusion in the study was not required according to local policy. All patients have signed an informed consent form prior to their operations.

Statistical analysis. Bivariate associations between scale and binominal variables were assessed using Mann-Whitney U test. Association of categorical variables in 4-fold tables were assessed using Fisher's exact test (2-tailed). P<0.05 was considered to indicate a statistically significant difference. Sensitivity and specificity analysis was performed, and subsequent ROC curve was produced. All aforementioned were carried out using SPSS v20 software (IBM Corporation).

Results

Descriptive statistics. Out of 311 appendicectomies, either open or laparoscopic, recorded in the study period, eleven cases have been excluded from the study. Median age at time of surgery was 27 years. 48.3% of cases were male patients. The negative appendicectomy rate reached 23% (69 patients). Acute appendicitis was confirmed in 77% of cases. Median TSB value, as well as median WCC and CRP on admission were 12.9, 25 and 12, respectively (Table I).

Comparisons. All median TSB, WCC and CRP on admission were significantly higher in patients with acute appendicitis compared to patients with negative histology (9 vs. 13, 9 vs. 13.9 and 4 vs. 37, respectively; P<0.001 for all three parameters) (Table II). Furthermore, after comparing the respective rates of normal and abnormal TSB, WCC and CRP on admission between the two groups of positive and negative histology, these were overall found to have a significant statistical association with final histology (P<0.014, P<0.001 and P<0.001, respectively) (Table III). The relative diagnostic values are shown in Table IV, with TSB demonstrating compared to WCC and CRP a clearly higher specificity for AA (0.88 vs. 0.67 for WCC and 0.42 for CRP), but with lower sensitivity (0.26 vs. 0.77 and 0.89, respectively) and overall accuracy (0.40 vs. 0.74 and 0.78, respectively).

ROC analysis. Fig. 1 presents cumulatively the ROC curves for the three biochemical predictors, namely WCC, CRP and TSB. Area under curve was 0.743, 0.755 and 0.703, standard error was 0.04, 0.037 and 0.045, respectively (P<0.001 for all three variables) (referring to equal probabilities).

Discussion

The relation between hyperbilirubinaemia and severe acute appendicitis has been reported since 1969 (12). It is postulated that elevated serum bilirubin occurs as a result of portal sepsis or empyema resulting in liver hepatocellular dysfunction or damage. This is thought to be caused by bacterial endotoxins or cytokines. The result is either a direct damage to hepatocytes, cholestasis, or both leading to hyperbilirubinaemia. Furthermore, endotoxins are shown to result in haemolysis, which then adds further increase in bilirubin levels. Other authors have reported that the elevated load of bacteria in appendicitis causes either direct invasion or translocation into the portal venous system. Direct invasion of bacteria into the hepatic parenchyma interferes with the excretion of bilirubin into the bile canaliculi by a mechanism that is thought to be caused by the bacterial endotoxin and is biochemical in nature rather than obstructive. On the other hand bacteria may transmigrate and produce portal bacteraemia, hepatocellular dysfunction or pyogenic liver abscess. The depression of hepatocellular function in the early, hyper-dynamic stage of sepsis does not appear to be due to reduction in hepatic perfusion but is associated with elevated levels of circulating pro-inflammatory cytokines such as TNF and IL-6. Thus up regulation of TNF and/or IL-6 may be responsible for producing hepatocellular dysfunction during the early hyper-dynamic stage of sepsis (13). Various studies have correlated elevated serum bilirubin and acute appendicitis, but most of them refer only

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male sex¹</td>
<td>145 (48.3)</td>
</tr>
<tr>
<td>Age at time of surgery²</td>
<td>27 (6-93)</td>
</tr>
<tr>
<td>WCC on admission²</td>
<td>12.9 (3.4-26.1)</td>
</tr>
<tr>
<td>CRP on admission²</td>
<td>25 (4-615)</td>
</tr>
<tr>
<td>TSB on admission²</td>
<td>12 (4-263)</td>
</tr>
<tr>
<td>Positive final histology²</td>
<td>231 (77)</td>
</tr>
</tbody>
</table>

¹Data are presented as count (%); ²Data are presented as median (range). WCC, white cell count; CRP, C-reactive protein; TSB, total serum bilirubin.
The rates of sensitivity, specificity and PPV in literature range respectively 8‑80, 56‑100 and 80‑100%, in accordance with the present results (2,3,6,7,11,15,17‑20). The present study significantly contributes to current scientific knowledge by providing additional data and evidence to further support the findings of the aforementioned studies and furthermore it makes a step forward by presenting a ROC curve analysis that clearly visualizes the advantages of the use of bilirubin.

WCC and CRP have been considered significant parameters for the diagnosis of AA (18) with comparable sensitivity, specificity, PPV, and NPV (4) and although they have been studied extensively for the diagnosis of AA, they lack sufficient specificity, either alone or in combination (4,21). There are studies that demonstrate clear differences in these values between patients with non‑surgical abdominal pain or early appendicitis and patients with phlegmons or perforated appendicitis (3,4,18), however the generally low sensitivity and specificity reported, does not allow for them to be considered as optimal indicators for AA (4,18). This is in line with the current study, with results generally comparable to the rates of sensitivity and specificity reported in literature for WCC (43‑93 and 50‑87% respectively) (4,15,17,18,22,23) and CRP (53‑97.2 and 46‑99.3% respectively) (15,24‑27). Elevated WCC demonstrates an acceptable sensitivity and at the same time a relatively low specificity as it can be triggered by various conditions (4). CRP levels have been reported as significantly higher in patients with perforated appendicitis (4,6,15) and CRP has been considered a more sensitive test in discriminating the pathological severity of appendicitis (24,28), with higher accuracy than WCC (15,27). Nevertheless, still its relative specificity is low as revealed by the present study and unable to enhance the role of CRP in the diagnosis of AA, unless integrated to other parameters.

Limitations of the study include its retrospective nature, the inclusion only of patients who underwent appendicectomies which may cause an increase of the specificities of the study and the non‑discrimination of the severity of disease across the cases. Furthermore, the possibility of inclusion of patients with Gilbert's syndrome in the cohort studied should be considered, although it represents an uncommon condition reported in 3‑10% the population (3). The cohort consisted of all consecutive patients operated for suspected appendicitis, therefore there were obviously patients that were eventually proven not to have appendicitis; thus cases were not previously confirmed. The negative rate is similar to that reported in literature, as discussed, so it is considered representative. Although specificity is indeed of less importance to diagnose the disease, this is the actual clinical question addressed, i.e., a way to reduce operations in to perforated appendicitis (14). While hyperbilirubinaemia in these cases can contribute to the diagnosis, the clinical presentation is more diagnostic, therefore the role of elevated serum bilirubin is probably limited. In the present study a significant relation between TSB and AA was demonstrated. The increase of TSB was significantly overall noted in patients with AA irrespective of perforation. Even though median TSB was within normal range in both groups with positive and negative histology, like in other studies (2,15), the values of median TSB varied significantly between these two groups, in line with other authors (3,15). Furthermore, similarly to other reports, the incidence of abnormal bilirubin among patients in the positive histology group was more than double compared to the negative appendicectomy group (3,16). As Table III demonstrates, TSB was significantly associated with the histology results. Interestingly, although presenting relatively low sensitivity and overall diagnostic accuracy, TSB was found in the current study to have a high specificity, as well as a high Positive Predictive Value (PPV) and this is in line with other publications that demonstrate a high specificity for TSB in acute appendicitis, yet with considered low sensitivity. The rates of sensitivity, specificity and PPV in literature range respectively 8‑80, 56‑100 and 80‑100%, in accordance with the histology results. Interestingly, although presenting relatively low sensitivity and overall diagnostic accuracy, TSB was found in the current study to have a high specificity, as well as a high Positive Predictive Value (PPV) and this is in line with other publications that demonstrate a high specificity for TSB in acute appendicitis, yet with considered low sensitivity. The rates of sensitivity, specificity and PPV in literature range respectively 8‑80, 56‑100 and 80‑100%, in accordance with the histology results. Interestingly, although presenting relatively low sensitivity and overall diagnostic accuracy, TSB was found in the current study to have a high specificity, as well as a high Positive Predictive Value (PPV) and this is in line with other publications that demonstrate a high specificity for TSB in acute appendicitis, yet with considered low sensitivity. The rates of sensitivity, specificity and PPV in literature range respectively 8‑80, 56‑100 and 80‑100%, in accordance with

## Table II. Comparison of values on admission with final histology.

<table>
<thead>
<tr>
<th>Biochemical parameters</th>
<th>Negative</th>
<th>Positive</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>WCC on admission</td>
<td>9 (3.8‑18.3)</td>
<td>13.9 (5.4‑26.1)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>CRP on admission</td>
<td>4 (4‑256)</td>
<td>37 (4‑615)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>TSB on admission</td>
<td>9 (4‑71)</td>
<td>13 (4‑263)</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

Data are presented as median (range). WCC, white cell count; CRP, C‑reactive protein; TSB, total serum bilirubin.

## Table III. Comparison of patients with normal and abnormal values, with final histology results.

<table>
<thead>
<tr>
<th>Biochemical markers</th>
<th>Final histology</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Biochemical markers</td>
<td>Negative</td>
<td>Positive</td>
</tr>
<tr>
<td>WCC</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>46 (66.7)</td>
<td>54 (23.4)</td>
</tr>
<tr>
<td>Raised</td>
<td>23 (33.3)</td>
<td>177 (76.6)</td>
</tr>
<tr>
<td>CRP</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>29 (42.0)</td>
<td>25 (19.1)</td>
</tr>
<tr>
<td>Raised</td>
<td>40 (58.0)</td>
<td>206 (80.9)</td>
</tr>
<tr>
<td>TSB</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>61 (88.4)</td>
<td>172 (74.5)</td>
</tr>
<tr>
<td>Raised</td>
<td>8 (11.6)</td>
<td>59 (25.5)</td>
</tr>
</tbody>
</table>

Data are presented as count (%). WCC, white cell count; CRP, C‑reactive protein; TSB, total serum bilirubin.
the absence of appendicitis. The authors of this study agree that a diagnostic test should not be implemented solely on the basis of high specificity, but in this particular context, where current diagnostic algorithms have achieved good sensitivity, such a test could be a useful adjunct for the reduction of unnecessary operations. A comprehensive assessment of the whole extent of diagnostic usefulness of TSB would require performing the full diagnostic algorithm to all patients that attended emergency department with the cardinal symptom, but that would rightly raise the concern of running expensive tests for patients where history taking and objective examination have excluded the diagnosis of appendicitis. Therefore, the authors believe that this study accurately represents the population and the relevant conditions in which the clinical question arises.

With the present study, the known association of WCC and CRP with acute appendicitis is again highlighted although the diagnostic accuracy of each single factor is limited. On the other hand, although hyperbilirubinaemia in the assessment of AA is not widely used in daily clinical practice, TSB had a clearly higher specificity than CRP and WCC overall in patients with appendicitis, not only in the complicated cases. This highlights the role of TSB as a diagnostic parameter irrespective of the severity of the disease and probably indicates that surgeons do not ignore TSB level in the diagnostic work up as it could improve specificity. This suggested addition of TSB and the anticipated increase in the accuracy of the overall diagnostic approach can lead to decrease of unnecessary operations with a subsequent significant benefit for patients and healthcare systems.

In conclusion, the low sensitivity and overall diagnostic accuracy of TSB compared to WCC and CRP indicates that the single diagnostic laboratory parameters should not been considered independently but in combination to each other, along with the clinical picture and eventual radiological adjuncts. Because of its high specificity, TSB can be considered useful to confirm rather than exclude the diagnosis of AA. A potential role as independent diagnostic marker could

### Table IV. Statistic values of WCC, CRP and TSB in predicting acute appendicitis.

<table>
<thead>
<tr>
<th>Statistic values</th>
<th>WCC&lt;sup&gt;a&lt;/sup&gt;</th>
<th>CRP&lt;sup&gt;a&lt;/sup&gt;</th>
<th>TSB&lt;sup&gt;a&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sensitivity</td>
<td>0.77 (0.71-0.82)</td>
<td>0.89 (0.85-0.93)</td>
<td>0.26 (0.20-0.31)</td>
</tr>
<tr>
<td>Specificity</td>
<td>0.67 (0.56-0.78)</td>
<td>0.42 (0.30-0.54)</td>
<td>0.88 (0.81-0.96)</td>
</tr>
<tr>
<td>Positive predictive value</td>
<td>0.89 (0.84-0.93)</td>
<td>0.84 (0.79-0.88)</td>
<td>0.88 (0.80-0.96)</td>
</tr>
<tr>
<td>Negative predictive value</td>
<td>0.46 (0.36-0.56)</td>
<td>0.54 (0.40-0.67)</td>
<td>0.26 (0.21-0.32)</td>
</tr>
<tr>
<td>Odds ratio</td>
<td>7.70 (4.98-11.88)</td>
<td>5.15 (3.67-7.23)</td>
<td>2.62 (1.18-5.79)</td>
</tr>
<tr>
<td>Diagnostic accuracy</td>
<td>0.74 (0.69-0.79)</td>
<td>0.78 (0.74-0.83)</td>
<td>0.40 (0.35-0.46)</td>
</tr>
</tbody>
</table>

<sup>a</sup>(95% confidence interval). WCC, white cell count; CRP, C-reactive protein; TSB, total serum bilirubin.

### Figure 1. ROC curve for WCC, CRP and TBS. WCC, white cell count; CRP, C-reactive protein; TBS, total serum bilirubin.
be investigated further with larger multicentre prospective studies considering clinical scoring systems too.

Acknowledgements

Not applicable.

Funding

No funding was received.

Availability of data and materials

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Author’s contributions

DZ, PML, JB, GS, EAC and VS made substantial contributions to conception and design. DZ, JB, GS and EAC contributed to acquisition of data. DZ, GS, PML and VS contributed to analysis and interpretation of data. DZ, JB, GS, and EAC participated in drafting the article. DZ, PML and VS participated in revising the article critically for important intellectual content. DZ, PML, GS, JB, EAC and VS have given final approval of the version to be submitted and any revised version. All authors read and approved the final manuscript.

Ethics approval and consent to participate

Conduction of this work is in full compliance with local Ethical Regulations and Anonymization standards. Approval from local ethical committee was not required as this was not an interventional study, involving only retrospective analysis of clinical data associated with diagnostic and therapeutic techniques performed without any deviation from institute’s local guidelines. The study analysed data retrospectively thus informed consent from the patients prior to their inclusion was not required according to local policy.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

References