INTRODUCTION

Liver diseases, particularly hepatocellular carcinoma, are common in Kenya. Resources for needle biopsy with subsequent histological diagnoses are not available in most district hospitals. Fine-needle aspiration (FNA) and cytology of aspirated cellular material is a cost effective technique of diagnosis that is easy to use in smaller health facilities (1-4). In many centres, FNA has largely replaced conventional large needle core biopsy in the diagnosis of focal lesions in the liver (1-7). It has the advantage of less discomfort, very low risk of complications, hospitalisation of patient not necessary and enables a quick decision in options for clinical management (1-4). Its additional advantage is that it can be repeated from different sites until a satisfactory sample is obtained, without undue risks to the patient (1,7). Use of imaging techniques make possible sampling of lesions, which may be inaccessible by other sampling methods, thereby...
making possible a tissue diagnosis in patients whose clinical conditions preclude other forms of obtaining specimens for investigation. Repeating of FNA from different sites has been reported to increase diagnostic sensitivity to over 90% (1-3).

In the evaluation of liver disease, the main indications for FNA cytology are single or multiple focal abnormalities demonstrated by palpation, CT scan or by ultrasonography (4).

The aim of this study was to test the utility of FNA cytology in excluding malignant disease of the liver at district hospitals, where histopathology facilities are not established. The accuracy of FNA cytology in the specific diagnosis of primary hepatocellular carcinoma has been reported to increase to over 93% when combined with cell blocks (5,6). In this study only smears were used.

MATERIALS AND METHODS

Patients with suspected liver disease visiting Murang’a and Machakos District Hospitals were evaluated and those with nodular hepatomegaly were indentified for study. All the patients evaluated had right upper abdominal complaint and all had hepatomegaly. The lesions were located by palpation or ultrasonography. Focal lesions were aspirated by transcutaneous fine needle aspiration (FNA) using a 10ml syringe and 21 gauge 1½ inch length needle. The cellular material aspirated was dropped onto a microscope slide, spread thinly and then immediately fixed while still wet in 95% ethanol for 30 minutes.

The fixed smears were then air-dried, packed in slide-mailers and transported to the cytology laboratory at KEMRI in Nairobi. The fixed smears were stained by the Papanicolaou technique and interpreted on a light microscope by a cytologist.

RESULTS

Fifty patients were evaluated. Twenty one of the 50 (42%) specimens had malignant cells; fifteen of the 50 (30%) had necrotic cells; two of the 50 (4%) had fatty change; one of the 50 (2%) had reactive cell changes later confirmed to be cirrhosis on histology, and 11 of the 50 (22%) showed normal hepatocytes.

<table>
<thead>
<tr>
<th>Diagnosis</th>
<th>No.</th>
<th>(%)</th>
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<tbody>
<tr>
<td>Hepatocellular carcinoma</td>
<td>21</td>
<td>42</td>
</tr>
<tr>
<td>Necrosis</td>
<td>15</td>
<td>30</td>
</tr>
<tr>
<td>Fatty change</td>
<td>2</td>
<td>4</td>
</tr>
<tr>
<td>Reactive cell changes</td>
<td>1</td>
<td>2</td>
</tr>
<tr>
<td>Normal hepatocytes</td>
<td>11</td>
<td>22</td>
</tr>
</tbody>
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*Malignancy could not be excluded in the cases of necrosis.

Figure 1
Liver aspirate showing ductal carcinoma in situ, cribriform type. Large monolayered sheet of uniform epithelial cells with rounded spaces Pap. Stain x 40.
Figure 2
Papillary clusters of pleomorphic malignant cells from a liver aspirate. Pap. Stain x40

Figure 3
Dysesive, pleomorphic malignant cells exhibiting coarse chromatin and prominent nucleoli. Note a cluster of cholangiocytes with small hyperchromatic nuclei at the top of the photomicrograph. Pap. Stain. x40

Figure 4
Necrosis of liver cells from an aspirate exhibiting degenerative nuclei whose cytoplasmic borders cannot be discerned surrounded by erythrocytes. Pap. Stain. x40
DISCUSSION

Diagnosis is an essential component of the management of any disease. FNACytology established diagnoses on which the management of the 50 cases of liver disease were based: 21 hepatocellular carcinomas, 15 cases of necrotic liver, 2 cases of fatty change, 1 case of reactive hepatocytes and 11 normal livers. To get these diagnoses on tissue biopsies would have required many times the resources used for FNACytology.

CONCLUSIONS

Diagnostic utility of FNACytology is high, especially when the lesions are precisely identified clinically and augmented by liver ultrasound. It is less invasive, and can be done as an outpatient procedure. Clinical management options in these patients were made based on the cytological diagnosis. The results supported the diagnostic utility of FNACytology in the management of liver disease, especially in institutions without histopathology facilities.

REFERENCES