The diagnosis of chronic pancreatitis depends on evidence of permanently abnormal pancreatic morphology, impaired function, or both. The best available methods for the study of pancreatic exocrine capacity and assessment of pancreatic insufficiency are tests requiring duodenal intubation with measurement of pancreatic enzyme and bicarbonate secretion. However, these tests are expensive, time consuming, and invasive. Faecal fat analysis is also time consuming, requires special laboratory equipment, and is distasteful to the patient, ward, and laboratory staff. The indirect (tubeless) pancreatic function tests, for example, para-amonobenzoic acid test and pancreolauryl test (PLT) have been increasingly used to assess for pancreatic insufficiency, particularly in non-specialist units. They are easily performed by the patient, require standard laboratory equipment for analysis, are not expensive and have a high negative predictive value. The British Society of Gastroenterology guidelines for investigation of suspected malabsorption suggest direct tests should be restricted to specialist units and that PLT and para-amonobenzoic acid test are adequate screening tests.

Over the last few years, the PLT has been used in our unit to assess cases of chronic diarrhoea with suspected malabsorption. We carried out an audit of the use of the PLT to determine:

1. The frequency of abnormal tests and whether subsequent clinical events matched these results.
2. If any symptoms or simple blood tests predict an abnormal clinical event.
3. If PLT could be better targeted.

**METHODS**

**Patients**

Patients were identified from the hospital records system and pathology laboratory reports. Retrospective analysis of patients’ case notes was performed by two independent investigators.

The patients’ demographic details, symptoms (diarrhoea, steatorrhoea, abdominal pain, and weight loss), alcohol intake, biochemical and haematological blood tests, radiological and endoscopy investigations with relevant histology were recorded.

The audit included all PLTs performed over a three year period.

**Statistics**

Results are expressed as mean (SD). Regression analysis was performed to determine the significance of relations between clinical parameters and test results. A p value of <0.05 was considered significant.

**RESULTS**

Forty seven patients, who had a PLT over a three year period, were included in the audit. The age range was 17–76 years (48 (13) years). Twenty six were male (55%). Seven patients had a repeat test for equivocal results. Alcohol consumption was recorded in 40 patients (mean 33.1 units/week) with 11 men and five women were taking more than safe recommended limits (28 units/week and 21 units/week respectively).

Thirty eight patients (81%) had diarrhoea of three months duration. Nineteen patients had a weight loss of 1–5 stones.

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Thirty eight patients (81%) had diarrhoea of three months to 12 years’ duration, of whom 10 described stools suggestive of steatorrhoea. The remainder had watery stools. Twenty two patients (47%) had abdominal pain of three months to several years’ duration. Nineteen patients had a weight loss of 1–5 stones.

Twenty three patients (49%) had an abnormal PLT. Ten of 21 females (48%) and 13 of 26 males (50%) had abnormal results. Table 1 shows the number of abnormal and normal PLT results in relation to the patients’ symptoms. Age was not correlated with PLT results (fig 1) and alcohol intake was poorly correlated (fig 2).
Of the 28 patients with watery diarrhoea, 17 (61%) had an abnormal PLT while four of 10 patients with steatorrhoea (40%) had an abnormal test. In the absence of diarrhoea, weight loss and abdominal pain were not associated with an abnormal PLT.

Table 2 shows the biochemical results in relation to the PLT. Abnormal full blood count, calcium, and haematinics tended to predict an abnormal PLT but normal results did not indicate normal pancreatic function. Liver function tests were not predictive of the PLT result.

Duodenal biopsies were performed in 12 of the abnormal PLT patients. Two showed villous atrophy. Eleven patients with an abnormal PLT had an abdominal ultrasound, of whom seven (64%) had structural pancreatic damage.

All the patients with normal PLT were thought clinically not to have pancreatic insufficiency. Nineteen of the 23 patients with abnormal PLT were thought to be suffering from pancreatic malabsorption. Two with an abnormal PLT had a final diagnosis of coeliac disease.

Eighteen patients had an excellent response to pancreatic enzyme supplements (Creon), with marked improvement in the diarrhoea and weight gain. One patient improved on no therapy and another patient failed to attend for follow up.

DISCUSSION

Our audit demonstrates that the PLT can be a useful test to screen for pancreatic insufficiency in a district general hospital unit. Twenty three out of 47 patients (49%) had abnormal results, of whom 19 were believed to be suffering from pancreatic malabsorption. On the other hand, all patients with a normal PLT were thought not have pancreatic insufficiency, indicating a high specificity.

Diarrhoea was the most common presentation in the group with an abnormal PLT. Interestingly, watery stools were more frequently associated with a positive test than steatorrhoea. Abnormal results were not seen with other presentations (weight loss and abdominal pain) in the absence of diarrhoea.

Males and females were equally likely to have an abnormal PLT and alcohol history did not predict the result.

Two patients with abnormal PLT results were found to have coeliac disease. Their abnormal result could be due to concomitant pancreatic insufficiency or to fat malabsorption secondary to villous atrophy. Forty percent of newly diagnosed coeliac patients have a positive PLT while only 18% of those on a gluten-free diet had a positive test, demonstrating that fat malabsorption may respond to diet. As the PLT was not repeated after starting a gluten-free diet, it is not known if these two tests were “false positives”. Low serum calcium and haematinics with an abnormal PLT should prompt distal duodenal biopsies. The PLT has a high negative predictive value in the patients with coeliac disease and small bowel disease without pancreatic dysfunction.

Abdominal ultrasound was not requested for all patients because it has poor sensitivity and is operator dependent in the investigation of pancreatic disease. Abdominal computed tomography, endoscopic retrograde cholangiopancreatography, or magnetic resonance cholangiopancreatography may be more appropriate, although many cases of pancreatic insufficiency are associated with normal imaging.

All patients who had an abnormal PLT received pancreatic enzyme supplements (Creon) and showed marked improvement in their response to treatment was considered sufficient to confirm the diagnosis of pancreatic insufficiency.

The PLT was easy for patients to carry out. Verbal instructions were given to each patient by the laboratory and pharmacy staff.

| Table 1: Relation between the pancreolauryl test (PLT) and patients’ symptoms |
|-----------------------------------|-------------------|-------------------|------------------|-------------------|
| Diarrhoea (n=38)                  | Watery diarrhoea  (n=28) | Steatorrhoea (n=10) | Weight loss (n=19) | Pain (n=22)       |
| Abnormal PLT                      | 21                | 17                | 4                | 10               | 10               |
| Normal PLT                        | 17                | 11                | 6                | 9                | 12               |

| Table 2: Pancreolauryl test (PLT) results and blood tests results |
|------------------|------------------|------------------|
| Abnormal PLT     | Positive test    | Negative test    |
| Full blood count | 10/47            | 7/10             | 14/37            |
| Liver function tests | 12/42           | 7/13             | 11/29            |
| Calcium          | 1/32             | 1/1              | 15/31            |
| Haematinics      | 3/31             | 3/3              | 12/28            |
Pancreolauryl test in a district general hospital

complemented by a typewritten instruction sheet. The test was suitable for use with outpatients as has previously been demonstrated and can be easily repeated. The audit was carried out for a period of three years (1996–98) before faecal pancreatic elastase was widely used. We are now conducting another audit to compare both tests in investigating pancreatic insufficiency in the unit over the last two years. The diagnosis of early chronic pancreatitis can be difficult and sometimes patients with pancreatic insufficiency may not present with classic symptoms. Although this audit confirms the high specificity of the PLT in a district general hospital setting, it does not determine the sensitivity. This has previously been reported as 30%–50% for mild pancreatic insufficiency with minimal anatomical changes on endoscopic retrograde cholangiopancreatography to >95% for severe insufficiency with minimal anatomical changes on endoscopic retrograde cholangiopancreatography to >95% for severe chronic pancreatitis. It should be emphasised that the PLT would not exclude chronic pancreatitis in patients who do not have loss of exocrine pancreatic function.

The diagnosis of pancreatic insufficiency should be considered in all patients with unexplained chronic diarrhoea and weight loss. Pancreatic insufficiency may be seen in the absence of steatorrhoea and without a history of high alcohol intake. This audit shows that the PLT is a simple, inexpensive, and non-invasive screening test with specificity of 83%–91% for diagnosing pancreatic insufficiency in a district general hospital setting.

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