Studies on the mechanism of non-visualization of diseased human gallbladders during oral cholecystography

M.R. Jacyna, P.E. Ross, D. Hopwood, and I.A.D. Bouchier*

Department of Medicine, Ninewells Hospital and Medical School, Dundee DD1 9SY, Scotland, UK.

Summary: Oral cholecystography is a well established method for studying the human gallbladder and radiological non-visualization of the gallbladder has been shown to correlate highly with the presence of disease. The exact mechanism by which diseased gallbladders fail to visualize is unclear, but may be due to a failure of the gallbladder to concentrate the luminal contents. Concentration of gallbladder contents is achieved by the reabsorption of water, the driving force for which is active sodium (Na+) absorption. Therefore Na+ transport was studied by measuring the flux of Na22 across isolated human gallbladder mucosa (obtained at cholecystectomy) and compared with the results of oral cholecystography and histological grading.

In 27 gallbladders studied, 59% absorbed Na+, whilst the remainder secreted Na+. Comparison with histological grading showed that as gallbladders became more diseased they absorbed less Na+ and were more likely to secrete Na+. In addition, gallbladders that absorbed Na+ were significantly more likely to visualize on cholecystography than those that secreted Na+. These results indicate that some diseased human gallbladders secrete, rather than absorb, Na+ and suggest that the mechanism for radiological non-visualization is failure of fluid absorption and the development of active fluid secretion.

Introduction

Oral cholecystography is a simple and effective method for studying gallbladder function and disease. The technique relies on the ability of the gallbladder mucosa to concentrate adequately the luminal bile containing orally-administered contrast media. Concentration of the bile is achieved by the absorption of water, and successful concentration of this medium results in radiological opacification of the gallbladder. Failure of the gallbladder to visualize has previously been shown to be a very good indicator of underlying gallbladder disease. Although the majority of diseased gallbladders fail to opacify because of a stone blocking the cystic duct, some have a patent cystic duct and the reason why these gallbladders fail to opacify is unclear. Although it would seem likely to be due to an inability of the inflamed gallbladder mucosa to concentrate adequately the contrast medium, previous work has suggested that it may be due to abnormal absorption of the contrast medium by the diseased gallbladder mucosa. Studies on fluid absorption in the human gallbladder, however, are difficult to perform and prone to inaccuracy. As the driving force for water absorption in the gallbladder is sodium ion (Na+) transport via the mucosal enzyme sodium, potassium-dependent adenosine triphosphatase, we studied this process in diseased human gallbladders obtained at cholecystectomy using a sensitive radio-isotopic technique. By comparing the findings with the results of oral cholecystography and histology we hoped to gain more information about the mechanism of radiological non-visualization of diseased gallbladders.

Methods

Human gallbladders

Twenty-seven chronically inflamed human gallbladders, obtained at cholecystectomy from patients with cholelithiasis, were studied. The gallbladders were received within 15 minutes of ligation of the cystic artery and immediately placed in oxygenated saline (NaCl solution 150mmol/l) at 4°C. Portions of the fundus of the gallbladder were gently washed...
with saline to remove surface bile, and the serosal tissue was removed by blunt dissection and the isolated mucosa mounted in the flux chamber. The remainder of the gallbladder was kept for histological assessment. The degree of chronic cholecystitis was assessed by one observer (DH), who was unaware of the results of sodium transport and radiology. The gallbladders were graded as mildly (grade 1), moderately (grade 2) or severely (grade 3) diseased based on the appearance of the epithelial cells, the muscle layer thickness, the size and site of Rokitansky-Aschoff sinuses and the amount of chronic inflammatory cell infiltrate.4

**Oral cholecystography**

Oral cholecystograms were performed up to 4 months prior to cholecystectomy. On the evening before the examination, the patient ate a light supper consisting of non-fatty foods and then swallowed six 500mg tablets of lopanoic acid ('Telepaque'; Sterling Research Laboratories) with water. They were then fasted until the examination which took place the following morning.

**Sodium transport**

Sodium transport was studied using a modified 'Ussing' flux apparatus,4 and the technique used is as previously described with slight modifications.6,7 The flux apparatus consists of two perspex hemi-chambers with a section of gallbladder musosa interposed, allowing 0.2cm² of both surfaces of the mucosa to be independently bathed. Experiments were performed under 'open circuit' conditions. The medium bathing both surfaces consisted of Krebs bicarbonate buffer (pH 7.4) containing glucose (28mmol/l), and a tracer dose of Na₂² was added to one side of the tissue. The solutions were gassed continuously with 95%/5% O₂/CO₂ and thermostatically maintained at 37°C. Experiments used four chambers per gallbladder; two for the determination of unidirectional mucosal to serosal flux and two for the determination of unidirectional serosal to mucosal flux. Samples were taken at 10 minute intervals over the 20 to 60 minute period (when a steady rate of Na₂² transfer was obtained) from the fluid which initially contained no isotope and the unidirectional flux was calculated as the average of 8 readings (i.e. 4 per chamber). The difference between the two unidirectional fluxes represents the net sodium flux. A net mucosal to serosal sodium flux (sodium absorption) was designated as positive and a net serosal to mucosal sodium flux (sodium secretion) was designated as negative. Data is presented for the experiments as the mean±s.e.m. and statistical analysis was performed using Student's t-test.

**Results**

**Sodium transport and histological grading**

Figure 1 shows the net sodium flux of each gallbladder compared to its histological grading. As the severity of cholecystitis increased, the net flux for the gallbladders became less positive (indicating reduced Na⁺ absorption). In the moderately and severely diseased groups, a large proportion of the gallbladders had a net negative Na⁺ flux indicating active secretion of Na⁺.

**Sodium flux and radiological visualization**

Net sodium flux was compared to the results of oral cholecystography. Figure 2 shows that the group of gallbladders that opacified had a net positive (absorptive) Na⁺ flux that was significantly higher than the gallbladders that did not opacify, which had a net negative (secretory) Na⁺ flux.

**Discussion**

The main function of the gallbladder is storage of bile during inter-digestive periods, and concentration of the stored bile.8 Concentration of the bile occurs by the absorption of fluid and gallbladder mucosa has one of the highest reported rates of

---

**Figure 1** Net sodium flux of gallbladders compared to the histological grade: 1, mildly diseased; 2, moderately diseased; 3, severely diseased. Lines represent the mean and s.e.m. of each group.
fluid absorption. The driving force for this absorption is sodium transport via sodium, potassium-dependent adenosine triphosphatase, which actively extrudes Na\(^+\) across the basolateral cell membrane, creating an electrochemical gradient along which water and anions (bicarbonate and chloride) pass. Studies have confirmed that the rate of fluid transport correlates directly with sodium, potassium-dependent adenosine triphosphatase activity.

Our results indicate that increasing disease of gallbladders not only reduces Na\(^+\) absorption (and hence concentrating ability) but also initiates a secretion of sodium by the mucosa. As Na\(^+\) and water absorption are so intimately linked, it is likely therefore that the reason these gallbladders fail to opacify on cholecystography is because of active fluid secretion into the gallbladder lumen. The effect of this secretion will be a ‘washing out’ of the gallbladder contents, which will also prevent any contrast medium from entering the gallbladder.

Na\(^+\) and water secretion in inflamed animal gallbladders is well described and is an energy-requiring process which can take place against an osmotic gradient. In cholecystitis, fluid secretion is believed to be mediated via prostaglandins and reversal to the more usual absorption by prostaglandin inhibitors has been demonstrated in both animals and man. This is the first study to compare Na\(^+\) transport in human gallbladders with radiological function. However, Nahrwold et al. have compared ‘short-circuit current’ (which is an indirect measure of Na\(^+\) transport) of 26 isolated gallbladders with the results of oral cholecystography and found a similar correlation, although there was considerable overlap. Conversely, another study of water transport in the human gallbladder found no correlation between gross fluid absorption and radiological function and the authors postulated that the reason for non-visualization may be due to absorption of the contrast media by the inflamed gallbladder mucosa. The authors of this latter paper however are self-critical of their method and admit that ‘experimental error’ may limit the conclusions. Measurement of Na\(^+\) flux across epithelia is simple and sensitive and our data support the opposite contention that cholecystography does in fact assess the concentrating ability of the gallbladder. We believe that the reason that diseased gallbladders fail to opacify during cholecystography is not only due to failure of fluid absorption but also because of active fluid secretion.

Acknowledgements

We are grateful to the surgeons at Ninewells Hospital, Dundee for supplying cholecystectomy specimens.

References

Studies on the mechanism of non-visualization of diseased human gallbladders during oral cholecystography.

M. R. Jacyna, P. E. Ross, D. Hopwood and I. A. Bouchier

Postgrad Med J 1988 64: 931-934

doi: 10.1136/pgmj.64.758.931

Updated information and services can be found at:
http://pmj.bmj.com/content/64/758/931

These include:

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/