SUMMARY

During the past three decades, prolonged monitoring of oesophageal pH has become increasingly popular among physicians. Most of the studies carried out with this technique focused on oesophageal acid exposure, defined as percentage of time with pH below 4. However, in most cases acid exposure variables are not very useful in the evaluation of patients with symptoms and signs of gastro-oesophageal reflux disease (GORD). An often neglected aspect of 24 hour pH recording is assessment of the temporal relation between reflux and symptoms. Various methods to quantify this relationship were developed. Here, we review these methods and the contribution of symptom association analysis to the evaluation of patients with symptoms suggestive of GORD.

INTRODUCTION

The use of prolonged monitoring of oesophageal pH as a measure of gastro-oesophageal reflux was first described in 1969. More than a decade after its introduction as a hospital based technique for inpatients, oesophageal pH monitoring became rather popular when ambulatory data recorders became available. Most of the studies carried out with the technique focused on measurement of oesophageal acid exposure, defined as percentage of time with pH below 4, in patients with various manifestations of GORD. Later, the notion emerged that 24 hour pH recording also makes it possible to investigate the temporal relation between reflux and symptoms. In this review, we focus on the contribution of symptom association analysis to the evaluation of patients with symptoms suggestive of GORD and we discuss the various methods for symptom analysis that are currently used.

MEASUREMENT OF OESOPHAGEAL ACID EXPOSURE

Since the introduction of 24 hour oesophageal pH monitoring, numerous parameters have been proposed to provide a quantitative description of a patient’s pH profile. Simple (for example, number of reflux episodes, percentage of time with pH below 4, number of reflux episodes lasting more than five minutes) as well as composite (for example, the DeMeester score) parameters are in use. Of these, the percentage of time with pH below 4 is generally accepted as the most useful. Using this parameter, patients can be categorised as having either physiological or pathological (excessive) oesophageal acid exposure. However, several factors limit the usefulness of such a categorisation.

Firstly, significant day to day variation occurs. Wiener et al found that the total per cent of time with pH below 4 varied between two consecutive 24 hour tests by up to a factor of 3.2-fold (218% higher to 69% lower). Overall reproducibility for the diagnosis of GORD was 80%, while reproducibility was much lower for patients with an acid exposure time that was borderline physiological or pathological. Data from other studies confirmed these results and although some would consider a reproducibility of 70–80% to be reasonable, most would argue that it is not acceptable that 20–30% of patients will receive a different diagnosis on repeated testing.

Secondly, the published cut off values to distinguish pathological from physiological reflux differ considerably between various studies. For the parameter time with pH <4 in the upright position, values of 4.6–16.0% were published, and for supine reflux, values of 1.2–11.2%. Furthermore, a physiological increase in acid exposure time occurs with age. These data indicate that distinguishing between pathological and physiological reflux is fraught with a large margin of error.

Once a “diagnosis” of pathological acid exposure has been made, how does this finding relate to reflux symptoms or endoscopic abnormalities? In case of heartburn and regurgitation, a pathological pH profile provides no evidence of whether the patient’s complaints are indeed caused by acid reflux episodes. Howard et al found that patients with an abnormal oesophageal acid exposure often lack a positive correlation between symptom occurrence and reflux episodes.
Klauser et al showed that of several symptoms only heartburn and regurgitation were slightly more frequent in patients with a pathological acid exposure time. In addition, a subset of patients with reflux induced heartburn and regurgitation had a “normal” acid exposure time and can therefore not be identified solely on basis of their acid exposure time. Pathological reflux also appears to be a poor predictor of supraoesophageal symptoms.

Many studies have shown that a high acid exposure time predisposes to erosive reflux disease, and in patients with high degree oesophagitis and Barrett’s oesophagus, almost invariably a pathological pH profile is found. These findings clearly help in our understanding of the pathophysiology of oesophagitis but do not implicate pH monitoring as a useful diagnostic tool. Obviously, ambulatory pH monitoring is an alternative to endoscopy for assessment of oesophageal mucosal injury.

The importance of determining whether a patient’s reflux is pathological or still within physiological limits has often been overstressed and the presence of pathological reflux does not provide evidence about the cause of the symptoms.

SYMPTOM-REFLUX ASSOCIATION ANALYSIS

In addition to quantification of a patient’s reflux profile, 24 hour oesophageal pH recording also provides the opportunity to assess the temporal relationship between the occurrence of reflux and the onset of symptoms. In the early days of oesophageal pH monitoring, the patient’s symptoms were recorded by placing marks on the paper of the ink writer. Nowadays, portable digital dataloggers contain event marker buttons to be pressed by the patient when experiencing symptoms. Although the temporal relation between symptoms and reflux can be convincing during visual inspection of the pH tracings, a quantitative measure of such a relationship is desirable. In the process of quantification of symptom-reflux associations, several criteria should be accurately defined, such as the criteria for the definition of reflux episodes and the length of the time window after a reflux episode during which a symptom is defined as reflux related.

Reflux episodes

An essential step in the analysis of symptom-reflux association is to identify all reflux episodes that occurred during the measurement period. A reflux episode is usually defined as a drop in pH below 4 that lasts for more than 10 seconds. Reflux preceding the onset of symptoms does not necessarily have to reach values below 4. It has been shown that short pH falls of more than 1 unit but not below 4 do not contribute to time with pH <4, but are also capable of eliciting symptoms. Thus incorporation of these pH drops in the symptom association analysis is likely to increase its yield.

Time window

In the early days of analysis of symptom-reflux association, the choice of time window was arbitrary. The aim of a study by Lam et al was to determine the characteristics of the optimal time window. Repetitive symptom association analysis was carried out using time windows of various onsets and durations. One of the indices frequently used for symptom analysis, the symptom index (see below), showed a gradual increase for windows with increasingly early time onset, following a pattern that would be predicted on the basis of Poisson’s theory. However, a relatively sharp cut off point was found at the time window starting at two minutes before pain onset and ending at pain onset. The authors concluded that two minute time windows beginning at two minutes before the onset of symptoms are recommended for symptom-reflux association analysis. It should be borne in mind that this study was performed in patients with non-cardiac chest pain, most of whom had reflux as the cause of their symptoms. The two minute time window may not be optimal for the evaluation of other symptoms such as cough.

Methods for symptom analysis

The next question is how to express associations found between reflux and symptoms in a quantitative way. Ward et al were the first to introduce an index to describe the relation between symptoms and reflux. They proposed the symptom index (SI), which is defined as the percentage of symptom episodes that are related to reflux:

\[
\text{SI} = \frac{\text{Number of reflux related symptom episodes}}{\text{Total number of symptom episodes}} \times 100\%
\]

The distribution of SIs in a population of patients with heartburn appeared to be bimodal and the result of receiver operating characteristic analysis indicated that the optimal threshold for SI is 50%, but thresholds of 25% and 75% have also been suggested. The SI and oesophageal acid exposure do not necessarily correlate. It has been shown that a substantial proportion of patients with a positive SI have physiological acid exposure. Figure 1 shows examples of 24 hour pH recordings with a positive SI in patients with pathological and physiological reflux, respectively.

The major shortcoming of the SI is that this index does not take the total number of reflux episodes into account. The higher the frequency of gastro-oesophageal reflux, the higher the likelihood that a symptom is found to be associated with reflux by chance. Therefore, in patients with frequent gastro-oesophageal reflux who report only one symptom episode during 24 hour pH monitoring, an SI of 100% might well occur by chance.

For this reason the symptom sensitivity index (SSI) was proposed as an additional parameter. SSI is defined as:

\[
\text{SSI} = \frac{\text{Number of symptom associated reflux episodes}}{\text{Total number of reflux episodes}} \times 100\%
\]

SSI values of 10% or higher are considered to be positive. Calculation of both SI and SSI for one and the same pH monitoring test may yield discordant results. However, conclusions can be drawn from discordant test results. For example, the combination of a high SSI and a low SI indicates that the patient’s oesophagus is sensitive to reflux but causes other than acid reflux are likely to contribute to the symptoms.

Both the SI and SSI suffer from the disadvantage that they do not integrate all factors determining the relationship between symptoms and reflux. As a consequence, the probability that SI becomes positive increases with an increasingly high number of reflux episodes and SSI is more likely to be positive when the number of symptom episodes is high.

The Leuven group introduced the binomial symptom index (BSI), a complex formula that calculates the probability that symptoms are related to reflux episodes by summing a number of partial probabilities. Apart from its complexity, BSI suffers from the disadvantage that it only takes into account episodes with a pH drop below 4, whereas it has been shown that drops of more than one pH unit that do not reach
a pH of 4 constitute up to one third of the reflux related symptom episodes. Weusten et al subsequently developed the symptom association probability (SAP), a parameter that expresses the likelihood that the patient's symptoms are related to reflux. It is calculated by dividing 24 hour pH data into consecutive two minute segments. For each of these two minute segments, it is determined whether reflux occurred, providing the total number of two minutes segments with (total R+) and without (total R−) reflux. Then, for each symptom episode, it is determined whether reflux did (S+R+) or did not (S+R−) occur in the preceding two minute period (fig 2). Subtraction of S+R+ from total R+ results in S−R+ and subtraction of S+R− from total R− results in S−R−. A 2×2 contingency table is then constructed in which the number of two minute segments with and without symptoms and with and without reflux are tabulated. Fisher’s exact test is used to calculate the probability (p) that the observed distribution could have been brought about by chance. SAP is calculated as (1−p) × 100%. By statistical convention, SAP ≥ 95% is positive (fig 3). Manual calculation of SAP is cumbersome and time consuming but automatic calculation of this parameter is easy and fast. It should be borne in mind that SAP is a statistical parameter that quantifies the probability that the observed distribution is not brought about by chance. As with all other statistical tests for association, a statistically significant relationship between two parameters does not necessarily imply causality.

Armstrong et al described a method of symptom association analysis that used the Komolgorov-Smirnov test to compare the frequency distribution of the pH values recorded during symptom episodes with the frequency distribution of the pH values during the rest of the measurement. If the two distributions of pH are significantly different, it can be concluded that the patient’s pain is dependent on oesophageal pH. It should be noted however that with this technique the relation between oesophageal pH and symptoms is tested, not the relation between reflux episodes and symptoms. As short pH drops can elicit symptoms but will not have a great effect on pH frequency distributions, the technique is likely to
underestimate the impact of short lived reflux episodes. Furthermore, the statistical test involved is not adequately  
powered if only a few symptoms occur.

Symptom association analysis can also be carried out with  
the use of receiver operating characteristics. Essential to this  
type of analysis are plots in which on the X axis the number  
of symptom associated reflux episodes divided by the total  
number of symptoms (SI) is shown and on the Y axis the  
number of reflux episodes without heartburn divided by the  
total number of reflux episodes is displayed. The area under  
the curve is thus a measure of symptom association. As this  
method makes use of the SI, it suffers from the disadvantages  
herent with this parameter.

Practical issues of symptom analysis

Computerised symptom association analysis is incorporated  
in most commercial ambulatory pH monitoring software. As  
manual inspection of the tracings can sometimes reveal artefacts that are not identified by the computer, it is strongly  
recommanded that such a visual check is carried out. Furthermore, editing of symptom events by the investigator  
is useful as patients can indicate on their symptom diary  
whether they pressed the event marker accidentally or  
whether the event marker was pressed for symptoms  
considered irrelevant in the context of 24 hour oesophageal  
pH testing.

Thorough instruction of the patient is one of the most  
portant prerequisites of a successful 24 hour pH test. It has  
to be stressed that the patient has to press the event marker  
button each time a symptom is perceived. A diary on which  
the patient can fill in the nature of every symptom makes it  
possible to perform symptom analysis for different symp- 
toms. As a result, a patient can have a positive SI, SSI, and  
SAP for heartburn, but not for cough, and vice versa. Furthermore, the patient has to be instructed not to reduce  
his normal daily activity and to consume his usual meals in  
order to elicit symptoms. Obviously, a 24 hour pH study  
during which no symptoms occur does not contribute to the  
evaluation of these symptoms.

CONTRIBUTION TO THE DIAGNOSTIC WORKUP

Before the advent of 24 hour pH monitoring, the relation  
between symptoms and acid could only be investigated using  
provocation tests such as the acid perfusion test according to  
Bernstein and Baker. In this test, hydrochloric acid and  
saline solutions are instilled into the oesophagus of the  
patient who is blinded for the sequence of instillation. The  
test is considered to be positive when acid infusion  
reproduces the patient’s symptoms and saline does not. Several comparative studies showed that oesophageal pH  
monitoring is superior to provocation testing in identifying  
the cause of patients’ symptoms. Discordance between the  
acid perfusion test and pH monitoring implied that other  
factors of gastro-oesophageal reflux besides the acidity of the  
refluxate were responsible for symptom generation. Comparison of the two tests also led to the conclusion that  
symptomatic gastro-oesophageal reflux, abnormal oesopha- 
geal acid exposure, and mucosal acid sensitivity are three  
separate although related aspects of GORD. As illustrated in  
figs 1 and 4, there is a subset of patients with a normal  
oesophageal acid exposure time who exhibit a positive  
temporal association between symptoms and oesophageal  
acid exposure. Whereas in the Rome II classification  
this frequently encountered condition is classified as func- 
tional heartburn, the Rome III consensus will incorporate it  
into GORD. Watson et al have shown that in a group of patients with  
heartburn and regurgitation and physiological oesophageal  
acid exposure, those with a positive SI showed a considerably  
better symptomatic response to omeprazole than those with a  
negative SI, suggesting that SI is a good predictor of response
to proton pump inhibitor (PPI) therapy. A recent study by Taghavi et al compared the predictive value of symptom association analysis for outcome after a short term treatment with a high dose of omeprazole (PPI test). Whereas the positive predictive values of SI, SSI and SAP were high, the negative predictive values were relatively low. This implies that patients may react satisfactorily to treatment with PPIs while there is no demonstrable relation between the onset of their symptoms and reflux. Whereas the authors of this study interpret this finding as imperfection of the association indices, it seems more likely that false positive responses to PPIs played a role. Diaz et al also showed that SAP is a good and independent predictor of the success of antireflux surgery; failure to reach a positive SAP value for the principal symptom reduced the likelihood of responding favourably to antireflux surgery. It has also been suggested that patients with physiological acid exposure and a positive SI benefit as much from antireflux surgery as patients with pathological acid exposure.

Oesophageal pH monitoring is not only used for evaluation of typical oesophageal reflux symptoms such as heartburn and regurgitation, but also for evaluation of atypical reflux symptoms. Whereas the acute onset of heartburn and regurgitation makes it possible to investigate the temporal correlation with reflux events, this is not possible with symptoms lacking a crisp onset, such as globus sensation and hoarseness. In the case of unexplained chronic cough however, symptom association analysis showed a temporal relation between acid reflux and cough in one third of patients. Combined oesophageal pH and pressure monitoring makes it possible to distinguish between reflux-cough and cough-reflux sequences, as coughs spells can be recognised manometrically. Making this distinction is important because reflux induced coughing should be treated differently from cough induced reflux.

Studies using symptom association analysis of combined 24 hour pH and pressure signals in patients with non-cardiac chest pain have shown that an oesophageal origin of the pain can be identified in a small subset of patients only. Combined oesophageal pH and pressure monitoring makes it possible to distinguish between reflux-cough and cough-reflux sequences, as coughs spells can be recognised manometrically. Making this distinction is important because reflux induced coughing should be treated differently from cough induced reflux.

Figure 4 Diagram of the various elements of gastro-oesophageal reflux disease (GORD). When GORD is defined as either mucosal damage and/or symptoms induced by gastro-oesophageal reflux, all GORD patients fall in the circles “oesophagitis” and/or “positive symptom-reflux association”. Patients with symptoms of heartburn and/or regurgitation but no oesophagitis and no relation between symptoms and reflux on pHmetry do not fall under the definition of GORD. Patients who have neither pathological acid exposure nor oesophagitis but do have a positive symptom association are classified by the Rome II criteria as functional heartburn.

Figure 5 Combined oesophageal pH-impedance measurement. There is a decrease in impedance starting in the most distal impedance segment (arrow) and moving in a proximal direction, indicating liquid reflux. No drop in pH below 4 is measured and thus this episode would not have been identified on pHmetry alone. The symptom event marker (“pain”) is pressed by the patient less than two minutes after the onset of this reflux episode, classifying this symptom as non-acid reflux associated.
diagnosis contributes significantly to the patient’s well being.30

FUTURE PROSPECTS

Although acid gastro-oesophageal reflux is responsible for symptoms of heartburn and regurgitation in a large proportion of patients, these symptoms sometimes occur without any evidence of acid reflux. Several studies have shown that besides acid, heartburn can also be elicited by hyperosmolar solutions, certain foods, bile acids, oesophageal contractions, and oesophageal distension.31-33

Studies using the Bilitec 2000, a fibreoptic spectrophotometric probe that makes use of the absorbance spectrum of bilirubin for the detection of duodenogastro-oesophageal reflux, showed that reflux of bile can only be held responsible for a minor proportion of symptoms.34-35 Another study calculated that only 7% of patients with GORD had a positive SAP for bile reflux, as detected by the bilirubin absorbance technique, which is 2% above the percentage that is expected to be found by pure chance.36 However, reflux detection using the bilirubin absorbance technique has several limitations. The most important of these is that clearance of material trapped in the probe gap takes much longer than oesophageal clearance of bile. As a consequence, individual reflux episodes cannot be detected with sufficient accuracy. Furthermore, the signal-response relationship of this device is distorted in an acid environment, a situation that often occurs in the oesophagus of patients with GORD.37

Intraluminal impedance monitoring appears to hold much more promise for symptom association analysis. With this technique, movements of liquid and air within the oesophageal lumen, either in an antegrade or retrograde direction, can be monitored. Using impedance monitoring, reflux can be detected independently of the acidity of the refluxate.38 The temporal resolution of reflux detection by intraluminal impedance monitoring is high, which makes it suitable for symptom association studies (fig 5). Using this technique, Vela et al showed that heartburn and regurgitation in patients off acid secretion inhibiting medication were caused by both acid and non-acid reflux, while during acid suppression non-acid reflux was responsible for symptom generation.39 In patients using acid suppression, addition of impedance monitoring to pH monitoring leads to a higher SI, SSI, and SAP.40-42 Preliminary data also suggest that a positive SI for non-acid reflux predicts a good symptomatic response to fundoplication.43

As already mentioned, acid reflux can be the cause of chronic cough. A recent study showed that combined pH-impedance monitoring can identify reflux as a causative agent for cough in more patients compared with pH monitoring alone.44

One of the restrictions of 24 hour reflux monitoring is that the frequency of symptoms varies from day to day, which implies that a single 24 hour period may not be representative. Recently, a wireless system for ambulatory pH monitoring has become available. A capsule containing a pH electrode and a radiotransmitter placed in the distal oesophagus transmits data to an external datalogger.45 It appears that pH monitoring carried out with this technique is tolerated better by patients than conventional catheter based pHmetry, and measurement time can therefore be increased. It was shown that 48 hour ambulatory pH monitoring doubles the SI and SAP in patients with atypical reflux symptoms in comparison with 24 hour monitoring.46

CONCLUSIONS

Reflux of gastric content may lead to symptoms and mucosal inflammation. While the latter can be detected endoscopically, symptom-reflux association analysis is the only method that can adequately identify reflux as the cause of short lived symptoms with sudden onset, such as heartburn, regurgitation, chest pain, and cough. Such analysis should therefore be included in every 24 hour pH study evaluating these complaints. Indices that express the strength of the symptom-reflux association numerically, such as the SI and SAP, constitute the most important information that can be obtained from a 24 hour pH study.

ACKNOWLEDGEMENTS

AJ Bredenoord is supported by a clinical research trainee grant from Janssen Pharmaceuticals, Tilburg, the Netherlands.
REFERENCES


How can a migraine cause faecal incontinence?

Clinical presentation
A 31 year old woman presented with vaginal faecal discharge and anal pain. Her past medical history was significant for a 20 year history of severe migraine treated with ergotamine based suppositories. Two months prior to admission, she started reporting constipation and having anal pain on defecation. Three weeks prior to admission, she reported a vaginal faecal discharge initially on defecation but later even on passing flatus. Her systemic evaluation showed no other sign or symptom. Physical examination and laboratory data were all normal. Rectal examination revealed a painful circular stenosis 3 cm cephalad to the dentate line. Rectoscopy was performed (figs 1–3).

Question
What does rectoscopy show (figs 1–3)?
See page 1824 for answer
This case is submitted by:

K Honein, J Boujaoude, M Chemaly, C Yaghi, R Karam, R Sayegh
Department of Gastroenterology, Hotel-Dieu de France Hospital, Saint-Joseph University, Beirut, Lebanon

doi: 10.1136/gut.2005.068627
Symptom association analysis in ambulatory gastro-oesophageal reflux monitoring

A J Bredenoord, B L A M Weusten and A J P M Smout

Gut 2005 54: 1810-1817
doi: 10.1136/gut.2005.072629

Updated information and services can be found at:
http://gut.bmj.com/content/54/12/1810

These include:

References
This article cites 63 articles, 14 of which you can access for free at:
http://gut.bmj.com/content/54/12/1810#BIBL

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.

Topic Collections
Articles on similar topics can be found in the following collections
Gastro-oesophageal reflux (351)

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/