LETTERS TO THE EDITOR

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Squawks in pneumonia

Squawks are short inspiratory wheezes that have been described in hypersensitivity pneumonitis and other fibrotic disorders. Little attention has been paid to the fact that they also occur in patients with pneumonia. In the course of studying the correlation of automated lung sound analysis with disease states in patients at a community teaching hospital, we noticed that squawks appeared to be more common in patients with pneumonia than we expected. We therefore examined the occurrence of squawks more systematically in 500 subjects who had been examined with a multichannel lung sound analyser (Stethographics Model STG-1602), as previously described.1 Seventy eight of the subjects in this population had a clinical diagnosis of pneumonia. All participants had been asked to breathe more deeply than normal with their mouths open. Two 20 second samples were taken. The Institutional Review Board of the Faulkner Hospital approved the study. Two experienced observers, blinded to the clinical diagnosis, used playback and waveform displays to identify squawks. They were defined according to the criteria initially described by Earis et al and adopted by CORSA.2 3 All channels from each subject were replayed and the waveforms of the data in the time domain were simultaneously examined. Only those sounds which fit both the auditory and waveform characteristics were considered to be squawks for the purposes of this study.

Squawks were present in 12 of the 78 patients with pneumonia and in none of 224 patients considered to have no significant lung disease. They were found in four of 18 patients with interstitial pulmonary fibrosis, two of 41 patients with bronchial asthma, one of 79 patients with COPD, and in none of 86 patients with congestive heart failure. We also noted squawks in a patient with radiation pneumonitis and in one of the two patients in our database with hypersensitivity pneumonitis.

In nine of the 12 patients with pneumonia the squawks were in the same location as the radiological opacifications. In one patient the squawk was in a different location and in another the chest radiograph did not show evidence consistent with pneumonia until 4 days after the squawk was detected. In the remaining patient a squawk was heard over the left posterior mid chest. The portable chest radiograph was interpreted as technically suboptimal due to the patient’s body habitus. The clinicians caring for this patient made a diagnosis of pneumonia and treated him accordingly.

Interestingly, one patient who clearly had congestive heart failure on a number of occasions had a squawk when we examined him. On re-examining his record he also had a diagnosis of pneumonia on that day. Similarly, there were two patients with squawks (one with COPD and one with asthma) who, on the day that the squawks were noted, had acute febrile illnesses consistent with pneumonia. In one patient the presence of a squawk led to reinterpretation of the chest radiograph as showing an area of opacification consistent with pneumonia.

The squawks in this study all had a distinctive sound that is readily distinguished from crackles, rhonchi, rubs, and most wheezing noises. Occasionally wheezes can be short and have a similar sound, but this occurs rarely as an isolated finding in inspiration. The squawks in this study all had sinusoidal waveforms as illustrated in the time-amplitude plot shown in fig 1. The mean (SD) duration of these sounds was 64 (49) ms (range 16–228) and the mean (SD) frequency was 425 (110) Hz (range 200–667). These findings are similar to those of Earis et al.1

When a squawk is accurately identified, the question arises—what does it mean? In a patient who is not acutely ill, investigations to rule out hypersensitivity pneumonitis and the other fibrotic conditions mentioned above should be considered. In a patient with a clinical picture consistent with pneumonia, the presence of a squawk offers some objective evidence to support the diagnosis. It would seem reasonable to suggest that the patient should be followed up to be sure that the squawk disappears when the acute illness resolves to exclude the possibility that the acute illness was the mode of presentation of a chronic pulmonary disorder. Squawks can be helpful in providing evidence for pneumonia in lung areas where radiological visualisation may be suboptimal, such as below the dome of the diaphragm or in the retrocardiac region.

In summary, short inspiratory wheeze-like sounds are found in pneumonia. Other conditions that can cause them are chronic restrictive disorders but these are relatively uncommon compared with pneumonia. When there is no evidence of these restrictive disorders and an acute syndrome consistent

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Figure 1  Time-amplitude plot of a sound recorded at the lung bases posteriorly (channel 14) in a patient with pneumonia. Waveforms are presented in both the unexpanded (A) and expanded (B) modes. (A) The unexpanded waveform shows one full breath. The solid bars above the unexpanded waveform mark the respiratory cycle (the light bar indicates inspiration and the dark bar indicates expiration). The normal inspiratory sound can be seen as having almost random waveform fluctuations. Fine crackles (c) look like spikes on an unexpanded waveform. A squawk is present at the end of inspiration. One fine crackle and a wheeze are present during expiration. (B) The expanded squawk waveform exhibits monophonic sinusoidal fluctuations lasting approximately 60 ms. (C) The squawk sound in the frequency domain shows a single peak at 600 Hz. In this patient similar squawks appeared in three consecutive breaths during 20 s of deeper than normal breathing. They occurred approximately in the same location on the chest and at the same time in the respiratory cycle. These observations were typical of the squawks detected in our study.

FFT = fast fourier transform.  

References


with respiratory infection is present, squawks can provide relatively specific—although not very sensitive—evidence of pneumonia.

R Paciej, A Vyshedskiy, D Bana, R Murphy Faulkner/Brigham and Women Hospitals, Boston, MA, USA

Correspondence to: Dr R Murphy, 1153 Centre St, Suite 4990, Boston, MA 02130, USA; murphy@faulknerhospital.org

Supported in part by grants from NIH SBIR (1R43HL70480-01) and from Stethographics Inc.

References

BTS guidelines for the management of pleural infection

We have read the BTS guidelines for the management of pleural infection1 and are concerned about the proposed antibiotic choices for the initial treatment of culture negative or pending pleural infection. Section 2.8 of the text and table 2 detail the antibiotic choices but, in our opinion, leave considerable gaps in antibacterial cover against the likely pathogens. In particular:

- Amoxicillin (text) is not reliably active against Staphylococcus aureus.
- Clindamycin has no activity against Gram negative aerobic organisms (especially not Haemophilus influenzae as mentioned in the text).
- Benzyl penicillin (table) rarely now has activity against Staphylococcus aureus and we suggest that relying on ciprofloxacin is unwise. In addition, this combination will not cover any anaerobic bacteria.
- We do not consider chloramphenicol is an appropriate agent to use in this category of patients in view of the serious side effect profile.
- Third generation cephalosporins such as cefazidine and cefotaxime have unreliable activity against many anaerobic bacteria.
- Pneumococci are considerably less susceptible to cefazidine than to other cephalosporins and penicillins; the policy (table), however, suggests its use as a single agent.
- Piperacillin (text) is no longer available in the UK except in combination with a β-lactamase inhibitor.

We suggest that the antibiotic choices in the BTS guidelines for the management of pleural infection should be changed to the following:

- For community acquired pleural infection, either cefuroxime + metronidazole or co-amoxiclav or (for the penicillin/cephalosporin allergic individuals) clindamycin + ciprofloxacin, all administered intravenously. Oral treatment choices would be co-amoxiclav or clindamycin + ciprofloxacin.
- For hospital acquired infection, clinicians should seek guidance from the local microbiologists but the following choices would be reasonable in the interim: piperacillin + tazobactam or cefotaxime-ceftriaxone + metronidazole or meropenem.

Relating to the initial diagnosis of pleural infections, we were also concerned that mycobacteria were mentioned only once in the article. Pleural fluid has a poor yield for diagnosis of tuberculosis and more emphasis should have been placed on the routine use of pleural biopsy for histology and culture of tuberculosis which has much higher diagnostic rates. The algorithm should include the investigation of pleural tuberculosis.

In conclusion, we would commend the inclusion of a medical microbiologist in discussions leading to guidelines dealing with the diagnosis and treatment of infections.

H S A Roberts
Consultant Microbiologist, Princess Royal University Hospital, Farnborough Common, Orpington, Kent BR6 8ND, UK; honor.roberts@bramleyhospitals.nhs.uk

*On behalf of the Outer South East London Microbiology Group: M Viagappan, University Hospital, Lewisham; A Mackay, Queen Elizabeth Hospital, Woolwich; A Teall, Queen Elizabeth Hospital, Woolwich; H Roberts, Princess Royal University Hospital, Bromley; J Kinsit, Queen Mary’s Hospital, Sidcup; A Fowler, Queen Elizabeth Hospital, Woolwich; A Shaw, Darent Valley Hospital.

References

Authors’ reply

We would like to thank Dr Roberts and colleagues for their interest in the BTS guidelines on pleural infection1 and for their thoughtful letter. In our view, guidelines (particularly the first set in an area) exist partly to stimulate a debate which subsequently better informs clinical practice.

Since the advisory regimens in the guidelines were first drafted (and they are “advisory” and to be used in line with local microbiological advice), the microbiology of pleural infection has become more clearly defined (not least by the joint BTS/MRC trial in pleural infection). This same work is also partly to stimulate a debate which subsequently better informs clinical practice (not least by the joint BTS/MRC trial in pleural infection). This same work is also partly to stimulate a debate which subsequently better informs clinical practice.

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Authors’ reply

We thank Dr Cotes for his interest in our work, and his thoughtful comments. We did not include the suggested analysis as our purpose was to address the utility of a single bronchodilator reversibility test which is the way this is applied in routine practice. We agree that regression to the mean is a real possibility, but also believe that physiological variation in baseline airway calibre contributes to the “noise” in this particular signal. Unfortunately the number of times this test must be undertaken to minimise these confounding effects is not clear, but is likely to be too large to be possible in most clinics. Hence our caution about interpreting the significance of small changes in spirometry recorded on one visit.

P M A Calverley
Department of Medicine, Clinical Sciences Centre, University Hospital, Aintree, Liverpool L9 1AL, UK
pma@liverpool.ac.uk

Reference

Prednisolone response in patients with COPD

Our interpretation of the important data on corticosteroid responsiveness in chronic obstructive pulmonary disease (COPD) presented by Burge et al is quite different from that of the authors. They have shown that those with an improvement in forced expiratory volume in 1 second (FEV₁) of >20% following short term treatment with prednisolone had a more significant reduction in exacerbation frequency with longer term treatment with inhaled fluticasone than those without. Thus, arguably, the only important positive clinical outcome of long term treatment of COPD with inhaled corticosteroids is largely confined to those who are identified by a positive response to prednisolone. This is a powerful validation of the suggestion in current guidelines that decisions on the use of long term inhaled corticosteroid therapy are based on a careful assessment of the response to short term treatment with prednisolone.

Clinicians will continue to use short term prednisolone trials in patients with unclassified, apparently fixed, airflow obstruction since this is the only practical way to identify the minority with markedly corticosteroid responsive disease. The data presented by Burge et al are important in that they show that smaller responses to prednisolone can provide additional useful clinical guidance. They also suggest that the mechanism of the bronchodilator response to prednisolone might have features in common with the mechanism of inhibition of COPD exacerbations with corticosteroids. We have recently shown that the improvement in FEV₁ and quality of life scores seen with prednisolone compared with placebo in COPD increases progressively with increasing eosinophilic airway inflammation, and is associated with amelioration in eosinophilic airway inflammation as reflected by a sixfold reduction in the induced sputum eosinophil count. Future studies should investigate whether treatment targeting eosinophilic airway inflammation in COPD results in a reduction in exacerbations, as it has clearly been shown to do in asthma.

I D Pavord, R Siva, C E Brightling
Institute for Lung Health, Glenfield Hospital, Leicester, UK
Correspondence to: Dr I D Pavord, Institute for Lung Health, Glenfield Hospital, Leicester, UK, ian.pavord@ulh-tr.nhs.uk

Authors’ reply

We thank Dr Pavord and colleagues for their letter. We were asked to add the section relating the effect of prednisolone response to fluticasone related exacerbation prevention by the papers’ referees. We originally omitted this as there was no relationship between the effect of exacerbation reduction and predni- solone response when prednisolone response was expressed as a continuous variable (p = 0.84), and we were worried that the analysis based on responders and non-responders would result in misinterpretation, which unfortunately has been the case. The main conclusion from the paper is that any separation of the patients into responder and non-responder groups is statistically flawed. The response is unimodally distributed and individual changes between two visits (that is, before and after prednisolone) suffer from the regression to the mean effect. We showed good evidence that this was in fact occurring, with “responders” declining by a mean 127 ml in the 4 weeks before prednisolone, while those in whom forced expiratory volume in 1 second (FEV₁) was reduced by >20% after prednisolone had a 47 ml increase in FEV₁ in the preceding 4 weeks.

The most likely explanation for the spurious relationship between exacerbation reduction and prednisolone response related to confounding by a low FEV₁. Responders were classified as those with a 20% increase in FEV₁, as a percentage of the starting value favouring a “response” in those with the lowest FEV₁ (a change from 0.8 to 0.96 l in the lowest group) and exacerbations are also more frequent in those with a lower FEV₁. A clinical decision made on a steroid trial is not a reliable, reproducible, or valid way of separating out “responders” and so cannot be recommended.

P S Burge
Birmingham Heartlands Hospital, Bordesley Green East, Birmingham, UK

References

Breathing exercises in asthma

It was a pleasure to read a report of a well conducted trial into a complementary treatment for asthma. This is a subject which attracts considerable media attention. The authors of this report and the author of an accompanying editorial both introduce their articles with a reference to “a third of respondents in a recent asthma survey having tried one or more breathing techniques to relieve symptoms”. While their quote is correct, the report to which they refer is of a survey of members of the UK National Asthma Campaign. Such a membership may not be typical of those with asthma. In a more recent study of a stratified cross section of the asthma population we found only 6% of the study population to be current users of complementary therapies. That use was greatest among those who expressed most concern regarding their current medication.

M R Partridge
Imperial College London, NHU, Charing Cross Hospital Campus, London W6 8RF, UK; m.partridge@imperial.ac.uk

References

A1-Antitrypsin deficiency: paper by Stolk et al

In the paper entitled “Correlation between annual change in health status and computer tomography derived lung density in subjects with A1-antitrypsin deficiency” by J Stolk et al which appeared in the December 2003 issue of Thorax on pages 1027–30, the affiliation address for J Stolk, W H Ng and K F Rabe on page 1030 and the correspondence address on page 1027 should have been Department of Pulmonology, Leiden University Medical Center and not Department of Radiology and Division of Image Processing. The publishers apologise for this error.

P S Burge
Birmingham Heartlands Hospital, Bordesley Green East, Birmingham, UK

P M A Calverley
University Hospital, Aintree, Liverpool, UK

P W Jones, S Spencer
St George’s Hospital Medical School, London, UK

Correspondence to: Professor P S Burge, Birmingham Heartlands Hospital, Bordesley Green East, Birmingham B9 5ST, UK; sherwood.burge@heartsol.wmids.nhs.uk

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I D Pavord, R Siva and C E Brightling

Thorax 2004 59: 179
doi: 10.1136/thx.2003.019414

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