Pulmonary puzzle

A cavitating pulmonary lesion with eosinophilia

A middle-aged individual of Ethiopian origin attended the emergency department with haemoptysis. The patient had no other significant medical problems and had not travelled for several years. The patient drank moderately and did not smoke but later admitted to chewing khat (a narcotic leaf) frequently over the past 6 months.

There were no significant findings on physical examination. Renal and liver function tests and C reactive protein were within normal limits, but a peripheral blood eosinophilia was noted (2.1×10^9/l). Serological tests for HIV, hepatitis B and C were negative. A poorly defined opacity was seen in the right lung on chest x-ray, and a subsequent CT scan showed an area of consolidation with cavitation (see figure 1). An incidental finding of low-attenuation subcapsular lesions in the part of the liver imaged on this scan was also noted.

The patient was referred to the chest clinic for investigation of possible tuberculosis, but sputum cultures were negative for mycobacteria. On review two months later, the haemoptysis and the radiological findings had worsened. The eosinophil count remained raised. An abdominal CT scan revealed multiple low-density tracking liver lesions (figure 2). A serological test was performed.

Figure 1 Contrast-enhanced CT chest: consolidation seen in the right lower lobe, with the suggestion of early cavitation (arrow).

Figure 2 Axial contrast-enhanced CT abdomen: There are multiple low-attenuation lesions seen in the subcapsular region (arrows) extending towards the porta hepatitis. Mild intrahepatic bile duct dilatation is seen (arrowheads).

QUESTION

See page XXX

Alastair Charles McGregor, Nyree Griffin, Ronan A Breen, William Newsholme

1 Department of Infection, St Thomas' Hospital, London, UK; 2 Department of Radiology, St Thomas's Hospital, London, UK; 3 Thoracic Medicine, St Thomas Hospital, London, UK

Correspondence to Dr Alastair Charles McGregor, Department of Infection St Thomas’s Hospital Westminster Bridge Road, London SE1 7EH, UK; alastairmcgregor@yahoo.co.uk

Contributors All authors have contributed to drafting this case.

Competing interests None.

Provenance and peer review Not commissioned; externally peer reviewed.
ANSWER

From question on page above

DIAGNOSIS: PULMONARY FASCIOLIASIS

Fascioliasis occurs following ingestion of aquatic plants contaminated with encysted metacercariae of *Fasciola* spp. The parasites migrate to the liver after hatching in the bowel and grow into adult flukes. Egg production commences after several months, with excretion via the biliary tract into the faeces. A developmental cycle in the intermediate host (a freshwater snail) then results in contamination of further plants by metacercariae and a new cycle of infection.

*Fasciola* is found worldwide, and outbreaks associated, particularly, with the consumption of watercress have been well documented. An association with the chewing of *khat* has also been noticed.1 *Khat* is the leaf of the shrub *Catha edulis*. It has a mild narcotic effect when chewed and, although not illegal in most European states, its use is limited to certain ethnic communities.2 *Khat* may become contaminated when sprinkled with water to keep it fresh during transport.3

*Fasciola* usually affects only the liver but, as with all flukes, aberrant migration can lead to disease in a variety of organs. There are reports of *Fasciola* causing pleural effusions, pneumothorax and pulmonary lesions.4 5 Other flukes, such as *Paragonimus* and *Schistosoma* can also cause lung lesions. When there is an epidemiological risk, it is important to consider these unusual pathogens to avoid subjecting patients to unnecessary antituberculous therapy or surgery. In our case, the diagnosis was made on serological testing of serum (*Fasciola IFAT 1:128*) and review of the liver imaging, which showed tracking lesions suggestive of *Fasciola* migration. The patient was treated with 700 mg of triclabendazole twice. Two months later, the chest x-ray abnormalities had improved, the eosinophil count was within normal limits and the haemoptysis had completely resolved.

Thorax 2012; 67:2. doi:10.1136/thoraxjnl-2012-201848

REFERENCES

A cavitating pulmonary lesion with eosinophilia

Alastair Charles McGregor, Nyree Griffin, Ronan A Breen and William Newsholme

Thorax  published online July 21, 2012

Updated information and services can be found at:
http://thorax.bmj.com/content/early/2012/07/20/thoraxjnll-2012-201848

These include:

References
This article cites 5 articles, 1 of which you can access for free at:
http://thorax.bmj.com/content/early/2012/07/20/thoraxjnll-2012-201848#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

Radiology (diagnostics) (812)  
Hemoptysis (80)  
Lung infection (97)  
Pneumonia (infectious disease) (579)  
Pneumonia (respiratory medicine) (562)  
TB and other respiratory infections (1273)  
Emergency medicine (185)  
Drugs: infectious diseases (968)  
HIV/AIDS (194)  
Journalology (123)

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/