Article 3



Images in clinical medicine



A woman with recurrent respiratory tract infections

Petros Ioannou

Corresponding author: Petros Ioannou, Department of Internal Medicine, University Hospital of Heraklion,

Panepistimiou, Iraklio 715 00, Greece. p.ioannou@med.uoc.gr

Received: 16 Nov 2020 - **Accepted:** 23 Nov 2020 - **Published:** 26 Nov 2020

Keywords: Kartagener, primary ciliary dyskinesia, dyskinetic cilia syndrome

Copyright: Petros Ioannou et al. Pan African Medical Journal (ISSN: 1937-8688). This is an Open Access article distributed under the terms of the Creative Commons Attribution International 4.0 License (https://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Cite this article: Petros Ioannou et al. A woman with recurrent respiratory tract infections. Pan African Medical Journal. 2020;37(278). 10.11604/pamj.2020.37.278.27013

Available online at: https://www.panafrican-med-journal.com//content/article/37/278/full

A woman with recurrent respiratory tract infections

Petros Ioannou^{1,&}

¹Department of Internal Medicine, University Hospital of Heraklion, Panepistimiou, Iraklio 715 00, Greece

[&]Corresponding author

Petros Ioannou, Department of Internal Medicine, University Hospital of Heraklion, Panepistimiou, Iraklio 715 00, Greece

Image in medicine

A 76-year-old caucasian woman who has been repeatedly hospitalized with recurrent respiratory tract infections, presented to the emergency department with fever and productive cough during the last 24 hours. Physical examination revealed tachypnea and crackles at the lower right lobe. The chest X-ray revealed an infiltration at the right lower pulmonary field and dextrocardia. Laboratory data revealed leukocytosis and increased C-reactive protein. The electrocardiogram revealed sinus rhythm, right axis deviation, inverted P waves in lead I, dominant R waves in V1 and V2 and dominant S waves in V3-6. Kartagener syndrome is an autosomal recessive disorder also known as "primary ciliary

Article 3



dyskinesia" and "dyskinetic cilia syndrome". It involves clinical features summarized in a triad including bronchiectasis, chronic sinusitis and dextrocardia/situs inversus. Kartagener syndrome affects equally males and females, with a prevalence of approximately 1 in 10,000-30,000. The underlying cause is impaired ciliary mobility resulting in reduced mucociliary clearance and mucus accumulation causing strong predisposition to recurrent bacterial respiratory tract infections that start during early childhood. Progressively, manifestations such as sinusitis and otitis media

increasingly appear along with respiratory tract infections and eventually bronchiectasis develops. manifestations usual clinical include Less telecanthus, infertility in males and subfertility in females. Differential diagnosis includes respiratory allergy, cystic fibrosis, immune disorders and a1-trypsin deficiency. Screening starts with mucociliary clearance measurements but the Kartagener syndrome diagnosis is confirmed with the presence of anatomic abnormalities of the cilia in biopsies of bronchial and nasal mucosa with the use of electron microscopy.

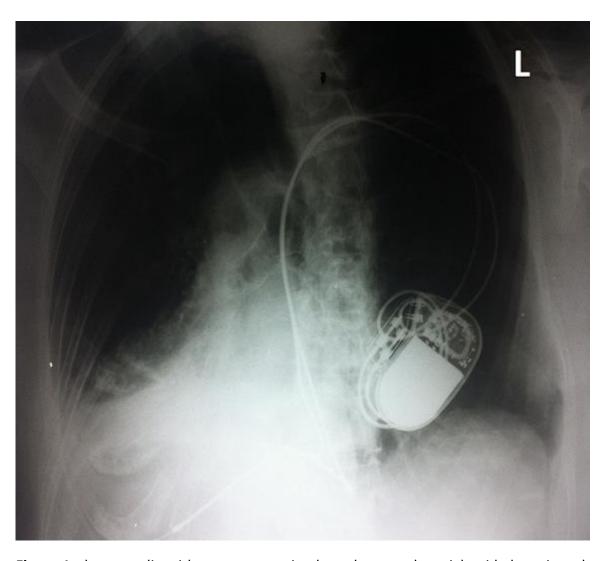


Figure 1: dextrocardia with a permanent implanted pacemaker, right-sided aortic arch suggesting situs inversus; consolidation of the right lower pulmonary field suggesting lower lobe pneumonia