A cyst within a cyst

Kist içinde kist

A 31-year-old woman was referred to our hospital for recurrent syncopal attacks. Physical examination was normal except for a 2/6 systolic murmur at pulmonary area. Her chest X-ray and electrocardiogram were also normal. The laboratory workout showed leukocytosis and hypereosinophilia. On transthoracic echocardiography, a cystic lesion with a smooth, distinct border was found next to the right ventricular outflow tract (RVOT) (Fig. 1A, Video 1. See corresponding video/movie images at www.anakarder.com). Inside the cystic structure, a smaller cyst of 1 x 1 cm dimension was seen (Fig. 1B, Video 2. See corresponding video/movie images at www.anakarder.com). The lesion was compressing the RVOT, and on Doppler examination, a gradient of 30 mmHg was measured at the RVOT (Fig. 1C, Video 2. See corresponding video/movie images at www.anakarder.com). The same lesion was confirmed by 2-D transesophageal echocardiography (TEE) (Fig. 1D, E, Video 3-4. See corresponding video/movie images at www.anakarder.com) but 3-D TEE study could not provide further details about the nature of the cyst (Fig. 1F, Video 5. See corresponding video/movie images at www.anakarder.com). Being endemic in Turkey, Echinococcosis was suspected because of the characteristic appearance of the cystic lesion. On cardiac magnetic resonance imaging, the cyst was found to possess a smooth border and no invasion into neighboring structures was noted (Fig. 1G, H). On surgery, median sternotomy was done, and the cyst was found to be situated on the right ventricle under the pericardium (Fig. 1I). The wall was punctured and hypertonic saline and iodine was injected. The same procedure was also applied to the inner cyst, and the two cysts were removed together. Treatment with albendazole was continued for 4 weeks after the operation. The postoperative follow-up was uneventful.

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Available Online Date/Çevrimiçi Yayın Tarihi: 10.01.2012
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A giant mediastinal carcinoid tumor that compresses the pulmonary artery and vein

Pulmoner arter ve vene bassi yapan dev mediyastinal karsinoid tümör

A 39-year-old man was admitted with recently initiated complaints of chest pain and dyspnea. He had a history of hypophysis and thyroid operation after being diagnosed with MEN type 1 syndrome including prolactinoma, parathyroid adenoma, gastrinoma and a nonfunctional adenoma on suprarenal glands 3 years before admission. Chest X-ray (Fig. 1, 2) displayed a large mediastinum, computed tomography showed 13x9x11 cm lobulating, heterogeneous hypodense mass in superior mediastinum that compressed left superior pulmonary vein (Fig. 3). There were no metastatic masses in liver or in any other localization.
Transthoracic echocardiography demonstrated a huge mass that compressed pulmonary artery resulting in 25 mmHg transpulmonary gradient (Fig. 4, 5). Pathological evaluation showed parathormone negative, chromogranin positive stained neoplastic cells that eventually proved to be a carcinoid tumor.

Carcinoid tumors are the most common neuroendocrine tumors. They grow insidiously and usually do not cause any symptom. Our case was an extreme sample of carcinoid tumor which was extended to a massive size that caused large vessel compression, and eventually treated surgically without complication.

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Available Online Date/Cevrimiçi Yayın Tarihi: 10.01.2012

Fig. 4. Transthoracic echocardiography view of a huge hyperechogenic mass
Fig. 5. Continuous wave Doppler examination revealing a maximal 25 mmHg of transpulmonary gradient due to the tumoral compression

Giant pulmonary artery aneurysm due to chronic pulmonary embolus associated with pulmonary hypertension

Aneurysm of the pulmonary artery (PAA) is a rare pathology with unknown natural history. The main causes of PAA are pulmonary hypertension (PHT) secondary to pulmonary embolus or congenital heart diseases with left-to-right shunts. We report a case of giant PAA due to chronic pulmonary embolus associated with PHT in an elderly patient.

An 83-year old male with a known history of multiple episodes of deep venous thrombosis, chronic pulmonary embolism associated with PHT and chronic atrial fibrillation in last five years was admitted with NYHA-3 exertional dyspnea. The physical examination revealed orthopnea, jugular venous distention, ascites and bilateral pretilial edema. Electrocardiography revea-