CASE REPORT OLGU SUNUMU



# Carbamazepine-Induced Lymphocytic Interstitial Pneumonitis

# Karbamazepine Bağlı Lenfositik İnterstisyel Pnömoni

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#### **Abstract**

A twenty-seven-year-old male was admitted with cough, sputum production, dyspnea and weight loss. Postero-anterior chest x-ray showed bilateral, diffuse reticulonoduler infiltration. He had received tuberculosis treatment in the past and he had been on carbamazepine for 8 years for a seizure disorder. Three sputum smear examinations for acid-fast bacilli were negative. Pulmonary function tests revealed FVC of 2.35L (51 %), FEV1 of 2.12 L (54 %), FEV1/FVC of 90 %, DLCO of 39 % and DLCO/VA of 75 %. Fiberoptic bronchoscopy was normal. Transbronchial lung biopsy and bronchial washing were nondiagnostic. Right minithoracotomy and open lung biopsy were performed. Pathologic examination revealed diagnosis of lymphocytic interstitial pneumonitis. Carbamazepine was withdrawn after neurological consultation. The patient was clinically and functionally well two months later. Computed tomography of the thorax was completely normal. The patient was considered to be compatible with carbamazepine-induced lymphocytic interstitial pneumonitis.

**Key words:** carbamazepine, drug-induced, lymphocytic interstitial pneumonitis.

# Özet

Yirmi yedi yaşında erkek hasta öksürük, balgam çıkarma, nefes darlığı ve kilo kaybı yakınmaları ile başvurdu. Arka-ön akciğer grafisi bilatera, yaygın retikülonodüler infiltrasyon gösteriyordu. Hasta geçmişte tüberküloz tanısı ile tedavi almıştı ve epilepsi nedeniyle 8 yıldır karbamazepin kullanıyordu. Üç balgam örneği aside dirençli basil açısından negatif idi. Solunum fonksiyon testinde FVC 2.35L (% 51), FEV1 2.12 L (% 54), FEV1/FVC % 90, DLCO % 39 ve DLCO/VA % 75 bulundu. Fiberoptik bronkoskopi normal idi. Transbronşiyal akciğer biyopsisi ve bronşiyal lavaj tanısal değildi. Sağ mini torakotomi ile açık akciğer biyopsisi yapıldı. Patolojik inceleme lenfositik interstisyel pnömoni olarak bildirildi. Nöroloji konsültasyonu sonrası karbamazepin kesildi. İki ay sonra hasta klinik ve fonksiyonel olarak iyi idi. Toraks bilgisayarlı tomografi normal olarak değerlendirildi. Hasta karbamazepine bağlı lenfositik interstisyel pnömoni olarak kabul edildi.

**Anahtar Sözcükler:** karbamazepin, ilaca bağlı, lenfositik interstisyel pnömoni.

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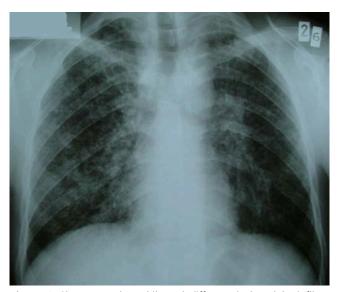
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Drug-induced disease is a fairly common problem in routine medical practice. Interstitial pneumonitis is the most common form of drug-induced pulmonary interstitial disease (1). Carbamazepine is widely used in the treatment of epilepsy, neuropathic pain and bipolar affective disorders (2). A large number of side effects have been reported, including hematological, hepatic, neurological, cardiac, dermatological and renal disorders (2,3). Pulmonary complications have rarely been recognized with carbamazepine (3). We present a case of lymphocytic interstitial pneumonitis induced by carbamazepine.

## **CASE**

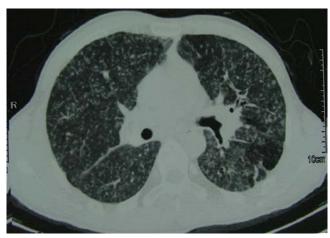
A twenty-seven-year-old male was admitted with cough, sputum production, dyspnea and weight loss. Postero-anterior chest x-ray showed bilateral, diffuse reticulonodular infiltration (Figure 1).



**Figure 1.** Chest x-ray shows bilateral, diffuse reticulonoduler infiltration.

He had received tuberculosis treatment in the past and he had been taking carbamazepine for 8 years for a seizure disorder. The physical examination was normal except for coarse crackles heard in both hemithoraces. Routine laboratory tests were within normal limits. Erythrocyte sedimentation ratio was 30 mm/hour. Three sputum smear and culture examinations for acid-fast bacilli were negative. Computed tomography of the thorax showed diffuse

reticulonodular infiltration and ground glass appearance in both lungs (Figure 2).



**Figure 2.** Computed tomography of the thorax shows diffuse reticulonoduler infiltration and ground glass appearance in both lungs

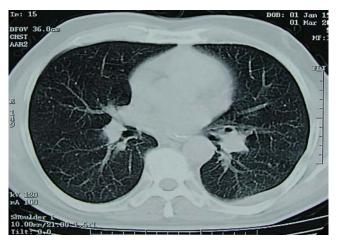
Mediastinal or hilar lymphadenopathy was not observed. Anti-nuclear antibody, anti-DNA antibody, rheumatoid factor and serologic tests for HIV infection were all negative. Fiberoptic bronchoscopy was normal. Transbronchial lung biopsy and bronchial washing were nondiagnostic. Direct examination of bronchial lavage and culture for acid-fast bacilli were negative. Right mini-thoracotomy and open lung biopsy were performed. Pathologic examination revealed diagnosis of lymphocytic interstitial pneumonitis. There was no granuloma in biopsy material. Carbamazepine was withdrawn after neurological consultation. The patient was clinically and functionally well two months later. Computed tomography of the thorax was completely normal (Figure 3).

The patient was considered to be compatible with carbamazepine-induced lymphocytic interstitial pneumonitis.

### **DISCUSSION**

Lymphocytic interstitial pneumonia (LIP) is a benign lymphoproliferative disorder characterized by a diffuse and exquisitely interstitial proliferation of small lymphocytes and plasma cells (4). LIP occurs most commonly in patients who have Sjögren syndrome, autoimmune thyroid disease, acquired immuno-deficiency syndrome, or Castleman disease (5,6).

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**Figure 3.** Computed tomography of the thorax was completely normal two months later.

Several drugs such as carbamazepine and diphenylhydantoin can induce interstitial pneumonia (1-3,7-9). Carbamazepine is a commonly used anticonvulsant drug. While carbamazepine-induced pulmonary sideeffects are very uncommon, interstitial pneumonitis, bronchiolitis obliterans organizing pneumonia, bronchospasm, pulmonary oedema and pulmonary nodules have all been reported (2,3). The mechanism of lung injury by carbamazepine was recognized as an immune-mediated hypersensitivity (3). knowledge, there have been few cases of carbamazepine-induced interstitial pneumonia in medical literature in English. We present a case of lymphocytic interstitial pneumonitis induced by carbamazepine. Our patient presented all clinical, radiological and functional features of interstitial disease. The diagnosis of LIP was established by open lung biopsy. The withdrawal of carbamazepine was followed by rapid clinical, radiological and functional improvement.

In conclusion, carbamazepine can produce lymphocytic interstitial pneumonitis. Our case points to the importance of the past medical history of patients.

### **CONFLICTS OF INTEREST**

None declared.

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