

Unilateral Diaphragm Paralysis Following Vaccination

George S. Habib

Abstract

Unilateral diaphragm paralysis (UDP) is relatively a rare entity. Currently, phrenic nerve injury following cardiac surgery is probably the most common cause. Here we report a case of symptomatic UDP that developed 5 days following vaccination against polio, Diphtheria (both subcutaneously), hepatitis A and abdominal typhus (intramuscularly). He had no fever or other neurological symptoms or signs. There is only one case report in the literature of UDP following immunization or vaccination with oral polio and triple DTP in a 6 month-old female. In contrast to our patient, here there was evidence of active polio disease with isolation of polio strains from her stool.

Keywords: Diaphragmatic paralysis; Unilateral; Vaccination

Introduction

Unilateral diaphragm paralysis (UDP) is relatively a rare entity. Currently, phrenic nerve injury following cardiac surgery is probably the most common cause [1]. Other causes include cervical spondylosis [2], compressive tumors [3], trauma [4], herpes zoster [5], poliomyelitis [6], pneumonia and others. However many cases of UDP remain idiopathic [4].

Patients with UDP but without underlying lung disease are usually asymptomatic at rest, however may have dyspnea on effort or while in supine position. An elevated hemi-diaphragm on chest x-ray is suggestive of the diagnosis that can

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Department of Medicine, Carmel Medical Center, Haifa 34362, Israel. Email: georgeha@clalit.org.il

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be confirmed with fluoroscopic sniff test with paradoxical movement of the paralyzed hemi-diaphragm with inspiration. Pulmonary function test usually shows reduced vital capacity of 70-80% of predicted normal values. The prognosis of UDP in idiopathic cases is usually fair. Most cases generally lead a normal life and some cases may recover spontaneously.

Here we report a case of UDP following vaccination against polio, Diphtheria, hepatitis A virus and typhus prior to a planned trip to India in a 62-year-old male physician.

Case Report

Sixty-two year old man, who works as a physician and has controlled diabetes (HgA1C < 6.6), hypertension and gout, received vaccination against polio (IPV- immovax, Aventis, France) given subcutaneously, Diphtheria (dT-immovax, Aventis, France) given intramuscularly and against hepatitis A virus and abdominal typhus (Hepatyrix, GSK, Belgium) given intramuscularly before a planned visit to India, 5 days following the vaccination, patient started to complain of orthopnea and shortness of breath on effort without cough, fever or chest pain. Eventually, the patient traveled to India for one week and came back with the same complains without progression, 3 weeks after the start of symptoms patient was referred to the hospital and admitted for evaluation. His medications included aturvastatin, metformin, doxazocin, allopurinol, amlodipine, vitamin E, and candisartan. On physical examination, while in sitting position, patient was without stress or cyanosis and had breath count of 12 per minute. Jugular venous pressure was normal. Heart sounds normal. Decreased breath sounds at the right lung base and dullness on percussion. The rest of the physical examination was negative. Routine complete blood count and chemistry were normal. Blood gases showed arterial oxygen saturation of 95% in room air, arterial PCO2 of 41 mmHg. Electrocardiogram and echocardiogram were normal.

Chest x-ray showed elevated right diaphragm with evidence of no movement on fluoroscopy (Fig. 1). Computerized tomography of the neck and chest showed no evidence of tumor but right diaphragm paralysis. Electromyogram



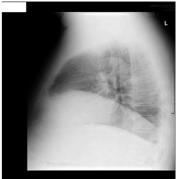


Figure 1. Antero-posterior and lateral views of chest x-ray of the patient.

(EMG) of the upper extremities was negative for neuropathy or myopathy.

Pulmonary function test showed an abnormal restrictive pattern with FEV1 of 1.52 liters (43% of predicted value), and FVC 2.12 liters (46.7% of predicted value). FEV1: FVC ratio of 71.82%, total lung capacity of 4.42 liters (59.2% of predictive value), and residual volume of 2.07 liters (82.3% of predictive value).

Patient was evaluated by neurologist who found no evidence neuropathy or myopathy of any source.

Follow-up after 3 months without subjective or chest xray changes and repeated pulmonary function tests showed no change in volume or flow parameters.

Discussion

The cause of the UDP in this patient is not clear. The temporal relationship with the vaccination makes the association very suspicious.

There is only a single case report of UDP following immunization in the literature [7]. A 6-month-old female who developed fever and flaccid paralysis of all limbs and right diaphragm following immunization with oral polio and triple (DTP) vaccines. Poliovirus types isolated from her stool matched with the vaccine strains. This case illustrates the development of full blown polio. The polio virus immunization in our patient is a dead virus and there was no evidence of the development of acute infection of polio or other infection in our patient.

There are no reports of diaphragmatic paralysis following Typhus or hepatitis A virus infection. There is one report of diaphragmatic paralysis in a 3-week-old infant following acute infection with diphtheria [8].

Our patient had diabetes, a disease that could be complicated with neuropathy or myopathy. However, he had no symptoms, sings or EMG findings suggestive of neuropathy. In the literature, there is one report of reversible unilateral right diaphragmatic hemi-paralysis in a 52-year-old female

with diabetic neuropathy [9].

In my opinion, the most likely explanation in our case is a neurological adverse effect affecting the phrenic nerve following vaccination. Neurological adverse effect post vaccination were reported, especially encephalomyelitis following different types of immunization including diphtheriatetanus-polio and others [10]. A coincidental event cannot be ruled out.

Conflict of Interests

None.

Funding

None.

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