Persistent Hiccups as a Rare Presenting Symptom of Pulmonary Embolism

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Pulmonary embolism (PE) is a life-threatening condition that may present as dyspnea, chest pain, cough or hemoptysis, but often occurs without symptoms. It is not typically associated with hiccups. Hiccups are generally self-limiting benign contractions of the diaphragm that may be associated with medications or food but may also be symptomatic of serious disease when persistent. We report 3 cases of PE presenting as persistent hiccups. [West J Emerg Med. 2012;13(6):479-483]

INTRODUCTION

Pulmonary embolism (PE) is a potentially lethal condition that can be difficult to diagnose. The incidence in the United States is as high as 1 per 1,000 persons per year, and it is the third commonest cause of death in hospitalized patients.1 Early diagnosis and treatment are crucial in ensuring a good outcome, but symptoms are non-specific and the index of suspicion often low.2-4 As a result the diagnosis is often made post-mortem.5

Hiccups are generally self-limiting benign contractions of the diaphragm that may be associated with medications or food but may also be symptomatic of serious disease when persistent. We report 3 cases of PE presenting as persistent hiccups.

CASE DESCRIPTION

A search of the medical literature (search for PE and intractable hiccups through Medline, the Cochrane library database and Google search engine) did not reveal any case series of hiccups as a presenting symptom of PE, although a single citation describing this entity was identified in the lay literature and one case report.33-34 In 2 of our 3 patients, recent surgery was a risk factor for PE.

Patient # 1

A 52-year-old African-American male without significant prior medical or surgical history presented to the emergency department (ED) with a 3-day history of hiccups. He had had progressively worsening dyspnea for 4 weeks, a dry cough and pain in the upper right back associated with inspiration and coughing. He had increasing leg edema for 2 weeks and was told he had congestive heart failure at another hospital.
institution, where he was advised admission, but left against medical advice. The rest of his review of systems (ROS) was unremarkable and he had no risk factors for deep venous thrombosis (DVT), such as family history of clotting disorder or personal history of immobilization, long journeys, surgery or trauma. He had quit occasional smoking many years ago and did not use alcohol or narcotic drugs.

Physical examination revealed a well-built male in mild respiratory distress with a respiratory rate (RR) of 20 breaths/minute, pulse rate (PR) of 108 beats/minute and blood pressure (BP) 113/76 mmHg. His oxygen saturation was 89–91% on room air. His electrocardiogram (EKG) was normal except for sinus tachycardia, and a duplex ultrasound of his lower extremities was negative for DVT. His portable chest radiograph (CXR) was the first hint of abnormality (Figure 1), showing a prominent pulmonary central artery (early Fleishner’s sign, red arrow heads) and a pruned tree /cut off of the pulmonary arteries (Westermark’s sign, black arrows). With the working diagnosis of PE, a computed tomography (CT) of the chest with intravenous (IV) contrast was performed which revealed a massive saddle embolus of the pulmonary artery (Figure 2A, black arrow). Patient was heparinized and transferred to a tertiary center where pulmonary thrombectomy was performed. He made a full recovery with resolution of hiccups and dyspnea. His laboratory examination with complete blood count (CBC), complete metabolic panel (CMP), prothrombin time (PT)/international normalize ration (INR) and partial thromboplastin time (PTT) was essentially normal. Workup for a coagulation disorder failed to reveal a reason for his embolus.

**Figure 1.** Chest radiograph demonstrating a prominent central pulmonary artery (early Fleishner’s Sign, red arrows) and a cut-off of the pulmonary arteries bilaterally (Westermark sign, black arrows).

**Figure 2.** A. Computed tomography (CT) showing a saddle embolus (black arrow). B. CT showing a large left pulmonary artery embolus (filling defect, red arrow). C. CT showing a left pulmonary artery embolus (filling defect, red arrows).
Patient #2

A 40-year-old Hispanic female came for her follow-up gynecology appointment after undergoing total abdominal hysterectomy for fibroids under general anesthesia 12 days earlier. She complained of severe left lower back pain of 1-day duration and was referred to the ED to rule out renal colic. Her postoperative course was uneventful and she had been ambulating on post-operative day #1 and discharged home shortly after surgery. In the ED she spoke in fragmented sentences due to persistent hiccups. ROS and past medical history were unremarkable.

Physical examination revealed a young female with hiccups and mild lower abdominal tenderness, but no signs of rhonchi, rales or crepitations in the chest. She was given dilaudid and morphine for back pain, but her PR was 110-120 beats/minute and BP 112/78 mmHg. Temperature was 100.7°F Fahrenheit and RR 20 breaths/minute. Based on the previous case of PE presenting with hiccups, a diagnosis of pulmonary embolism was entertained. CXR was unremarkable, but EKG revealed S1T3Q3 changes (Figure 3). CT chest revealed a large left pulmonary embolus (Figure 2B, red arrow). Bedside duplex ultrasound did not show DVT. Her laboratory examination with CBC, CMP, PT, INR, and PTT was essentially normal. Workup for a coagulation disorder failed to reveal a reason for his embolus. She was started on IV heparin infusion and warfarin and discharged home with resolution of hiccups and back pain.

Patient #3

A 46-year old African-American male presented to the ED with persistent hiccups 2 days after having left inguinal herniorrhaphy under general anesthesia. Surgery was uncomplicated and performed as an ambulatory procedure. He had been ambulating at home, but one day after surgery developed hiccups. His past medical history was unremarkable, but he had undergone right inguinal herniorrhaphy in 2008 without any adverse events or symptoms. He was a nonsmoker and drank no alcohol and denied any other risk factors for DVT.

Physical examination revealed a middle-aged man in no apparent distress except for hiccups. At presentation his temperature was 98°F Fahrenheit, PR 81 beats/minute, BP 128/77 mmHg and RR 18 breaths/minute. ECG showed normal sinus rhythm. He was given chlorpromazine, which resolved his hiccups, and he was discharged home. On his way home the hiccups returned and he came back to the ED. A second dose of chlorpromazine failed to resolve the hiccups this time. He also complained of feeling dizzy and his BP was found to be low. A chest CT was ordered and revealed a pulmonary embolus (Figure 2C, red arrow heads). His laboratory examination with CBC, CMP, PT, INR, and PTT was essentially normal. Workup for a coagulation disorder failed to reveal a reason for his embolus.
DISCUSSION

The symptoms and signs of PE are highly variable and non-specific and a high index of suspicion is critical in the diagnosis of this potentially lethal condition. The most common symptoms of PE are dyspnea at rest or on exertion (73%), pleuritic chest pain (44%), and cough (34%) with calf pain (44%) and calf or thigh swelling (41%) indicating DVT preceding PE. Patients with such symptoms or signs often need additional testing to confirm or exclude the diagnosis. Hiccups on the other hand are rarely associated with PE. Patients may also be completely asymptomatic, making the diagnosis more challenging. A meta-analysis of 28 studies found that among 5,233 patients who had DVT, 1,665 (32%) had asymptomatic PE, highlighting the importance of a high index of suspicion. Risk stratification or pretest probability assessment thus becomes paramount in raising awareness of a possible diagnosis of PE.

All patients with chest pain or shortness of breath should have CXR and ECG. In patients with high pretest probability and non-specific symptoms, even if CXR and EKG are normal d-dimer level, lower extremity duplex scan, ventilation-perfusion (VQ) scans and CT angiogram may be needed.

Numerous prediction models and algorithms have been developed for assessment of patients with possible PE, including the Geneva, Kline, Pulmonary Embolism Rule-out Criteria (PERC), Pisa and Wells systems. Of these the Wells criteria and the PERC rules are the most popular. The different elements included in these scoring systems are clinical history, physical examination, and diagnostic tests, such as arterial blood gas, D-dimer, ECG, and CXR with points ascribed to each element. Based on the total score and the clinical picture, the need for additional testing, such as VQ scans and CT angiograms, is determined by the clinician. Thus, by PERC rules if a patient meets all 8 criteria and falls into a low-risk category the probability of PE is < 2% and further testing may not be required. By Wells’ criteria, a patient with low probability of PE and a negative D-dimer test may be discharged home without more testing.

Zylicz reported a case of intractable hiccups due to PE suggesting that a thrombus in the inferior vena cava (shown on ultrasound) caused PE and then hiccups, but no objective evidence of PE is presented. Hiccups resolved with low molecular weight heparin, but recurred when heparin was discontinued. The patient also had underlying non-small cell lung cancer and it is unclear if lung cancer was the source of the hiccups. All of our patients had CT-documented PE and 2 had the risk factor of recent surgery.

Patient #1 had a low Wells score of 1.5 (HR > 100), making him low risk, but he failed 3 of 8 PERC criteria and the CXR and overall gestalt raised suspicion for PE. Also the Kline decision rule (age/hypoxemia) put him in the unsafe category with a pretest probability of 45.2%. The revised Geneva score placed him in a moderate risk group based on tachycardia alone. Patient #2 had a score of 3 on Wells criteria (HR > 100; recent surgery) and failed 2 of 8 PERC categories. In patient #3 Wells score was 1.5 (recent surgery), and he failed only 1 of 8 PERC categories.

Different scoring systems put patients in different risk categories highlighting the difficulty in relying only on one scoring system. Intractable hiccups in 2 patients raised our level of suspicion after our experience with patient #1, although both of these patients had recent surgery. CXR in patient #1 and EKG in patient #2 also helped point us in the right direction. Our series of 3 patients with PE who had hiccups as a presenting symptom highlights the importance of including hiccups in the constellation of symptoms and signs associated with PE. If pretest probability testing is also taken into account, it will help us decide which patients need more advanced imaging and testing to confirm or exclude the diagnosis.

CONCLUSION

Hiccups are common and often idiopathic, but persistent hiccups should be taken seriously. They may be manifestations of immediately life-threatening conditions like myocardial infarction or even pulmonary embolism. Including hiccups in the pantheon of symptoms associated with PE will raise awareness as demonstrated by our small series.

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