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Epinephrine soaked tampons induced transient acute dilated cardiomyopathy during FESS procedure



Sari Naddaf¹, Scott Ehrenberg¹, Rony Hakim², Muhamad Mahamid³, Yoav Turgeman^{1,3} and Ofir Koren^{1,3}*

Abstract

Background: Epinephrine, in all modes of use, may pose a wide range of cardiotoxic events, ranging from sinus tachycardia to heart failure, life threatening arrhythmias, and even death. Because of daily and extensive use of epinephrine, these unusual and rare events tend to be forgotten by physicians. We present a case of dilated cardiomyopathy that developed following routine use of epinephrine-impregnated tampons during function endoscopic sinus (FESS) surgery.

Case presentation: A healthy, 24-year-old man with no family history of heart disease has undergone elective surgery under general anesthesia to repair the paranasal sinuses using endoscopic approach. During surgery, soon after being treated with 1: 1000 diluted epinephrine-soaked tampons, an hypertensive crisis was noticed followed by pulseless electrical activity. An extensive examination led to the diagnosis of non-ischemic dilated cardiomyopathy. After several days of heart failure medical therapy, complete resolution of all structural and functional changes was achieved.

Conclusion: In our case, we present an unusual and rare event of acute dilated cardiomyopathy following the use of epinephrine-soaked tampons during elective FESS surgery. A prompt response was observed after several days of heart failure treatment. Awareness of the epinephrine cardiotoxic potential even in the form of soaked tampons is essential for proper diagnosis and prompt treatment.

Keywords: Epinephrine, Dilated cardiomyopathy, Heart failure, FESS, Surgery

Background

Dilated cardiomyopathy (DCM) is recognized by the dilation of the right, left, or both ventricles in the absence of abnormal loading conditions as hypertension and valve disease, or significant coronary artery disease [1–5].

The etiology of DCM is extremely heterogeneous sometimes classified based on known genetic mutation. Among the non-genetics cause are different etiologies,

tain toxins as alcohol or allergens; complication of pregnancy, systemic endocrine or autoimmune diseases, and infection as HIV [6].

including myocarditis, exposure to drugs as cocaine, cer-

One of the rarest etiologies regarding toxin and metabolic-related cause of DCM is catecholamines, and its various derivates, which has been reported mainly in case reports or short series review [7-13].

Pheochromocytoma, as a rare neuroendocrine catecholamine-producing tumor, has also been described as an etiology of reversible DCM and was found in \sim 39% of pheochromocytoma-related cardiomyopathies [14]. Paul et al. proposed that catecholamine-induced vasoconstriction, a direct toxic effect of the by-products

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of catecholamine degradation and direct receptormediated mechanisms, contribute to cardiomyopathy in subjects with pheochromocytoma [15].

Case description

A 24-year old male with chronic rhinosinusitis was admitted for an elective Functional endoscopic sinus surgery (FESS) procedure under general anesthesia. During the operation, the surgeon applied several tampons soaked in 1:1000 dilution epinephrine to the nasal mucosa. Ninety minutes into the surgery, unexpectedly, blood pressure rose to 210/130 mmHg followed by pulseless electrical activity. CPR was initiated, with the administration of 2 mg of IV epinephrine in consecutive doses, leading to the return of spontaneous circulation. ECG showed sinus tachycardia and a prolonged QTc interval of 486 ms, Inverted T waves in leads I and aVL, and no signs of acute ischemic changes (Fig. 1). A chest x-ray demonstrated pulmonary edema and a borderline enlarged cardiac silhouette (Fig. 2).

The patient was placed on mechanical ventilation. Transthoracic echocardiogram showed a dilated left ventricle with an increased end-diastolic dimension (124% of normal value) with a mild reduction in LV mass, a severe reduction in systolic function, apical akinesis, and hyperdynamic base. The estimated systolic left ventricular ejection fraction was 30% (Fig. 3). Cardiac troponin and CPK were elevated. NT or NT pro-BNP were not taken. A head CT was performed and demonstrated mild global cerebral edema, with multiple maxillary sinus fractures (Fig. 4).

Medical information gathered from family members indicated no family history of heart diseases. The patient did not smoke, use illegal drugs nor consumed alcohol on daily basis. The patient was diagnosed with dilated cardiomyopathy and admitted to the intensive cardiac care unit. Upon arrival, upload

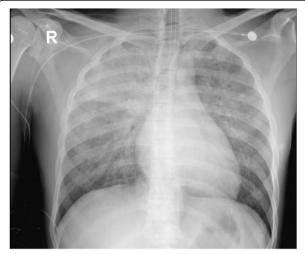


Fig. 2 Chest X-ray at admission indicate bilateral pulmonary edema and normal heart silhouette

titration of ACE inhibitors, β -blockers, and diuretics initiated. The following ECG strips indicate normal sinus rhythm, QTc interval of 420 ms, and normal T wave in lateral leads. We decided not to perform a diagnostic coronary catheterization since the patient's risk profile for ischemic cardiomyopathy was low. Objectively, there were no ischemic changes on ECGs strips and no regional wall motional abnormalities were seen on TTE upon arrival.

Three days later, a second echocardiogram was performed which showed a normal-sized left ventricle, with preserved systolic function (Fig. 5).

The patient was discharged in full functional capacity. An ambulatory cardiac MRI performed two weeks after discharge revealed mildly dilated LV cavity, good global systolic function, and no signs of late gadolinium enhancement or edema.

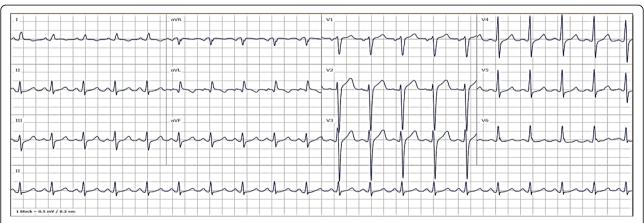


Fig. 1 ECG at admission indicate sinus tachycardia, prolonged QTc (QTc = 486 ms), Inverted T waves in leads I and aVL, and no signs of acute Ischemic changes



Fig. 3 Transthoracic Echocardiography at admission demonstrates a dilated left ventricle dimension. Apical 4 Chambers view at the end of the diastole

Discussion

The relationship between exogenous catecholamines and dilated cardiomyopathy is not fully understood. Some propose that elevated catecholamine levels decrease myocardial viability via cyclic -AMP mediated Ca + overload [16, 17].

While reductions in ejection fraction and left ventricular dysfunction was found to be reversible, mild histologic changes were found in dog studies indicate chronic perivascular fibrosis which seems to be irreversible [18].

Since activation of alpha or adrenergic receptors has been shown to induce stress-induced cardiomyopathy, it has been shown in studies that these cardiac effects can be attenuated by pretreatment with the use of alpha and beta blockers [19–23]. In addition, increasing levels of

estrogen have shown partial attenuation of these cardiac changes [24, 25].

Dilated cardiomyopathy is usually characterized by long-standing processes or conditions, its appearance in the acute setting is not common and a detailed evaluation of acute onset dilated cardiomyopathy (ADCM) typically does not elucidate a specific etiology in most cases [26].

Conclusion

We report a rare case of a transient acute non-ischemic transient dilated cardiomyopathy following exposure to sstandard diluted dose epinephrine soaked-tampons during ENT procedure. Prompt heart failure treatment resulted in complete resolution. Our report provides

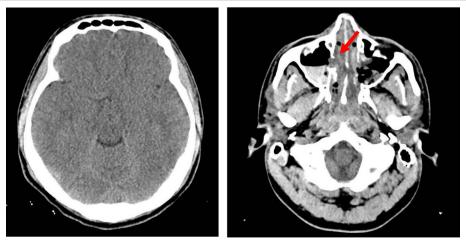


Fig. 4 Head CT at admission shown Global Edema (Left-sided) and multiple Maxillary Sinuses fractures (Right-sided red arrow)

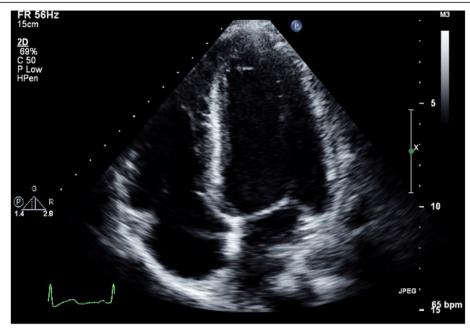


Fig. 5 Transthoracic Echocardiography at discharge demonstrates a normal-sized left ventricle dimension with normal LV mass. Apical 4 Chambers view at the end of the diastole

supporting evidence of the cardiotoxic devastating potential effect of epinephrine and its role in acute dilated cardiomyopathy. We hope that this paper will raise awareness among physicians and surgeons to the relevance of cardiotoxic effect of epinephrine, especially in the form of soaked tampons.

Abbreviations

FESS: Functional endoscopic sinus surgery; ENT: Ear, nose and throat; DCM: Dilated cardiomyopathy; ECG: Electrocardiography; TTE: Transthoracic echocardiography; CT: Computerized tomography; MRI: Magnetic resonance imaging; PEA: Pulseless electrical activity; CPR: Cardiopulmonary resuscitation; ACEI: Angiotensin-converting enzyme inhibitor; BB: Beta-blocker

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Authors' contributions

SN, SE, RH, MM, and OK contributed to the writing, editing, formatting of the main manuscript, and production of the Figs. MM, OK and YT provided care to the patient and revised the manuscript. All authors have read and approved the manuscript.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Ethics approval and consent to participate

Informed consent was waived due to the use of anonymous patient's data as well as due to the retrospective nature of the case study.

Consent for publication

Written consent to publish the information presented in the clinical case was obtained from the patient.

Competing interests

The authors declare that they have no competing interests.

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