Prolonged oral sildenafil use-induced Mondor disease: a case report

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Introduction

Mondor disease (MD) is a palpable, painful, subcutaneous induration caused by superficial venous thrombosis [1,2]. In 1939, the first report involved a female patient with symptoms in the thoracoepigastric region. Penile MD was first reported in 1958 [2,3]. MD is rare and its etiology remains to be established. However, a relationship with Virchow’s triad is strongly suspected [1,3]. Moreover, rare cases of MD related to oral contraceptives have been reported [4]. Sildenafil, also known as Viagra, is currently used for the management of male erectile dysfunction and has been approved for the treatment of pulmonary hypertension [5]. We report a case of penile MD that was suspected to be related to prolonged oral sildenafil use.

Case

A 46-year-old man visited the emergency department (ED) with sustained penile pain and swelling that began 7 hours after sexual intercourse. He had a history of hypertension, gastroesophageal reflux disease, and posttraumatic stress disorder. He had used oral sildenafil intermittently for 11 years for erectile dysfunction, and
the night prior to arrival at the ED, he had engaged in sexual intercourse after taking sildenafil. No clinical signs of intercourse-related trauma to the genital area were observed. The following morning, he noted a sustained, painful penile erection with swelling. Using a visual analogue scale, the patient’s pain intensity was scored at approximately 2 points. His initial vital signs were as follows: blood pressure, 100/60 mmHg; heart rate, 110 beats/min; respiratory rate, 20 breaths/min; and body temperature, normal. He reported no urinary symptoms and declined biochemical laboratory tests. On physical examination, the penis was swollen, with no increase in skin temperature. He complained of tenderness over the dorsal penile vein area and a mild protrusion was noted (Fig. 1). The differential diagnosis included suspected unidentified priapism, soft tissue infection, or penile fracture, and an ED ultrasound examination was performed. No abnormal disruption of the tunica albuginea was noted in the penile shaft. However, a dilated vein with hyperechoic thrombosis was observed in the right superficial dorsal vein of the penis with soft tissue swelling. Moreover, the absence of flow signal in the thrombotic lesion was identified on the color Doppler ultrasound image (Fig. 2). Following ED-administered pain management, he was advised to avoid sexual intercourse and was discharged from the ED with a prescription for nonsteroidal anti-inflammatory medication. A follow-up appointment at the Department of Urology in outpatient clinic was made to determine whether surgical thrombectomy would be required. During the next 2 weeks of outpatient follow-up, the patient’s symptoms improved.

**Discussion**

MD is a rare, painful, superficial venous thrombosis that has been reported to predominantly occur in the chest wall or penis [2]. Penile MD comprises venous thrombosis of the dorsal superficial penile vein and is typically benign and self-limiting. However, its pathophysiology remains unclear [1]. Virchow’s triad, involving vascular wall injury, venous stasis, and hypercoagulation, has been reported to cause venous thrombosis [1]. The most likely cause of penile MD is vascular wall injury, which is one factor in Virchow’s triad, due to lengthening or splitting of the vessel from prolonged vigorous sexual activity, typically within 24 to 48 hours [3]. Hypercoagulation due to protein S, protein C, and antithrombin III deficiency has also been reported to cause penile thrombosis [6]. Di-
Sildenafil-induced transient arrhythmic cardiac embolism

Supposed mechanism

Bilateral blurred vision

Right side weakness

Superior sagittal sinus

Left middle cerebral artery

Right transverse and sigmoid sinus

Site of thrombus

Platelet aggregation activation

Platelet aggregation activation

Jugular vein

Sildenafil-induced transient arrhythmic cardiac embolism

Table 1. Previous reports of sildenafil-associat ed vascular thrombosis

<table>
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<tr>
<th>Study</th>
<th>Symptom</th>
<th>Site of thrombus</th>
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<tr>
<td>Rupa et al.</td>
<td>Bilateral blurred vision</td>
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<td>Kim et al.</td>
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<td></td>
<td>Headache</td>
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Sildenafil acts by blocking phosphodiesterase-5, an enzyme that promotes the breakdown of cyclic guanosine monophosphate (cGMP), causing relaxation of the corpora cavernosa and penile arteriolar smooth muscle. Owing to the localized vasodilatory effect of the nitric oxide–guanosine monophosphate pathway, the U.S. Food and Drug Administration has approved its use in the treatment of pulmonary hypertension [5]. However, sildenafil has been reported to have various adverse effects, including venous thrombosis. Table 1 shows previous reports on the association between venous thrombosis and oral sildenafil [5,8,9].

According to previous reports, the mechanisms of vascular thrombosis may be due to vascular stasis and/or platelet aggregation resulting from long-term sildenafil use. cGMP can inhibit platelet aggregation. Initially, cGMP causes platelets to clump to seal a wound and later reverses this action to stop an excessive build-up of cells that might then block a blood vessel. Sildenafil-enhanced intracellular cGMP activates cGMP-dependent protein kinase (PKG) [10]. PKG promotes von Willebrand factor-induced activation of human platelets [11]. Rupa et al. [5] suggested that the chronic use of sildenafil causes high cGMP levels, which might interfere with normal endothelial function. Moreover, sildenafil lowers systolic blood pressure by 8 to 10 mmHg owing to its vascular relaxation effect, which could be the cause of vascular insufficiency [12]. Decreased blood flow owing to blood vessel relaxation may be an additional cause of thrombus formation.

Penile MD is not life-threatening, but it can negatively affect a patient’s quality of life. Considering the adverse effects of sildenafil, prolonged oral administration could cause abnormal venous thrombosis of the superficial dorsal penile vein. It is also reason-


