Inadvertent Injection of Sodium Tetradecyl Sulfate During Placement of Mental Nerve Block: Discussion of Management and Outcome

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Sclerotherapy involves the injection of a foreign substance into the lumen of a vessel to destroy the wall of that vessel and lead to its occlusion. Sodium tetradecyl sulfate (STS) is a detergent solution commonly used in sclerotherapy to treat varicose and telangiectatic veins. Its actions lead to destruction of endothelial cells.1,2 We report a case of the inadvertent injection of STS into the mental nerve of a patient who was supposed to be treated with an anesthetic agent for a mental nerve block before hyaluronic acid treatment. We discuss the case presentation, management, and outcome.

Case Report

A 39-year-old woman presented to a clinic elsewhere for hyaluronic acid treatment. Local anesthesia consisting of 1% lidocaine with epinephrine and bicarbonate was used to anesthetize the right mental nerve. Inadvertently, 2 mL of sodium tetradecyl sulfate 0.2% was injected into the left mental nerve. The patient complained of mild tenderness to the left mental nerve. The site was then infiltrated with 7 mL of normal saline and 10 U of hyaluronidase. On Day 2, the patient complained of moderate pain but denied any paresthesia. On physical examination, she had a 2- × 2-cm mild blue discoloration at the site of injection. Pentoxifylline 400 mg three times a day was started, and cephalexin 250 mg four times a day for 7 days was prescribed. On Day 3, the patient called complaining of severe pain and increased discoloration. Her physical examination showed yellow–blue discoloration. (Figure 1). On Day 4, hyperbaric oxygen treatment was started. She tolerated two treatments (2 hours each at 2 atm for 2 consecutive days) well without complications. On Day 7, she returned to clinic. Her pain was completely resolved, but a yellow–blue discoloration persisted (Figure 2). On Day 14, she returned to clinic with her discoloration completely resolved (Figure 3).

Discussion

Inadvertent injection or extravasation of STS can result in cutaneous necrosis and nerve damage. In this patient, the error was fortunately recognized early, and she was promptly treated with copious amounts of normal saline and hyaluronidase. Pentoxifylline was started on Day 2 to help prevent necrosis. Hyperbaric oxygen therapy was also started once she began to complain of pain and worsening discoloration.

Cutaneous necrosis is a known and relatively uncommon yet feared side effect that may occur after the injection of a sclerosing agent.3 Inadvertent subcutaneous injection of up to 1 mL of hypertonic saline 23.4% in lieu of lidocaine into the neck or cheek has been reported to result in no adverse

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sequelae. In this case, cutaneous necrosis was likely avoided by rapid physiologic dilution of the hypertonic saline. Certain sclerosants are associated with a lower risk of cutaneous necrosis. Duffy has reported injecting 0.4 mL of 3% pol into his own forearm without ulceration (Figure 4). Injection of STS into a nerve has also been reported to be painful and can result in anesthesia and sometimes permanent nerve impairment. Multiple cases of inadvertent injection into a nerve have been reported with resulting variable degrees of nerve paralysis or paresis.

Tissue necrosis can occur with any sclerosant, but it is more commonly associated with high concentrations of sclerosants and in particular with certain types of sclerosants. It is common with hypertonic saline and high-strength sodium tetradecyl sulfate. Sodium tetradecyl sulfate 1% will cause contact ulceration, whereas sodium tetradecyl sulfate 0.2% is dilute and will not cause contact ulceration. Tissue
necrosis also occurs more commonly after sclerotherapy for vessels involving the medial malleoli, telangiectasia below the ankle (Figure 5), and on the face (Figure 6).8–10 Nevertheless, prompt treatment is critical to avoid or lessen the potential for necrosis and nerve damage.

Management of extravasated sclerosing solution includes prompt dilution of the site. Dilution with hyaluronidase in normal saline has been shown to limit the extent and prevent cutaneous necrosis from 3% STS.11 Hyaluronidase helps to prevent tissue necrosis through accelerated dilution, cellular stabilization, and improved wound repair,12–15 although at least one study has shown that hyaluronidase solution must be injected within 60 minutes of extravasation to be effective.16 Although hyaluronidase is supposedly ineffective when used more than 60 minutes after extravasation, it has been shown to be of value in avoiding tissue necrosis after the use of hyaluronic acid17 even when the treatment had been performed several days before. Hyaluronidase and a protocol for its employment should be immediately accessible to anyone who uses dermal filling agents or performs sclerotherapy. Hyaluronidase (Vitrase is available at Ista Pharmaceuticals, Inc., Irvine, CA). It must be kept refrigerated and should not be used in patients with allergies to bee stings; skin test should be performed before use.

Pentoxifylline also shows promise in the treatment of extravasated sclerosing solution to prevent cutaneous necrosis. Pentoxifylline decreases tissue injury of ischemia-reperfusion by increasing the deformability of red blood cells and decreasing blood viscosity.18 Hyperbaric oxygen treatment has become a mainstay for the treatment of chronic and nonhealing wounds. It has been shown to promote wound healing by increasing tissue oxygen and stimulating angiogenesis, fibroblast proliferation, and collagen synthesis.19,20 Hyperbaric oxygen therapy was used successfully in this patient to help reverse her deteriorating condition and avoid ulceration or permanent nerve damage.
In summary, inadvertent injection of a sclerosing agent has the potential for a devastating outcome. Safeguards must be in place to avoid this potentially disastrous problem. A simple way to clearly identify sclerosant-filled syringes is exemplified in Figure 7, in which colored gummed stickers attached to the piston of the syringes identify the presence and concentration of sclerosants in the syringes.

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References


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