Possible local anesthetic resistance in Emery–Dreifuss muscular dystrophy during regional anesthesia

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We recently provided perioperative anesthetic care for a 31-year-old man with Emery–Dreifuss muscular dystrophy (EDMD) presenting for bilateral Achilles lengthening and equinovarus deformity correction. We performed bilateral popliteal sciatic and femoral nerve blockades, using ultrasound guidance and nerve stimulator confirmation, mainly to avoid some potential pitfalls of general anesthesia and airway instrumentation. However, the patient appeared resistant to the effects of local anesthetic. The few published reports of anesthesia management for such patients describe general and neuraxial techniques [1,2]. To our knowledge, this is the first written description of regional anesthesia in this rare patient population.

EDMD is a rare X-linked condition that primarily affects skeletal muscle, leading to joint contractures with progressive weakness, and cardiac muscle, leading to malignant arrhythmias and cardiomyopathy [3]. EDMD presents unique challenges to the anesthetist, including arrhythmias (i.e., atrioventricular heart block), cardiomyopathy, possible compromised airway mechanics leading to difficult intubation (i.e., cervical paraspinal contractures and hypoplasia of the third to fifth cervical vertebral bodies that can influence neck flexion and extension), potential for difficult neuraxial access due to paraspinal muscle contractures, and concern for adverse medication reactions (i.e., succinylcholine exaggerated hyperkalemic response or risk of rhabdomyolysis [3] and negative volatile anesthetic reactions [1]). Considering the challenges of anesthetic management in this patient, it seemed reasonable that bilateral sciatic and femoral nerve blockades could avoid airway instrumentation and general anesthesia. Notably, however, the patient commented preoperatively that “…every time I received a local anesthetic injection at the dentist’s office, I required multiple injections in order to feel any effect.”

A popliteal approach nerve block of the sciatic nerves was performed bilaterally with ultrasound, visualizing the circumferential perineural spread of ropivacaine at the junction of the tibial and common peroneal nerves. Bilateral perineural catheters were then placed. Then, a single-shot block of the femoral nerves at the inguinal crease was performed bilaterally with ultrasound, visualizing the circumferential perineural spread of ropivacaine. Despite nerve stimulation confirmation (Stimuplex® HNS12, B. Braun Medical, Bethlehem, PA, USA) with less than 0.5 milliamperes and ultrasound imaging of perineural local anesthetic spread at all block sites (Fig. 1), the patient reported intact sensation to temperature (alcohol pad testing) and nociception (blunt needle prick) within all affected nerve distributions. Both sciatic catheters were then bolused with lidocaine, yet the patient reported achieving only a minimal degree of temperature sensation change. Owing to inadequate surgical anesthesia with the regional technique, we decided to administer total intravenous anesthesia (TIVA) using a propofol infusion and intermittent boluses of...
The patient had atypical requirements for local anesthetics (testing upper limit guidelines), while clinically achieving minimal efficacy with peripheral nerve blocks. Guidelines for the recommended local anesthetic maximum are based on intravenous administration (i.e., ropivacaine 3 mg/kg and lidocaine 4–5 mg/kg) and anesthesia providers often extrapolate these same local anesthetic limits when performing regional techniques. Resistance to local anesthetic effects has rarely been reported, with only one report of decreased efficacy and duration of topical and intradermal local anesthetic administration in patients with Ehler–Danlos syndrome (it was postulated that this was due to the laxity and density of abnormal connective tissues [4], which is clearly not related to muscular dystrophy). There is also research on specific sodium channel variants that may be linked to resistance to local anesthetics [5]. We acknowledge that there are many points to confirm before declaring that our patient clearly demonstrated resistance to local anesthetic. However, he expressed interest in genetic testing of his sodium channels to look for abnormalities related to local anesthetic resistance. If an atypical sodium channel variant were discovered, it could provide valuable insight toward determining if such a variant had a higher incidence in those with muscular dystrophy.

References