

Anesthesiology  
71:322, 1989

## The Lack of Effect of Succinylcholine on Serum Potassium in Patients with Parkinson's Disease

*To the Editor:*—Depolarizing muscle relaxants are known to cause hyperkalemia in a variety of neurological diseases. The effect of succinylcholine (SCh) on serum potassium in patients with Parkinson's disease has not been adequately studied and reported.<sup>1</sup> The etiology of hyperkalemia associated with SCh and Parkinson's disease in a single patient report was clouded by the complex nature of the case.<sup>2</sup>

With approval from the internal review board, seven patients (38–58 yr) with severe Parkinson's disease who underwent elective adrenal medullary to caudate transplantation were studied. They were surgical candidates for the procedure because of poor control of their disease with medical therapy. Preoperatively, their medications for Parkinson's disease were discontinued for approximately 12 h before induction of general anesthesia.

Baseline arterial blood gases and serum K<sup>+</sup> concentration were determined, after which anesthesia was induced with thiopental 5 mg/kg and fentanyl 2–3 µg/kg, and tracheal intubation was facilitated with SCh 1.5 mg/kg. Patients' lungs were ventilated with 100% oxygen and end-tidal carbon dioxide levels maintained within ± 2 mmHg of preinduction values. Arterial blood gases and the serum K<sup>+</sup> concentration were measured 3–5 min following administration of succinylcholine.

In five of the seven patients there was no change in serum K<sup>+</sup> concentration. One patient had an increase of serum K<sup>+</sup> of 0.2 meq/l, and one patient had a decrease of serum K<sup>+</sup> of 0.2 meq/l. In all, patients' arterial oxygen saturation remained above 99%. Arterial car-

bon dioxide was within ± 2 mmHg between the two sampling periods, and arterial pH was within ± 0.02 in all seven patients during this time period. None of the seven patients had any of the EKG abnormalities associated with hyperkalemia during induction of anesthesia or during the first 60 min of the case. All seven patients successfully recovered from general anesthesia without neurologic or cardiac sequelae.

In summary, based upon the finding in this series of seven patients with Parkinson's disease, SCh-induced hyperkalemia was not an identifiable clinical problem.

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(Accepted for publication May 1, 1989.)

Anesthesiology  
71:322–323, 1989

## Source of Specialized Endotracheal Tubes

*To the Editor:*—It was with interest and an appreciation of the circumstances that I read Dr. Holzman's letter regarding fabrication of an elongated endotracheal tube for intubation in a patient with tracheal resection.<sup>1</sup> Because of the size of some species encountered in veterinary anesthesia, veterinary anesthesiologists frequently use specialized endotracheal tubes for maintenance of inhalation anesthesia. One of the vendors of anesthesiology and respiratory care devices for human use entered the veterinary market by applying the materials and expertise utilized for manufacture of human endotracheal tubes to needs of veterinary anesthesia (tubes of up to 30 mm ID and up to 90 cm length). They have manufactured their veterinary product line from the same materials and under the same Current Good Manufacturing Practices guidelines promulgated by the Food and Drug Administration as they have their human product line.\* The only difference is that the human tubes are packaged sterilely before shipment, while the veterinary tubes are not.

When anesthetizing neonatal foals, tracheal tube diameters of 6–10 mm ID are sufficient. However, because of the length of the foal's head, use of commercial human endotracheal tubes is not possible because they are too short, particularly if intubation is performed nasally.<sup>2</sup> One of the vendor's products (fig. 1) is used for these cases and could have potential application in the case described by Holzman. It

is an endotracheal tube made of silicone in sizes ranging from 7–14 mm internal diameter with a 55 cm length (FT-70 to FT-14, Bivona Inc., Gary, Indiana). While a length of 55 cm would have been excessive for that particular case, the tube could be shortened, either to a pre-

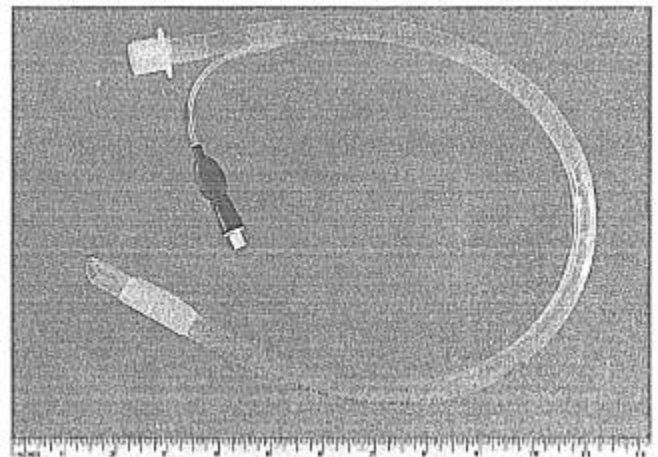


FIG. 1. Commercial 7.0 mm I.D. by 55-cm length endotracheal tube.

\* H. M. Kaufman, Bivona Inc., Gary, Indiana.

determined length by the company; or, following a change by the company in the point where the cuff inflation tube exits the tube wall,† by the anesthesiologist, to a length appropriate for the case. Either method would provide a commercially available endotracheal tube for this type of case and help eliminate the need for improvisation as the case progresses. Should one of these elongated tubes be purchased, it would improve convenience to request that sterilization be performed prior to shipment.

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*(Accepted for publication May 2, 1989.)*