

devices more tempting to investigate and advocate than older, already well-explored instruments. Slavish acceptance of novelty can lead to unfortunate 'fads', later recognized to resemble the emperor who is, if not naked, at least ill-clad.

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REPLY:

The intent of our project was to review the literature since the ASA Guidelines and to determine if previous recommendations should be modified, and to offer guidance regarding newer airway devices.

Cole and Mallon¹ studied eight residents who, after 1000 tracheal intubations (quarter with the flexible fiberoptic endoscope [FFE]), were assessed during tracheal intubation in a further elective 131 patients, 59 with the direct laryngoscope, 72 with the FFE. In the case of FFE-facilitated tracheal intubations, a catheter was placed in the oropharynx before the procedure began for

continuous oxygen insufflation. It took nearly twice as long to complete fiberoptic intubation; only 73% were performed within one minute of apnea. Would this be reproducible in the unanticipated failed intubation, without dedicated assistance?

We acknowledged the value of the FFE in the management of the unanticipated difficult intubation. We suggested that its use was more difficult in the setting of a paralyzed, apneic patient, particularly when the airway might be soiled by blood or secretions. Finucane also took exception to our reservations: "I tend to differ with them on this issue. Fiberoptic assisted intubation can be extremely useful even in anesthetized patients, but ... it requires practice, experience and an assistant to maintain the airway,"² (my italics) We cited evidence that tracheal intubation can be achieved rapidly and safely, without the need for an assistant, with the alternate devices reviewed.

With regard to newer devices, we did not mention the intubating LMA because, at the time, there was no evidence demonstrating its effectiveness. However, the other devices reviewed, with the exception of the McCoy laryngoscope blade, have been available for more than a decade.

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The Charlottetown Click

To the Editor:

We read with concern the letter by Dubois and colleagues describing insertion of the LMA using a semi-inflated rotational technique, the so called "Charlottetown Twist" (*Can J Anaesth* 1998;45:823). The authors make the unsupported statement that this technique is less traumatic than the standard recommended technique, but we consider this to be incorrect. There is no doubt that placement of the LMA with the mask aperture bars facing in the cephalad direction can occasionally be advantageous in moving the cuff from the mouth into the pharynx. The disadvantage is that the cuff must then be rotated back through 180° for the device to function. The authors state that this rota-