
In Reply.—It is interesting to know that Dr. Michael Rosen’s experience in terms of satisfaction with patient-controlled intravenous analgesia (PCA) using meperidine during labor in a population of mixed social and economic status was similar to our experience. In our obstetric unit, although we allow nurses to initiate and maintain PCA, an anesthesiologist is always available on an immediate basis.

Dr. Deborah Wilson’s concerns regarding the safety with PCA and its effectiveness have been well discussed in our manuscript and in the accompanying editorial by Dr. David Chestnut published in the September 1997 issue of Anesthesiology. Our manuscript has clearly suggested that epidural analgesia provides better analgesia than PCA during labor and that epidural analgesia should be preferred to PCA. However, in view of the complete satisfaction expressed by 65-70% of women in the PCA group and minimal counterfeit, it was reasonable to state that PCA is also an effective method of pain relief during labor. Professor Rosen has supported this view in his letter to the editor. Regarding neonatal outcome, we have also clearly indicated in our manuscript that our primary purpose in the study was to evaluate the rate of cesarean section. However, in our manuscript there was no difference between the two groups with regards to immediate neonatal outcome in terms of objective criteria such as Apgar scores and cord pH, PCO₂, and PO₂ except for an increased requirement for naloxone in the PCA group. Further, we had enough power to draw this conclusion (to determine a two-tailed significant difference of 0.05 between a mean \pm SD umbilical arterial pH of 7.30 ± 0.06 and 7.25 ± 0.06 in the two groups for 80% power, only 24 acid base measurements per group are required). We did not compare neonatal neurobehavioral changes, so interpreting it either way would be incorrect. The clinical significance of subtle differences in neurobehavior is unclear. Overall our manuscript and the accompanying editorial clearly suggest that PCA using meperidine is a reasonable option in some circumstances.

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Preinsertion Pulmonary Artery Catheter Flushing

To the Editor—Since the introduction of central venous and pulmo-
mary artery catheters into clinical practice, foreign body embolization
from either guidewires or catheter shearing has been reported. We
present this occurrence as the first reported case of potential foreign
body embolization unrelated to wires or catheter shearing, during pul-
monary artery catheter insertion.

A 63-yr-old man was scheduled for sternal debridement, rewiring, and
flap closure. The patient had sustained a recent myocardial infarction
and a ventricular fibrillation arrest before surgery. He was now 10 days
status after coronary artery bypass grafting (CABG), which was compli-
cated by postoperative congestive heart failure (CHF). Current medica-
tions included lasix, digoxin, vasotec, potassium chloride, and aspirin.
Given the patient’s significant cardiac history and also the magnitude
of the surgery, pulmonary artery catheter insertion was planned. After
Fig. 1. Paceport® pulmonary artery catheter with 2.6 cm × 1 mm cylindrical core extruded from the right ventricular lumen (Lot No. 3S12A 7522, Model No. 93A-931H-7.5F).

securing adequate intravenous and intraarterial access, the right internal jugular vein was cannulated uneventfully. While routinely flushing the catheter ports before insertion into the patient, a yellow core was noted to be protruding from the distal right ventricular lumen of a Paceport® pulmonary artery catheter (Baxter, Irvine, CA). This core did not come out with flushing using the flush valve of the transducer setup, rather it took a manual “power flush” from a syringe to fully dislodge the core from the distal right ventricular lumen (Fig. 1). The core was a solid cylindrical object that was 2.6 cm in length, 1 mm in diameter, with smooth flat ends. The core appeared to be made of the same yellow plastic used to make the catheter itself. This core was most likely a remnant from the manufacturing process. Another catheter was obtained, flushed, and inserted without incident. The rest of the anesthetic and surgery were uneventful.

We were fortunate that the lumen core was extruded during the flushing of the catheter ports before insertion into the patient. Had this not occurred until the catheter was inserted into the patient, it could have resulted in any one of several complications. Closer examination of the catheter revealed no additional problems. The manufacturer has no reports of a similar nature involving the same model and lot number. Moreover, the manufacturer has implemented several process improvements to prevent the type of occurrence reported herein. The Product Insert Data Sheet, included with each catheter, strongly recommends preinsertion testing of the catheter.

It is imperative that we pay close attention to the pulmonary artery catheter itself when flushing the ports because they may contain more that just heparinized saline solution.

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References


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In Reply.—Baxter appreciates the opportunity to provide additional comment regarding the above referenced manuscript. As a responsible medical device company, Baxter is interested in learning of experiences with its products. The occurrence reported in the manuscript is an isolated one. Baxter has no reports of a similar nature involving the same model and lot number of the device. In the interest of patient safety, Baxter implemented several process improvements to prevent the type of occurrence reported. Of importance, Baxter’s product insert data sheet, which is included with each catheter, recommends preinsertion testing of the catheter.

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