bupivacaine is difficult to justify. On the contrary, sales and revenue from liposomal bupivacaine continue to grow, undoubtedly owing to an aggressive marketing campaign. We have seen first-hand the results of this campaign, as many physician colleagues who are grounded in evidence-based medicine have shifted their practice to anecdotal medicine, insisting that liposomal bupivacaine exhibits superior pain control and leads to clinically significant reduced length of stay. Perhaps the dose influences their observations: We suspect that they are injecting the maximum dose of liposomal bupivacaine (i.e., 266mg), while using a lower dose of nonliposomal bupivacaine. We have additionally observed that liposomal bupivacaine usage tends to be an institutional decision: If the hospital system has decided to purchase this expensive product, then surely it ought to be used. And so, as more practices and hospitals are infiltrated with liposomal bupivacaine despite an absence of strong evidence, where do we go from here?

As physicians, we must strive to practice evidence-based medicine and use evidence such as that presented by Ilfeld et al. and Hussain et al. to defend against inappropriate and wasteful healthcare costs. However, distinguishing between objective data and marketing bias may pose a challenge for providers seeking to stay abreast of the current evidence in their field, given the prominent role the industry plays in medical education. Not too long ago, we witnessed the perils of the pharmaceutical industry’s influence on medical education and practitioners and how this ultimately helped fuel the opioid epidemic. Nowadays, in our determination to optimize postoperative pain control with nonopioid alternatives, are we repeating the missteps of the past by allowing the industry to again influence our practice without high-quality evidence? The continued intersection of the industry with medical education places us at risk of propagating non–evidence-based practices that may translate into little benefit, potential unforeseen harm, and unnecessary costs on an already taxed healthcare system.

Competing Interests

The authors declare no competing interests.

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DOI: 10.1097/ALN.0000000000003882

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(Accepted for publication June 14, 2021. Published online first on July 13, 2021.)

The corresponding author of the original article referenced above has read the letter and does not have anything to add in a published reply.

Personal Protective Equipment: Comment

To the Editor:

uskin et al. describe how personal protective equipment used because of COVID–19 impairs the performance of anesthesia clinicians and teams. They detail how reductions in the senses of sight, sound, and touch challenge
CORRESPONDENCE

We thank Dr. Johnstone for his insightful comments, in response to our article, about the practical implications of the loss of smell caused by personal protective equipment. We agree that healthcare professionals commonly rely on their sense of smell to facilitate diagnosis of some conditions. As Dr. Johnstone reported, the first indication that a vaporizer is leaking might be when a member of the surgical team detects the characteristic odor of a potent volatile anesthetic. Physicians' use of smell has also been evaluated to diagnose pseudomonas infections, and this ability might also be impaired by personal protective equipment.

N95 filtering facepiece respirators are designed to eliminate 95% of “most transmissible” particles from inspired air but are not resistant to oil and do not eliminate nuisance organic vapors. A person wearing an N95 mask may therefore be able to detect the odor of potent volatile anesthetics. P100 filters remove 99.97% of airborne particles, are strongly resistant to oil, and are commonly used with elastomeric half-facepiece respirators. Manufacturers commonly include a layer of activated carbon to eliminate nuisance organic compounds. Although there are currently no studies of how various respirators affect a person's ability to detect volatile anesthetic agents (e.g., from a leaking vaporizer), a P100 filter would reasonably be expected to eliminate the odor of sevoflurane in low concentrations. In the authors' personal experience, P100 filters are highly effective in eliminating other offensive odors that may be found in the operating room.

We agree that use of the extensive personal protective equipment that is required to care for patients with COVID-19 or other respiratory illnesses can impair the user’s sense of smell. This reduction in a critical sense suggests an important research opportunity. New technologies that detect contamination by volatile organic compounds or infectious agents without relying on a person’s ability to detect an odor may help users of personal protective equipment during a future pandemic. As we stated in our review, personal protective equipment impairs human performance in sometimes unpredictable ways. We thank Dr. Johnstone for pointing out yet another opportunity for improvement.

Competing Interests

The authors declare no competing interests.

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DOI: 10.1097/ALN.0000000000003902

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(Accepted for publication June 25, 2021. Published online first on August 3, 2021.)

Personal Protective Equipment: Reply

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Competing Interests

The authors declare no competing interests.

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DOI: 10.1097/ALN.0000000000003901

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Clinical Use of Lactate Measurements: Comment

To the Editor:

We read with great interest the recent review discussing the clinical interpretation of lactate measurements by Drs. Pino and Singh.1 The article covered the topic in great detail; however, there are several points which we also feel warrant inclusion when discussing this topic.

An additional source of lactate in surgical patients can be from red blood cell transfusion. As the authors state, red cells are completely anaerobic because they lack mitochondria and require glycolysis for maintenance of adenosine triphosphate. As a result, stored red blood cells are a source of lactate from transfusions; lactate levels and resultant patient lactate loads can increase with the use of stored red blood cells. Although many patients have adequate capability to metabolize lactate loads from transfusion, this may not be the case in massive transfusions, liver transplantation, or pediatric cardiac surgery. In particular, sudden massive boluses of red blood cell–based lactate in pediatric cardiac surgery from older stored red blood cells can significantly impact more traditional and conventional interpretations of lactate concentration.2

Additionally, endogenous overproduction of lactate may represent more than a metabolic waste product. Proponents of the concept contend that lactate can function as a metabolic glucose regulator and regulator of insulin release and insulin resistance.3 Brooks4 reviews compelling evidence that endogenous L-lactate is actively involved in aerobic intermediary energy metabolism and signaling effects through hydroxycarboxylic acid receptor 1. In the population with obesity, new evidence suggests hormonally triggered overproduction of lactate is an essential and obligatory feature of adipocytes, even in the absence of hypoxia.5 Adipocyte lactate production may therefore alter interpretations of elevated lactate levels in obesity and insulin resistance, although this concept is too recent for correlative studies in the perioperative period to define the magnitude and significance of these metabolic pathways.

The authors state that typical measurements of lactate, as conventionally performed with a blood gas machine, do not detect D-lactate, which has been shown to produce numerous deleterious effects. Some formulations of lactated Ringer’s solution have historically contained DL lactate,6 and although most formulations today are likely to contain only L-lactate, compositions may vary between countries and manufacturers. Additionally, although the consumption of fermented foods may contribute only a small proportion of total lactate as D-lactate, the presence and significance of D-lactate in pathologic states is not limited to gut ischemia or short bowel syndrome. Small intestinal bacterial overgrowth, in the absence of short bowel syndrome, has been associated with D-lactic acidosis with central nervous system symptoms. In fact, half of probiotic strains have been shown to ferment carbohydrates to D or DL lactate and may be responsible for gastrointestinal and central nervous system symptoms from D-lactic acidemia in some patients.7 Because D-lactate levels are not customary measurements, it is conceivable that many cases can be missed with routine laboratory investigations.

Competing Interests

The authors declare no competing interests.

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DOI: 10.1097/ALN.0000000000003905

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