DISCUSSION

Methods developed to separate 99mTc from 99Mo were recently reviewed by Dash et al. (12). In this study, we showed that ChemMatrix resins were efficient at selectively trapping 99mTc without retaining molybdenum in the presence of 4N NaOH, with quantitative elution using water. The overall Na^{99m}TcO₄ purification was implemented using a compact commercial automated synthesis system designed for use with disposable kits. We achieved high trapping efficiency with reproducible results using cyclotron-produced ^{99m}Tc. It should be noted that as the counts in the waste solution were not corrected for the presence of isotope contaminants that are known to occur with cyclotron-produced technetium (such as 99Mo and 95Nb) and were effectively separated from ^{99m}Tc, the recovery percentage of ^{99m}Tc was likely slightly underestimated. Because of the experimental design using cyclotron-produced pertechnetate, we did not evaluate the fraction of molybdenum recovered in the waste vial after technetium extraction by the ChemMatrix resin. Although molybdenum is not retained in high-pH, high-salt solutions by biphasic exchange chromatography, the determination of precise recovery yields will require further studies by testing this system with low-specificactivity ⁹⁹Mo solutions that are quantified before and after purifi-

The entire process is amenable to meeting good manufacturing practice–compliant production. This method is applicable for separation of $^{99\mathrm{m}}\mathrm{TcO_4}^-$ from any solution containing excess molybdate ions and should be suitable for use with low-specific-activity $^{99}\mathrm{Mo}$, with minor modifications to recirculate the $^{99}\mathrm{Mo}$ solution for daily reuse.

CONCLUSION

Cross-linked PEG beads were highly effective at separating and purifying ^{99m}Tc from low-specific-activity molybdate solutions, with quantitative trapping and elution of ^{99m}Tc. ^{99m}Tc-pertechnetate was purified in high yields using a simple disposable kit system.

DISCLOSURE

The costs of publication of this article were defrayed in part by the payment of page charges. Therefore, and solely to indicate this fact, this article is hereby marked "advertisement" in accordance with 18 USC section 1734. This work was supported by the Natural Sciences and Engineering Research Council, the Canadian Institutes of Health Research, and Natural Resources Canada. A provisional application for a patent was filed for some of the material presented in this article. No other potential conflict of interest relevant to this article was reported.

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Erratum

In the article "Preclinical Evaluation of 3- 18 F-Fluoro-2,2-Dimethylpropionic Acid as an Imaging Agent for Tumor Detection," by Witney et al. (*J Nucl Med.* 2014;55:1506–1512), reference to part D of Figure 1 is missing from the legend. The legend for part D is as follows: "(D) Effect of exogenous 19 F-FPIA and 19 F-FAC on intracellular metabolite concentrations of acyl-carnitine esters as analyzed by liquid chromatography–mass spectrometry (n=4)." We regret the error.