with currently available MR techniques [5, 41]. Given that microSPECT provides these added benefits, while also delivering a functional analysis equivalent to state-of-the-art microCT, microSPECT should be considered as a valid alternative to other preclinical imaging modalities when studying mouse models of cardiac disease.

Acknowledgements

All work was performed by the Duke Center for In Vivo Microscopy, an NIH/NIBIB Biomedical Technology Resource Center (P41 EB015897). Special thanks to Yi Qi for help with animal setup, and to Sidney Simon and Sally Zimney for editorial assistance.

Supplementary data

Supplementary material for this article can be found online at http://www.civm.duhs.duke.edu/4DmicroSpectCT2013/.