Fast Multiparametric 3D Spine MRI: Color Me Dixon!

Kenneth L Weiss¹, Judd M Storrs², and Manohar S Roda²

¹Radiology, University of Miami Miller School of Medicine, Miami, FL, United States, ²Radiology, University of Mississippi Medical Center, Jackson, MS, United States

Purpose

Optimize single and multi-station 3D Dixon¹ spine MRI; and develop multiparametric color encoding techniques to facilitate interpretation of the information rich opposed-phase (OP), in-phase (IP), and derived fat (F) and water (W) series.

Methods

Composed multi-station (head to tail) and single large FOV volumetric sagittal Dixon series were acquired at 3T (TR 4.1-4.2 ms, TEs 1.2 -1.4 and 2.4 -2.6 ms) on 48 -128 channel systems, maximizing available array coils to optimize S/N. Novel multiparametric color encoding programs were created in Matlab to combine OP, IP, and W series so that tissues containing only W signal (e.g. brain, cord, discs) appear gray scale, F appears yellow (red + green), tissues or "Indian ink artifact" containing both W and F appear red-magenta, and tissues exhibiting significant T2*decay appear cyan-green. To achieve this in RGB color space; IP, OP, and W series were respectively assigned to the red (R), green (G), and blue (B) channels. To achieve this in Cielab (L*a*b*) color space, designed to more closely parallel human perception, we set: L = IP + OP, a = (IP - OP)/(IP = OP), and b = (IP + OP - 2W) / (IP + OP). Sagittal color composites and subsequent coronal and axial reformats were generated from near mm³ isotropic volumes. (Fig 1)

Results

Rapid high-resolution volumetric Dixon imaging of the spine and other structures was achieved at 3T, with short echo times, high BW, and short acquisition times mitigating motion and susceptibility artifact. Even in the presence of metallic ACD&F hardware (T2* effects appreciated as cyan-green), fat water separation was true and adjacent disc pathology well appreciated. Both RGB and L*a*b* color composites behaved similarly, were intuitive (e.g. yellow marrow – Y, red marrow – R), and facilitated interpretation. (Fig 1)

Conclusion

Multiparametric color-encoded fast Dixon spine imaging demonstrates significant promise and warrants further systematic investigation.

References

- 1. Dixon, W.T. (1984). Simple proton spectroscopic imaging. Radiology 153, 189–194.
- Bray TJP, Singh S, Latifoltojar A, Rajesparan K, Rahman F, Narayanan P, et al. Diagnostic utility of whole-body Dixon MRI in multiple myeloma: A multi-reader study. PLoS ONE. 2017;12(7): e0180562.
- Kather JN, Weidner A, Attenberger U, Bukschat Y, Weis C-A, Weis M, et al. Color-coded visualization of magnetic resonance imaging multiparametric maps. Sci Rep. 2017 23; 7:41107.
- 4. Weiss KL, Richards CR, Sun D, Weiss JL. Subminute Fat-Water-Separated Dual-Echo Automated Spine Survey Iterative Scan Technique. American Journal of

Neuroradiology. 2009 Nov 1;30(10):1840-6.

5. Boll DT, Marin D, Redmon GM, Zink SI, Merkle EM. Pilot Study Assessing Differentiation of Steatosis Hepatis, Hepatic Iron Overload, and Combined Disease Using Two-Point Dixon MRI at 3 T: In Vitro and In Vivo Results of a 2D Decomposition Technique. American Journal of Roentgenology. 2010 Apr 1;194(4):964–71.



Figure 1