## Entropy of T2-weighted Imaging and Apparent Diffusion Coefficient of Uterine Leiomyoma in Prediction of Leiomyoma **Volume Reduction Following Uterine Artery Embolization**

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## **Introduction:**

Uterine artery embolization (UAE) has become an alternative option for symptomatic leiomyomas<sup>1</sup>. Many recent studies have investigated the role of MRI features of leiomyomas before UAE for prediction of leiomyoma volume reduction (VR) following UAE<sup>2, 3, 4</sup>. However, many of the results were contradictory and there is lack of a consensus on which is the most reliable feature to predict the outcome after embolization<sup>2, 3, 4</sup>. In this work, we for the first time employed entropy of T2-weighted imaging as the morphological parameter, together with apparent diffusion coefficient (ADC) as the functional parameter, to determine their utility for predicting the leiomyoma VR after UAE.

## **Materials and Methods:**

In this prospective study, 11 patients (age range 29 to 56 years; mean 42 years) with symptomatic uterine leiomyomas who underwent pelvic MRI including diffusion weighted imaging (DWI) before and 6 months after UAE were included. Pre-UAE and post-UAE MRI was all obtained on a 3.0 T system (HDxt; GE healthcare) equipped with a phased-array pelvic coil. Imaging sequences include fast spin-echo T2-weighted imaging, axial DWI using a single-shot spin-echo echo-planar sequence (b-values = 0, 1000 s/mm<sup>2</sup>) and axial contrast-enhanced T1-weighted imaging using a 3D volumetric interpolated technique (LAVA) prior to and after administration of intravenous Gd-DTPA contrast.

The volumes of each leiomyoma before and after UAE were determined using software ITK-SNAP on contrast-enhanced T1weighted images, and the percentage change in volume was calculated. Entropy of T2-weighted imaging and ADC before UAE were assessed using the following equations:  $ADC = -\ln (Sb/S0)/b$ , where b is the diffusion-sensitizing factor (b-value), and Sb and S0 the signal intensity at a non-zero b-value and zero b-value, respectively; Entropy =  $\sum i(-pi)(\log(pi))$ , where pi represents the probability of signal intensity (SI) i in the image and is calculated by dividing the pixel number of each SI by the total pixel number.

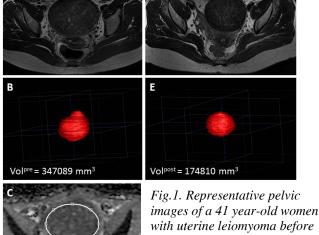
A total number of 16 leiomyomas larger than 2 cm in diameter were evaluated. The mean leiomyoma volume before UAE was 72.6 cm<sup>3</sup> (range 7.3–347.1 cm<sup>3</sup>), while the mean volume 6 month after UAE was 34.6 cm<sup>3</sup> (range 1.5–174.8 cm<sup>3</sup>), resulting in a mean leiomyoma VR of 58.9% (range 25.8%-95.0%). The mean ADC of leiomyomas was  $1.37 \times 10^{-3}$  mm<sup>2</sup>/s (range  $1.05 \times 10^{-3}$ )  $2.32 \times 10^{-3}$  mm<sup>2</sup>/s) and the mean entropy of T2-weighted imaging

was 5.36 (range 4.62–5.91) before UAE. ADC and entropy were significantly correlated with VR (Pearson correlation r = 0.61, P = 0.012; r= 0.73, P = 0.001). On multiple regression analysis, a combination of ADC and entropy constituted the best model for determining leiomyoma VR using the Akaike information criterion. For predicting  $\geq 50\%$  VR, receiver operating characteristic (ROC) curve analysis showed that the cutoff value of ADC was 1.39  $\times$  10<sup>-3</sup> mm<sup>2</sup>/s (sensitivity 45.5%, specificity 80.0%) and the cutoff value of entropy was 5.15 (sensitivity 90.9%, specificity 60.0%).



**Results:** 

Entropy of T2-weighted imaging and ADC of leiomyomas were significantly correlated with VR after UAE. A combination of entropy and ADC may have predictive value for VR after UAE.



(A-C) and after UAE(D, E).A,D: T2-weighted images; B,E: 3D volumetric maps; C:ADC map

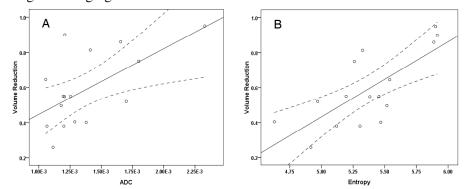


Fig.2. Relationships between ADC (A), entropy (B) and VR, respectively

## **References:**

- 1. Deshmukh SP,et al. Radiographics 2012;32(6):E251-E281.
- 2. deSouza N, et al. Radiology 2002;222(2):367.
- 3. Harman M, et al. Acta Radiol 2006;47(4):427-435.
- 4. Hecht EM, et al. J Magn Reson Imaging 2011;33(3):641-646.