

# Oxygen-enhanced lung imaging using rapid acquisition of T<sub>1</sub>-maps during free breathing

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## Purpose/Objective:

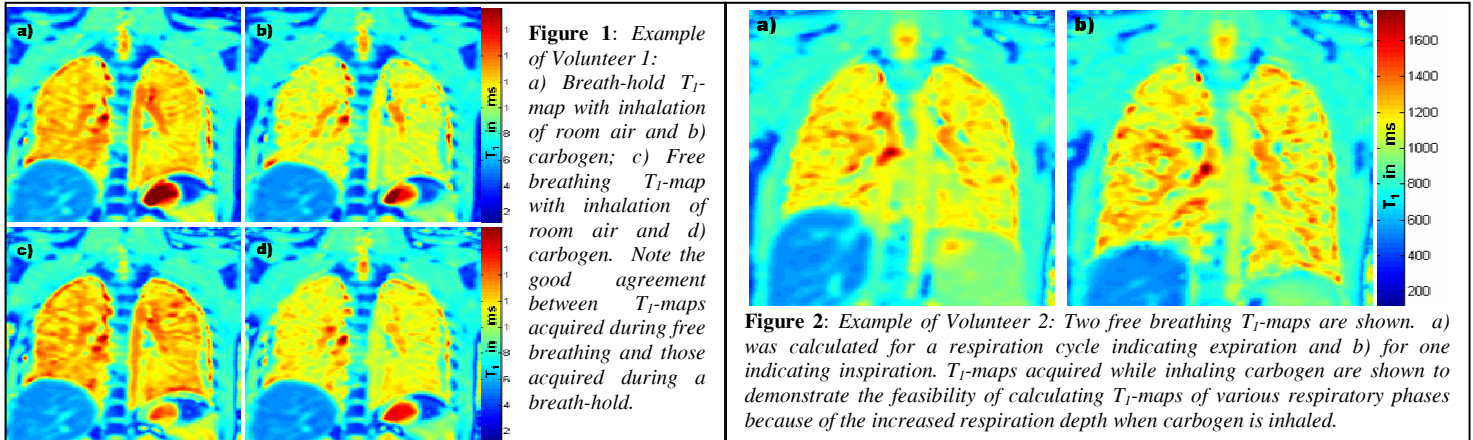
Oxygen-enhanced (OE) lung MRI was first proposed by Edelman et al. in 1996 [1] using a simple T<sub>1</sub>-weighted approach. The main drawback of this method is its lack of quantification and the potential for misinterpretation [2]. As an alternative, Jakob et al. proposed the acquisition of quantitative T<sub>1</sub>-maps [2]. Since T<sub>1</sub>-mapping requires the acquisition of several images, all of which should be obtained during the same breathing cycle, the measurement is usually performed during a breath-hold of several seconds. Since several slices under various breathing gas compositions often have to be acquired more than once, repeated breath-holds are necessary, which can cause severe discomfort in some patients. Therefore, this study presents a technique for rapid acquisition of T<sub>1</sub>-maps during free-breathing, suitable for quantitative OE lung imaging.

## Materials/Methods:

Seven healthy volunteers were examined on a 1.5 T whole-body scanner (Vision, Siemens, Germany). Lung T<sub>1</sub>-mapping was performed using a technique based on IR Snapshot FLASH [2]. Imaging parameters were TE=1.4 ms, TR=3.5ms, FA= 7°, FOV=(400mm)<sup>2</sup>, ST=15mm and an image matrix of 64x128, resulting in a total acquisition time of 4 seconds for all 16 images. The imaging protocol consisted of two phases: Firstly, 10 breath-hold T<sub>1</sub>-maps were acquired as a reference for comparison with the new technique. Secondly, 15 T<sub>1</sub>-map data sets were acquired during free breathing with a delay time of 3 seconds between consecutive T<sub>1</sub>-maps. In a retrospective gating procedure, the diaphragm position in each image was assessed. Images showing identical diaphragm positions were selected out of the 15 data sets in order to obtain a single data set where all images are within the same respiratory cycle. This data set was used for T<sub>1</sub>-map calculation. Because the short delay times used do not allow for full recovery of the magnetization between consecutive inversion pulses, the T<sub>1</sub>-values were corrected using the post-processing technique described in [3]. To demonstrate the feasibility for OE lung imaging, the procedure described above was performed while the subjects were breathing room air and again during inhalation of carbogen (95% oxygen, 5% carbon dioxide).

## Results:

T<sub>1</sub>-map acquisition during free breathing was better tolerated by the volunteers, especially when inhaling carbogen, where the diaphragm position in consecutive breath-holds varied substantially in some subjects. T<sub>1</sub>-maps acquired during free breathing were in good agreement with breath-hold acquired T<sub>1</sub>-maps in all cases (mean deviation of 1.5%). As an example, free-breathing T<sub>1</sub>-maps and breath-hold T<sub>1</sub>-maps of subject 1 are shown in Figure 1. In Figure 2, the feasibility of calculating T<sub>1</sub>-maps in various respiratory phases by choosing different retrospective navigator positions is demonstrated. The results for all volunteers are depicted in Table 1.



| Volunteer           | 1    |      | 2    |      | 3    |      | 4    |      | 5    |      | 6      |      | 7    |      |
|---------------------|------|------|------|------|------|------|------|------|------|------|--------|------|------|------|
|                     | Air  | Carb | Air  | Carb | Air  | Carb | Air  | Carb | Air  | Carb | Air    | Carb | Air  | Carb |
| T <sub>1</sub> BH   | 1213 | 1110 | 1240 | 1124 | 1214 | 1102 | 1231 | 1100 | 1339 | 1196 | 1234   | 1103 | 1182 | 1063 |
| T <sub>1</sub> free | 1253 | 1134 | 1220 | 1149 | 1217 | 1188 | 1225 | 1141 | 1373 | 1240 | 1237.0 | 1122 | 1172 | 1046 |
| % deviation         | 3.2  | 2.1  | -1.6 | 2.2  | 0.2  | 7.2  | -0.5 | 3.6  | 2.4  | 3.6  | 0.2    | 1.6  | -0.9 | -1.7 |

**Table 1:** T<sub>1</sub> values of the whole lung are shown for breath-hold T<sub>1</sub>-maps and free breathing T<sub>1</sub>-maps for all seven volunteers. The bottom row shows the mean deviation of the free breathing T<sub>1</sub>-maps from the breath-hold T<sub>1</sub>-maps in the lung.

## Conclusions:

The technique described provides for rapid and accurate T<sub>1</sub>-mapping during free breathing. T<sub>1</sub>-maps can be calculated for various respiratory phases and the diaphragm positions of several T<sub>1</sub>-maps, for instance acquired during the inhalation of room air and carbogen, are in good alignment as the same retrospective navigator placement is used. However, this can be problematic for different breath-hold acquired T<sub>1</sub>-maps as they can vary substantially in diaphragm position [4]. Since various respiratory phases can be calculated using the same data set, changes in T<sub>1</sub> caused by respiration can be easily studied. The entire measurement, i.e. acquiring 15 data sets with a 3-second delay time, takes less than 2 minutes. However, the measurement time can be decreased even more by further reduction of the delay time and by using less than 15 data sets, for instance when only T<sub>1</sub>-maps indicating expiration are needed. Thus, rapid free breathing quantitative OE T<sub>1</sub>-mapping is perfectly suited for routine clinical use.

## References:

- 1) Edelman RR, et al [1996] Nature Medicine 11:1236;
- 2) Jakob PM, et al [2001] JMRI 14:795;
- 3) Arnold JFT, et al [2004] MAGMA 16:246;
- 4) Molinari F, et al [2006] Invest Radiol 41 :476;