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Hyperoxia Challenge in Healthy and Anemic Subjects using BOLD MRI
Chau Vu, Julie Coloigner, and John Wood

Synopsis
This study explores the brain’s response to 100% oxygen inhalation in anemic subjects using BOLD MRI. Hyperoxic challenge has previously been used to identify brain regions with increased oxygen extraction fraction from inadequate perfusion. After controlling for changes in peripheral oxygen saturation, hyperoxic BOLD response was not significantly different between sickle cell disease patients, non-sickle anemic patients and healthy controls. Therefore, our results suggest that chronically anemic patients do not have increased oxygen extraction fraction from inadequate resting oxygen delivery under resting conditions.

Introduction
Hyperoxia is a popular medical technique employed in the emergency room for cases of stroke, cardiac arrest, respiratory failure and major trauma. It is also frequently used in MR brain studies to investigate cerebral metabolic rates and oxygen extraction. The addition of 100% oxygen increases oxygen content by approximately 7% in patients with normal hemoglobin levels and 11% in patients with chronic anemia syndromes such as sickle cell disease or thalassemia. If there are brain regions with flow limitation and higher oxygen extraction (OEF) under baseline conditions, hyperoxia would be expected to increase BOLD signal to a greater degree than normally perfused brain tissue. Therefore, in this study, we used a hyperoxic challenge to determine whether patients with chronic anemia syndromes are flow-limited under resting conditions.

Methods
A total of 82 subjects were studied; the demographics and hematologic results are detailed in Table 1. All subjects underwent an MR study using Philips 3T Achieva with an 8-element head coil accompanied by complete blood work.

MR acquisition: For each subject, anatomical 3D T1 MR and BOLD MR were acquired. Anatomical 3D T1-weighted images were acquired with the following parameters: TR = 8.20ms, TE = 3.77ms, flip angle = 8°, in-plane resolution = 1mm × 1mm, FOV = 256mm × 256mm, 160 slices and slice thickness = 1.0mm. BOLD images: TR = 2000ms, TE = 50ms, flip angle = 90°, in-plane resolution = 2.3mm × 2.3mm, FOV = 220mm × 220mm, 26 axial slices, slice thickness = 5mm and 150 volumes.

Hyperoxia challenge: At the start of the BOLD experiment, patients wore a breathing mask connected to a source of pressurized room air. After 50 seconds, the breathing circuit was switched to 100% oxygen gas until the end of the experiment.

Image processing: The BOLD datasets were pre-processed with FMRIB Software library (FSL) with a pipeline described elsewhere. Ratio in individual normalized BOLD signal between hyperoxia and normoxia was computed as illustrated in Figure 1. For linear regressions and statistical analyses, Ratio_{hyperoxia} was log-transformed to ensure a normal distribution.

Results
There was no statistically significant difference in BOLD hyperemic response between the three participating groups ($p = 0.11$). Hyperemic response was significantly higher in the grey matter than the white matter, while external veins exhibited the largest BOLD changes.
Significant correlation was observed between $\text{Ratio}_{\text{hyperoxia}}$ and $\Delta S_pO_2$ in Figure 2 ($r^2 = 0.10$, $p = 0.0072$). After regressing $\Delta S_pO_2$ from our model, the residuals did not show any between-group difference ($p = 0.26$) as shown in Figure 3. The residuals were also independent of sex, age, transfusion status, cell-free hemoglobin and all components of the complete blood count on univariate regressions.

**Discussion**

We used a hyperoxic challenge to probe for evidence of increased OEF in chronically anemic patients. Published works have validated increased OEF in flow-limited conditions such as in carotid artery stenosis as a compensation mechanism for impaired oxygen carrying capacity. Therefore, if SCD patients were flow-limited with increased OEF under resting conditions, we would have expected the hyperoxic BOLD effect to be larger in SCD subjects.

However, our results demonstrated no significant difference in the hyperoxic effect between SCD and control subjects after controlling for $\Delta S_pO_2$. Therefore, our results were not consistent with previous reports suggesting increased OEF in SCD. Additionally, our results also did not support previous hyperoxia experiments done in mice. However, these experiments did not control for the differences in oxygen saturation changes in response to hyperoxia. In our study, most of the hyperoxic difference could be explained by a lower resting $S_pO_2$ in the SCD subjects, so correcting for saturation was necessary to disentangle the OEF effect from the saturation effect. The differences in resting $SpO_2$ could be secondary to right shift in the hemoglobin dissociation curve or lung disease, both of which are relatively common in SCD patients.

In conclusion, we were unable to demonstrate evidence of increased OEF and flow limitations under resting conditions for SCD and chronically anemic patients. 100% oxygen does not appear to provide a useful imaging contrast, in contrast to carbon dioxide challenges.

**Acknowledgements**

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**Figures**
Figure 1: Representative global BOLD time course (blue) and the least-squares fit (red) during the hyperoxia challenge. Percent signal change $\Delta$BOLD was computed as the percent difference between the initial baseline and the stable value during hyperoxia.
Figure 2: The significant correlation between $\Delta$BOLD and $\Delta S\text{O}_2$ shows a large fraction of the hyperoxic response was derived from the change in resting arterial saturation during 100% oxygen inhalation.

Figure 3: Group mean maps for the hyperoxic BOLD effect show similar grey matter – white matter differentiation in all three subject groups.

Figure 4: Boxplot of percent signal change log-transformed $\text{Ratio}_{\text{hyperoxia}}$ shows no significant difference between the three subject groups.
<table>
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<th></th>
<th>ACTL (N=14)</th>
<th>CTL (N=34)</th>
<th>SCD (N=33)</th>
<th>p-value (SCD vs. CTL)</th>
<th>p-value (ACTL vs. CTL)</th>
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<td><strong>Age</strong></td>
<td>22.8 ± 6.8</td>
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<td><strong>Sex</strong></td>
<td>7 F, 7 M</td>
<td>25 F, 9 M</td>
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<td><strong>Hemoglobin</strong></td>
<td>11.4 ± 2.7</td>
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<td><strong>Hemoglobin S</strong></td>
<td>--</td>
<td>--</td>
<td>51.1 ± 30.8</td>
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<td><strong>Resting SpO$_2$</strong></td>
<td>95.8 ± 0.64</td>
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Table 1: Patient demographics and hematologic marker results.

**References**