Qualitative and quantitative lung perfusion imaging of children with congenital diaphragmatic hernia at 3T: initial results

F. G. Zoellner1, K. Zahn2, T. Schaible3, S. O. Schoenberg3, L. R. Schad4, and K. W. Neff1
1Computer Assisted Clinical Medicine, Heidelberg University, Mannheim, Germany, 2Dept. of Pediatric Surgery, University Medical Center Mannheim, Heidelberg University, Mannheim, 3Dept. of Paediatrics, University Medical Center Mannheim, Heidelberg University, Mannheim, 4Institute of Clinical Radiology and Nuclear Medicine, University Medical Center Mannheim, Heidelberg University, Mannheim, Germany

Introduction
Congenital diaphragmatic hernia (CDH) occurs in approximately one in 2000 to one in 4000 live births. Thereby, left-sided hernias are much more frequent (> 85%) than right-sided or bilateral hernias [1, 2]. In CDH, lung hypoplasia and secondary pulmonary hypertension are the major causes of death. After early surgical repair of CDH, long-term follow-up of paediatric patients is necessary. Recent studies involving MRI in the pre- and postnatal evaluation of CDH-patients comprised fetal MRI with lung volume determination and postnatal cerebral MRI, extra- and intracranial MR angiography [3], in particular after extracorporeal membrane oxygenation (ECMO-therapy) with carotid artery repair, and morphological lung analysis. To the best of our knowledge, quantitative perfusion imaging of the lung in CDH has not been utilized so far. Therefore, the aim of this study was to investigate whether dynamic contrast enhanced MR imaging (DCE-MRI) of the lung in survivors after CDH-repair at 3.0T is feasible.

Materials and Methods
Ten children (mean age 2 ± 0.2 years, 5 male, 5 female, 9-12 kg body weight) after CDH-repair were examined using a 3.0T scanner (Magnetom Trio, Siemens Healthcare Sector). Each patient underwent surgery to correct for CDH within the first 10 days after birth and was enrolled in our long-term follow-up program. Eight children had a left-sided hernia and two a right-sided hernia. Imaging was performed using a time-resolved angiography with stochastic trajectories (TWIST) sequence [4] with parameters TR/TE/FA=2.55ms/0.95ms/20°, FOV=320x260mm², parallel imaging (GRAPPA) factor 3 and 96 slices. Matrix size was 256x256 resulting in an isotropic voxel resolution of 1.3x1.3x1.3mm³. TWIST view sharing was set to 15% central region and 20% sampling density in the outer region. Temporal resolution was 3sec per volume. In total 25 volumes were acquired. 0.05 mmol/kg body weight of contrast agent (Dotarem, Guerbet, France) was administered manually after the fifth volume acquired. Patient related information was removed from the data set before offline processing. Image quality was assessed by calculating the signal-to-noise ratio (SNR) in pre-contrast images as well as peak SNR (PSNR) during the first pass [5]. Regions of interest (ROI) were placed in the apical lung parenchyma avoiding the inclusion of larger pulmonary arteries and veins. Quantification of lung perfusion was performed using a pixel-by-pixel deconvolution approach implemented in PMI 0.4 [6]. For calculations, the arterial input function was determined by placing a ROI in the pulmonary artery main stem. To access differences between left and right lungs, ROIs were placed in one slice on the calculated pulmonary blood flow (PBF) maps and average regional PBF (rPBF) was extracted.

Results
For all ten patients lung perfusion measurements could be obtained. Average SNR of the pre contrast images was 6.4±0.7. PSNR was 31.6±8.9. Figure 1 shows MIP image of time-resolved TWIST MRA, T2-weighted HASTE image and maps of PBF and pulmonary blood volume (PBV) of one of our patients. Table 1 demonstrates average PBF values and their standard deviation of ROIs placed in the left and right lung of the patients. In nine out of ten hypoplastic lungs a significant difference in the PBF compared to the contralateral side could be observed. Mean rPBF for the hypoplastic lungs was 34.3 ± 18 ml/100ml/min and 89.7 ± 27 ml/100ml/min for the contralateral side. In one case no differences could be detected (cf. Tab. 1, patient 4), however the patient is status post left-sided CDH repair.

Discussion
DCE-MRI of the lung in two-year old patients is feasible at 3.0T. Ipsilateral lung hypoplasia with reduced perfusion is reflected by significant lower rPBF values compared to the contralateral lung. MRI of lung is challenging as of low signal reception from the lung parenchyma. PSNR is only five times higher than baseline SNR. This might explain why the standard deviations within the ROI measurements of calculated PBF values are high. Certainly, also respiratory movement might contribute to the variations within the data, since no motion correction was applied so far. In our approach a higher spatial resolution was realized than having a high temporal resolution. In adult DCE-MRI of lungs, temporal resolutions between 1 and 1.5 sec are reported, however at lower in plane resolution and thicker slices. In a recent study, a temporal resolution of up to 3 sec was found not to influence the deconvolution analysis [5]. In conclusion, DCE-MRI of the lung in CDH can help characterizing lung hypoplasia initially and in long-term follow-up of children after CDH-repair.

References