

Optimizing Diffusion Weighted Imaging in Neonatal Vascular Territory Injuries.

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Introduction: Diffusion-weighted imaging (DWI) has been widely used to assess acute cerebral injury, but studies for the determination of the optimum value for the diffusion sensitivity factor, b , have mainly been performed in adult stroke patients.^{1,2} In neonatal brain, it has been suggested that the optimum b value to image ischemic injury is around 800s/mm².³ We studied the effects of varying the b factor during DWI assessment of neonatal focal ischemic brain injury.

Methods: The study group consisted of 8 neonates who presented with focal seizures within the first few days of life and a focal vascular territory abnormality on DWI. All patients underwent clinical examination on a 1.5T Signa Scanner which included multiple Diffusion Weighted Imaging (DWI) scans, each with a different b value.⁴ The sequence parameters were: TR =7500ms, TE = min for chosen maximum b value, $b = 0$ and at least 2 of 500, 700, 1000, 1500 or 2000 s/mm², single shot, FOV 200x200mm, matrix 128x128, 4 to 5mm thick, slice gap 0-1mm, NEX =3. The diffusion tensor images were eddy current corrected and processed to create DWI and ADC maps. Intra-subject data sets at different b values were co-registered using FLIRT software. The lesion was outlined manually on all DWI images using Analyze 5.0. A Region of interest (ROI) was also drawn in the normal tissue of the cerebellar hemisphere. The cerebellum was chosen for a normal ROI as it was not involved in any case and therefore served as a control ROI. The average DWI signal intensity (S) was determined in each ROI at all b values. ΔS was calculated as the difference in signal between DWI lesion and normal cerebellar tissue. The signal-to-noise ratio (SNR) was defined as S/σ , S being the signal intensity of the lesion and σ represented noise as calculated from an ROI placed in an area in the air surrounding the head that was free of artifact. The contrast-to-noise ratio (CNR) was defined as $\Delta S/\sigma$. The lesion was also outlined on each ADC map generated from the DWI at each b value. The mean ADC was recorded and the size of the ROI created was compared to the size of ROI of the corresponding DWI map.

Results: DWI maps: In all patients, noise remained approximately the same at all b values. SNR dropped with increasing b values. ΔS and CNR increased when b increased from 500 or 700 to 1000 s/mm² and then dropped with further increase of b to 1500 and 2000 s/mm². (Figure 1) The size of the lesion on the DWI maps decreased with increasing b values.

ADC maps: The size of the lesion on the ADC maps was the same at all b values and smaller than the lesion outlined on the corresponding DWI maps. This difference between ADC and DWI lesion volume was more pronounced at $b = 700$ s/mm² and almost nonexistent at $b = 1500$ s/mm². (Figure 2) The observed ADC values were lower at higher b values for both normal tissue and areas of infarction.

Discussion: At a $b = 1000$ s/mm², CNR is highest and SNR, although lower than at 700s/mm², is still acceptable. T2 shine through effects are lower at a $b = 1000$ s/mm² than at 700s/mm² and therefore the lesion size on the DWI map is closer to the ADC size. Although DWI lesion size was closer to ADC size at a $b = 1500$ s/mm², the lower SNR and CNR made this b value less favorable. In conclusion if a single b value is used to image focal ischemic injuries in neonates, a b value of 1000s/mm² is recommended. Also given changes in ADC that occur with b , the b value of the acquired data should be reported when providing quantitative ADC data.

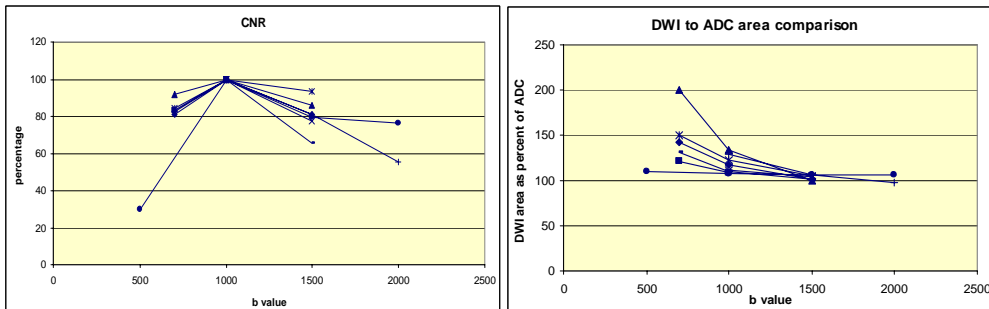


Figure 1: a. CNR at different b values presented as percentage of CNR at $b = 1000$ s/mm². b. Size of the lesion on the DWI maps at different b values presented as percentage of size on corresponding ADC map.

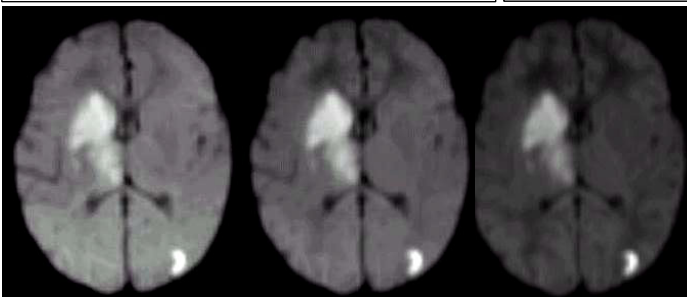


Figure 2: Newborn DWI images at b values of 700 (left), 1000 (middle) and 1500 (right) s/mm².

References: 1. Pereira RS, et al, J Magn Reson Imaging. 2002 May;15(5):591-6. 2. Meyer JR, et al, AJNR Am J Neuroradiol. 2000 Nov-Dec;21(10):1821-9. 3. Kingsley PB, et al, Magn Reson Med. 2004 May;51(5):996-1001. 4. Sorensen AG, et al, Radiology 1996 May;199(2):391-401.