Time course of wallerian degeneration after ischaemic stroke revealed by diffusion tensor imaging

G Thomalla, V Glauche, C Weiller, J Röther

Wallerian degeneration (WD) of descending fibre tracts after ischaemic stroke is a well known phenomenon reflecting severe fibre tract damage. WD represents a uniform answer to injury within the central and peripheral nervous systems, and disintegration of axonal structures within the first days after injury is followed by infiltration of macrophages, degradation of myelin after several weeks, and finally, fibrosis and atrophy of the affected fibre tracts. After ischaemic stroke it usually takes two to four weeks before WD can be detected by conventional magnetic resonance imaging (MRI), where the main pathological finding is a hyperintensity on T2-weighted images along the affected tracts in the chronic stage, weeks to months after stroke.

Recently, diffusion tensor imaging (DTI) has opened up new possibilities of imaging fibre tracts in the brain by estimation of the whole diffusion tensor—which provides information on the predominant direction and degree of water diffusion and thus allows conclusions to be drawn about the microstructural properties of tissue. The degree of anisotropy of diffusion reflects the integrity and the degree of organisation of the fibre tracts within the brain. DTI has been used to study WD in the chronic stage of stroke and it has been shown that fractional anisotropy (FA) was reduced along the pyramidal tract on the affected side below the primary lesion over time is shown in fig 1. In a previous study, we used DTI to study early WD of the pyramidal tract after acute ischaemic stroke. We found decreases in FA and characteristic changes in the principal diffusivities (eigenvectors), reflecting early WD in the cerebral peduncle of the affected side as early as 2–16 days after ischaemic stroke; at the same time T2-weighted images and maps of the orientationally averaged diffusivity did not reveal obvious changes.

No longitudinal DTI studies of WD have been published. Here we report the findings from diffusion tensor images obtained for two of our patients at three different time points during the time course from the early subacute to the chronic stage of stroke.

METHODS
Magnetic resonance images were acquired on a 1.5 Tesla MR system (Magnetom Vision, Siemens, Erlangen/Germany). A high resolution T1-weighted image data set (voxel size 1 × 1 × 1 mm) was acquired. For DTI we used a single shot STEAM sequence with matrix size 56 × 64, field of view 192 × 192 mm, slice thickness 3 mm without interslice gap, and voxel size 3 × 3 × 3 mm. Diffusion sensitising gradients (b = 750 s/mm²) were applied along six directions, and one image without diffusion weighting (b = 0 s/mm²) was obtained. The diffusion tensor (D) for each voxel was calculated, and maps of eigenvalues, averaged diffusivity (Dav) and FA were generated using SPM99 in Matlab 5.3 (The MathWorks, Natick, MA). Three dimensional regions of interest (ROIs) were manually defined for each side covering the medial anterior cerebral peduncle between the hypothalamus and the pons. Eigenvalues (λ₁, λ₂, λ₃), FA, and Dav were calculated within the ROI, and ratios between values of the affected and unaffected side were determined (rλ₁, rλ₂, rFA, rDav). Details of the MRI protocol and postprocessing have been reported elsewhere.

RESULTS
Two patients with striatocapsular infarction were examined at three time points after stroke (case 1: 12, 104, and 288 days after stroke; and case 2: 5, 35, and 92 days after stroke). DTI revealed a clear pattern of progressive structural changes, which corresponded well with histological findings on the temporal evolution of WD and with DTI findings of WD in the chronic stage after stroke as described above. In both patients the rDav decreased continuously (from 0.84 to 0.75 and from 0.83 to 0.62) and the rDav increased slightly (from 0.98 to 1.11 and from 0.96 to 1.02). Ratios for the second (rλ₂) and third (rλ₃) eigenvalues markedly increased over time in both patients (rλ₂: from 1.07 to 1.29 and from 1.02 to 1.10; rλ₃: from 1.05 to 1.27 and from 1.06 to 1.28).

An example of progressive FA decrease along the pyramidal tract below the primary lesion over time is shown in fig 1. Corresponding structural changes are clearly visible on the coregistered high resolution T1-weighted image in the late chronic stage, where a hypointensity resulting from degeneration of descending tracts in the mediolateral cerebral peduncle is easily identified.

DISCUSSION
We longitudinally studied the course of WD in two patients by DTI and found a pattern of progressive structural degeneration which corresponds well to findings from

Abbreviations: Dav, averaged diffusivity; DTI, diffusion tensor imaging; FA, fractional anisotropy; WD, wallerian degeneration
WD of the pyramidal tract after ischaemic stroke is known to reflect severe pyramidal tract damage associated with persisting impairment of motor functions. In patients with ischaemic stroke and motor impairment, the degree of WD of the pyramidal tract has been shown to be correlated to motor scales at different time points. In both our patients the DTI findings of a typical pattern of progressive WD were associated with persisting moderate to severe hemiparesis.

DTI allows evaluation of the time course of WD from the early subacute to the chronic stage. The findings on imaging reflect the different stages of WD that are well known from experimental and histological studies. Thus, DTI offers a way to detect and monitor the time course of severe degeneration of the pyramidal tract and may be a helpful tool in forecasting and monitoring recovery in patients with ischaemic stroke.

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REFERENCES
manifestation of MS. A crossed quadrant field defect may rarely occur as an initial quadrantanopia. Furthermore, our patient demonstrates that parietal lobe produced the left superior quadrantanopia, while optic radiations. The demyelinating lesion in the right temporal and left parieto-occipital lobes (fig 2). A (MRI) of the brain demonstrated high signal abnormality in the right temporal region and left parieto-occipital area, which account for the corresponding field defects.

lesions required to produce them and the tendency for rapid recovery in the vast majority of cases. Another factor that may limit the occurrence of homonymous field defects in MS is the tendency for lesions to occur along venules and non-fibre tracts. The chances of a checkerboard field defect appearing in MS are reduced by: (1) the rarity of the homonymous defects in this condition; (2) the necessity for bilateral field defects; (3) the occurrence of quadrantanopic defects rather hemianopic ones; and (4) the need to have one lesion above and one lesion below the representation of the horizontal meridian.

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Figure 1 Humphrey visual fields [30-2] demonstrate a left superior and right inferior quadrantanopia consistent with a crossed quadrant pattern of field defects.

Figure 2 Axial sections of brain MRI. Fluid attenuation inversion recovery [FLAIR] images demonstrate high signal abnormality (arrows) in the right temporal region and left parieto-occipital area, which account for the corresponding field defects.

A 43 year old previously healthy woman presented with headache and blurred vision. Visual acuity was 20/30 in both eyes, with normal colour vision, pupil, and funduscopic examinations. Computerised threshold perimetry (fig 1) revealed a left superior and right inferior homonymous quadrantanopia. Magnetic resonance imaging (MRI) of the brain demonstrated high signal abnormality in the right temporal and left parieto-occipital lobes (fig 2). A diagnosis of multiple sclerosis (MS) was established when the patient developed recurrent neurologic dysfunction.

Crossed quadrant hemianopias are rare field defects that are usually secondary to ischaemia and are typically attributed to bilateral injury to the calcarine fissure. Such defects are uncommon since bilateral quadrantopias secondary to vascular disease are usually either both superior or inferior, owing to watershed ischaemia or embolic disease. From 1891 to 1994 only nine cases with crossed quadrant field defects were reported worldwide and no-one suffered from MS. In 1995, the first report of crossed quadrant hemianopia occurring in a patient with clinically definite MS was documented to be secondary to lesions in the trigone areas bilaterally. We are not aware of any further reports of checkerboard field defects since 1995. In our patient, crossed-quadrant field defects resulted from bilateral lesions in the optic radiations. The demyelinating lesion in the right temporal lobe produced the left superior quadrantanopia, while the lesion in the left parietal area produced the right inferior quadrantanopia. Furthermore, our patient demonstrates that a crossed quadrat field defect may rarely occur as an initial manifestation of MS.

Symptomatic homonymous field defects in MS are uncommon, and may be related to the large retrochiasmal

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