Semiautomatic quantitation of macrophages in human renal biopsy specimens in proteinuric states

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Abstract

**Aims**—To develop and validate a rapid and economical semiautomated approach to the measurement of immunostainable tissue components which is applicable to routine diagnostic practice. To apply this approach to the measurement of macrophages in renal biopsy specimens in nephrotic states, as protein in the renal tubules may induce macrophage infiltration, and the morphology of macrophages in tissue sections does not lend itself to cell counting.

**Methods**—Macrophages were identified by immunostaining with a pan-macrophage marker, followed by digital image capture and analysis using a macro procedure written for the freeware image analysis program NIH-Image.

**Results**—The method was rapid, robust and accurate to within the limits imposed by sampling error inherent in the use of small needle biopsy specimens. Very few macrophages are found in normal kidney (mean volume fraction (±95% confidence limits) 0.04% (0.02%)) but infiltration of macrophages was detected in minimal change nephropathy (0.29% (0.12%)) and in membranous glomerulonephritis (0.42% (0.11%)). A statistically significant correlation was found between macrophage volume fraction and weight of proteinuria in minimal change nephropathy but not in membranous glomerulonephritis. Correlations were found in both diseases between macrophage volume fraction and serum creatinine at time of biopsy.

**Conclusions**—The equipment is inexpensive and measurement takes less than one minute per biopsy specimen. The results indicate that macrophage infiltration is part of the pathological process in minimal change nephropathy and membranous glomerulonephritis. The correlation with creatinine at time of biopsy suggests that renal impairment in minimal change nephropathy may result from infiltration by immunologically active cells and not merely from haemodynamic changes in nephrons. However, the correlation is not close, indicating that the relation between macrophage infiltration and disease severity is not a simple one.

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The role of the macrophage in the induction of fibrosis is becoming clearer in a variety of disease processes. In the kidney, most emphasis has been placed on macrophage infiltration in the glomerulus, with consequent production of cytokines and induction of excess deposition of extracellular matrix. The renal interstitium has been studied comparatively rarely, despite the acknowledged importance of interstitial fibrosis in renal impairment, and despite the observation of foam cells in the interstitium in biopsy specimens from patients with heavy proteinuria from a variety of causes.

Recently, there have been several attempts to predict the course of chronic human renal disease from the renal biopsy specimen by measuring the numbers of myofibroblasts in the interstitium. This has been quite successful in IgA nephropathy and in membranous glomerulonephritis, both of which are normally rather unpredictable in their course. However, the method used, manual counting of cells after immunocytochemical detection, is laborious and for this reason alone is unlikely to gain wide acceptance in routine diagnosis.

In general terms, one may postulate the following as a typical sequence of events in a fibrotic reaction: an insult leads to macrophage infiltration, which results in cytokine production, which stimulates proliferation and activity of matrix producing cells (such as myofibroblasts). If this is accepted, it seems possible that quantitation of interstitial macrophages rather than myofibroblasts might produce an earlier and potentially more sensitive marker of the severity of the insult to the kidney, although resolution may occur if the infiltration is short-lived. Therefore, we sought to measure the level of macrophage infiltration in renal biopsy specimens from adult patients with idiopathic membranous nephropathy, with minimal change nephropathy and, as a control group,
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Clinical data collected included serum creatinine concentrations and the 24 hour urine protein output at the time of biopsy.

**IMMUNOSTAINING**

Paraffin wax sections (3 μm) were dewaxed, and treated with 6% H2O2, to block endogenous peroxidase. Sections were exposed to 0.1% trypsin (Difco) in 0.12% CaCl2, pH 7.8, for 25 minutes at 37°C. A standard indirect immunoperoxidase method was used with PGM1 (Dako), a pan-macrophage marker, as the primary antibody applied overnight at 4°C at a dilution of 1 in 50 in TRIS-buffered saline. A counterstain was omitted in order to facilitate subsequent image analysis.

**IMAGE ANALYSIS**

Sections were viewed on a Zeiss photomicroscope III with a JVC TK 120E video camera attached. The camera was linked with a Y/C cable to the in-built framegrabber board of an Apple Macintosh 7100/80 AV microcomputer, and images were imported directly to the freeware image analysis program NIH-Image*, using the 'Plug-in digitizer' software PhotoShop-compatible plug-in.

Sequential images were grabbed using the ×10 objective, moving along the central line of each biopsy specimen from one end of the available cortex to the other, without overlapping. The presence of glomeruli in the fields was ignored. Renal medulla was not measured.

To calculate the volume fraction of the macrophages, each image was converted to grey-scale, and then a threshold was applied at a level which distinguished between the stained macrophages and the unstained background.

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*The image analysis program NIH-Image was written by Wayne Rasband (e-mail: wayne@helix.nih.gov) and is distributed as freeware (at: http://rsb.info.nih.gov/nih-image/ or by anonymous ftp from zippy.nimh.nih.gov).


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The proportion of black to white pixels in the image (excluding the edges) was then calculated as a percentage. As sequential fields were assessed, a running mean of the macrophage area fraction was maintained automatically. Processing of a typical image is illustrated in fig 1.

A set of macros was written in the NIH-Image programming language to automate this process, such that sequential fields can be measured with just two keystrokes per field. These macros are available as a simple text file, free by e-mail on request to pnfl@le.ac.uk.

**METHOD VALIDATION**

To test the reproducibility of the image analysis method, 10 sequential sections were cut from a single renal biopsy specimen. These were immunostained with PGM1 on 10 separate occasions, each occasion separated by at least three days. The volume fraction of macrophages in the resulting set of 10 slides was then measured, and the measurements were repeated the following day.

**STATISTICAL ANALYSIS**

The regression coefficients were calculated as part of the graphing process by the program CA-Cricket graph III; p values for these linear correlations were calculated using Systat. Comparisons between the mean macrophage volume fractions in normal biopsy specimens and those in the disease groups were made using Student’s t test (unpaired), as calculated by Microsoft Excel; 95% confidence limits were also calculated using Excel.

**Results**

**METHOD VALIDATION**

The results of measurements of 10 replicate sections are shown in fig 2. The means, quartiles and maximum/minimum figures show little variation between the two occasions on which they were measured. The overall variation is considerably less than one might expect with a quantitative immunocytochemical method, and we felt use of the method in the main part of the study was justified.

In most biopsy specimens, the exact depth of the cortical tissue could not be judged. In a few specimens, however, the renal capsule and cortico-medullary junction were both visible. Such cases were too few for statistical analysis, but there seemed to be a gradient of macrophage numbers which decreased from outer to inner cortex (fig 3). This finding probably justifies further study; if confirmed, it suggests that sampling error, rather than measurement error, is likely to be the biggest source of random variation with this method.

**MACROPHAGE VOLUME FRACTIONS**

The mean macrophage volume fractions are shown in fig 4. Macrophage infiltration in minimal change nephropathy and in membranous nephropathy was significantly greater.
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Although the mean level of macrophage infiltration in membranous glomerulonephritis was greater than in minimal change disease, the difference was not statistically significant (p = 0.13).

There was a significant correlation between macrophage volume fraction and serum creatinine at the time of biopsy in membranous nephropathy and minimal change disease (fig 5). A significant correlation was also found with the 24 hour urine protein output in minimal change disease (fig 6). However, it is obvious from the scatter plots that the regression value is low and the correlation is poor. These correlations have only just reached significance, even though our automated measuring system permitted the analysis of relatively large numbers of specimens.

Discussion

These results confirm that a significant interstitial infiltrate of macrophages is a part of the pathological process in membranous nephropathy and in minimal change nephropathy. The existence of a correlation between macrophage volume fraction and serum creatinine in both diseases implies that immunological mechanisms may be involved in the reduced glomerular filtration rate which is commonly seen at presentation of both diseases. This is not surprising in the case of membranous glomerulonephritis; but in minimal change nephropathy, the presence of interstitial macrophages is not widely recognised as part of the disease process. Indeed, one argument which has been used against the puromycin aminonucleoside (PAN) model of minimal change nephropathy is that PAN induces an interstitial

Figure 5  Scatter graphs showing correlations between macrophage volume fraction and creatinine at time of biopsy in (A) normal specimens, in (B) minimal change nephropathy and in (C) membranous glomerulonephritis.

Figure 6  Scatter graphs showing correlations between macrophage volume fraction and 24 hour urine protein output immediately before biopsy in (A) normal specimens, in (B) minimal change nephropathy and in (C) membranous glomerulonephritis.

\[ r = 0.299 \quad p = \text{NS} \]

\[ r = 0.398 \quad p < 0.05 \]

\[ r = 0.393 \quad p < 0.05 \]

\[ r = 0.127 \quad p = \text{NS} \]

\[ r = 0.433 \quad p = 0.05 \]

\[ r = 0.234 \quad p = \text{NS} \]
infiltrate of macrophages, on which the renal impairment seen in the model depends. Our results suggest that this criticism is invalid, and also argue against the widely held view that any renal impairment in minimal change nephropathy has a haemodynamic cause, reflecting circulation volume changes in nephrosis, rather than an immunological one. Furthermore, it shows that the presence of a moderately heavy interstitial infiltrate of macrophages does not necessarily indicate the presence of an irreversible fibrotic process, as minimal change nephropathy rarely, if ever, progresses to renal fibrosis.

A close correlation has previously been reported between macrophage infiltration and proteinuria in mesangial proliferative glomerulonephritis with mesangial IgM. Tubular expression of monocyte chemoattractant protein-1 (MCP-1) is known to be increased in membranous glomerulonephritis and to correlate with the interstitial macrophage infiltrate. It has been suggested that the mere presence of large quantities of filtered protein in the tubules represents an insult to the tubular epithelium which can result in chemokine-induced macrophage infiltration. This certainly seems to be the case in the protein overload model of proteinuria in the rat. We have shown that MCP-1 is released by cultured tubular epithelial cells after the addition of protein to the medium bathing the apical surface of the cells.

However, macrophage infiltration is induced in many animal models of fibrotic disease, in the kidney and other organs. This includes the fibrosis caused by ureteric obstruction, where proteinuria is not an issue. Of course, we do not know for how long our patients had proteinuria before biopsy was performed; variation in this time may account for some of the variation in macrophage numbers, but it seems likely that other aspects of the disease process are involved in the variation we have seen. If assessment of macrophages in renal biopsy specimens is to be relevant to the severity and progression of disease, it is evident that we should be measuring macrophage function rather than simply their presence. Assessment of cytokines such as tumour growth factor β may be more promising.

There has been considerable interest recently in quantitative or semiquantitative approaches to predicting the rate of disease progression from renal biopsy specimens, both native and transplant. Most of these studies have used either semiquantitative scoring on a scale of 0 to 3 by a ‘blinded’ observer, or laborious manual morphometry, or cell counting. Having previously undertaken a study requiring point-counting, we have been very favourably impressed with the method described here, which can be applied to any tissue component that can be localised by immunostaining. The high speed and low expense of the method mean that if the clinical significance of such methods can be demonstrated, it will be easy to convert this information into a routine quantitative assessment—a feature currently lacking in most diagnostic histopathology reports.

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