Immunoglobulin and complement deposition in glomeruli of 756 subjects who had committed suicide or met with a violent death

J Varis, I Rantala, A Pasternack, H Oksa, M Jäntti, E S Paunu, R Pirhonen

Abstract

Aims—To study immune deposits in renal glomeruli.

Methods—Tissue was obtained from 756 necropsy cases from people who had committed suicide or met with a violent death. Glomerular immune deposits were examined by immunofluorescence microscopy and a light microscopy. The clinical histories of all the deceased were studied to ascertain reasons for the deposits.

Results—Immune deposits were found in glomeruli in 91 (12%) cases. In 52 (6.8%) cases mesangial IgA was observed as a solitary finding in 34 (4.5%), and was accompanied by other immunoglobulins in 18 (2.4%). Mesangial IgM was present in 19 (2.5%) and IgG in 11 cases (1.5%). Two cases had capillary IgG (0.3%). Light microscopic examination showed mesangial enlargement in eight of the cases with mesangial IgA. These included one with IgA glomerulonephritis diagnosed before death. Two cases with normal glomerular morphology and mesangial IgA deposits had clinical laboratory evidence of renal disease. In two subjects with normal glomerular morphology, mesangial IgM and microscopic haematuria were present. In one case with capillary IgG membranous glomerulonephritis was detected.

Conclusions—Ten cases had mesangial IgA together with morphological or clinical laboratory findings suggestive of renal disease. If all these are regarded as IgA glomerulonephritis, then its prevalence can be estimated at 1.3%. For IgM glomerulonephritis, a prevalence of 0.3% was deduced.

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Glomerular immune deposits can occur without clinical evidence of glomerulonephritis.^{1,2} The prevalence of these deposits is surprisingly high: 4% for mesangial IgA and 1% for IgM in unselected necropsy material from Singapore¹; and 4.8% for IgA deposits in a hospital necropsy study from Germany.² Biopsy specimens from cadaveric renal transplants have also shown glomerular deposits in 10.9% of cases studied on immunofluorescence.³ Not much is known about the reasons why immunoglobulins are deposited.

Methods

Over three years (1985-1987) 1813 forensic necropsies (540 men and 216 women) were performed at Tampere University Hospital. In 756 cases death had been caused by violent death or suicide. The age distribution by sex is shown in fig 1.

Specimens of cortical kidney tissue for immunofluorescence and light microscopy studies were obtained at necropsy from all of the deceased. Eighty per cent of cases were necropsied within seven days of death, and 99% within 16 days of death. We have already shown that immunofluorescence from immunoglobulins and complement in necropsy tissue can be reliably interpreted up to 17 days after death.⁴

For immunofluorescence studies, the specimens were embedded in Ames OCT compound and frozen in liquid nitrogen. The frozen tissue was cut, fixed in absolute acetone, and stained with fluorescein isothiocyanate (FITC) conjugated antisera to human IgA, IgG, IgM, C1q, and C3b. Controls included incubation in FITC conjugated non-specific antiserum and incubation in an unconjugated blocking antiserum before the FITC-conjugate was used. All antisera were obtained from Behringwerke AG (Marburg-Lahn, Germany).

For light microscopy, the tissue was routinely embedded in paraffin wax. The speci-

Institute of Clinical Sciences, University of Tampere, Finland

J Varis
A Pasternack
H Oksa
M Jäntti
E S Paunu
R Pirhonen

Department of Pathology, Tampere University Hospital
I Rantala

Correspondence to:
Juha Varis, Institute of Clinical Sciences, University of Tampere, Box 607, SF 33101 Tampere, Finland

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Figure 1 Age and sex distribution of study material.

Table 1 Location and appearance of different immune reactants in relation to predominant immunofluorescence finding

Predominant Immunofluorescence finding	IgA	IgG	IgM	C1q	C3b	N=
IgA	M G					34
	M G	M G				5
	M G		M G			4
	M G				M G	3
	M G			M G		1
	M G	M G	M G			2
	M G	M G		M G		1
IgG			M G			6
		C G				1
		M G		M G		3
	M G		M G	M G		1
		M G		M G		1
		C G	C G			1
			M G			1
IgM			M G			17
			M G	M G		1
		M G	M G			1
C1q				M G		5
C3b					M G	2

C G = granular fluorescence along capillary walls, M G = mesangial and granular fluorescence.

Clinical data were collected from necropsy documents, the files of Tampere University Hospital, and the Discharge Reporting System. This register, organised by the Finnish National Board of Health, covers all hospitals in the country. The local Pirkanmaa Kidney Register was also examined. By combining data from these sources, clinical information was available for 590 (78%) cases. The correlations between immunofluorescence findings and the diseases encountered were analysed using the χ^2 test.

Results

Of the 756 necropsies, immunofluorescence showed renal deposits of immunoglobulin in 91 (12%) (table 1). There was no correlation between the prevalence of positive immunofluorescence findings and the age or sex of the deceased. The immune deposits were granular, diffusely distributed, and located in mesangial areas except in two cases with capillary IgG (fig 2).

Mesangial IgA was found in 52 (6.9%) cases. Isolated IgA deposits were present in

mens with positive immunofluorescence findings were stained by haematoxylin and eosin, periodic acid Schiff, and Jones's periodic acid silver-methenamine methods.

Figure 2 Cases with immunofluorescence findings: their light microscopic findings and clinical data.

Table 2 Data on previous clinical finding in the kidney in 38 cases

Renal diseases	Cases in total study population	Cases with immunofluorescence deposits		
		IgA	IgM	IgG
Renal cysts	13	—	—	—
Renal trauma	6	1	—	—
Chronic pyelonephritis	6	—	—	1
Hydronephrosis	2	1	—	—
Kidney anomaly	2	1	—	—
Interstitial nephropathy*	3	—	—	—
Glomerulonephritis†	2	1	—	1
Unknown nephropathy	2	—	—	—
Renal scars in urography	1	—	—	—
Renal calculi	1	—	—	—

*Two cases with rhabdomyolysis and transient failure during life and one with unknown interstitial nephropathy.

†One case with IgA glomerulonephritis and one with mesangial and granular fluorescence, both with pre-mortem diagnosis. In three cases two renal diseases were detected in the same person; one with glomerulonephritis and duplex ureter, one with chronic pyelonephritis and slight hydronephrosis, and one with hydronephrosis and renal cysts. The disease mentioned first was regarded as the primary diagnosis.

34 (4.5%) specimens. IgA was accompanied by complement in four cases, and by other immunoglobulins in 14 (table 1).

Of the cases with mesangial IgA (fig 2), light microscopic examination showed mesangial enlargement in eight. In two other cases previous clinical data had indicated renal disease but this was not detected at light microscopy. Of these 10 cases one had IgA glomerulonephritis diagnosed before death. If the remaining nine diagnoses are also taken to be IgA glomerulonephritis, on the basis of the diffuse and global occurrence of IgA, together with morphological or clinical laboratory signs of renal disease (fig 2), a prevalence of 1.3% for IgA glomerulonephritis can be deduced.

Mesangial IgM was found in 19 (2.5%) and IgG in 11 (1.5%) cases. Isolated C1q was present in five (0.7%) and C3b in two (0.3%) cases (table 1). In all of these specimens the glomerular morphology was normal.

Two cases with IgM deposits had microscopic haematuria (fig 2). According to current opinion, these represent cases of IgM glomerulonephritis. This gives a prevalence of 0.3% for IgM glomerulonephritis.

Granular IgG deposits along the capillary basement membranes were seen in two cases (0.3%). One of these had heavy proteinuria, and membranous glomerulonephritis was seen at light microscopy.

Previous data on clinical renal findings were found in 38 cases (table 2).

Among the 590 cases with available clinical data the percentage of positive immunofluorescence findings was 11.7%; in the remaining 166 cases it was 13.2%. There were no differences in the prevalence of positive immunofluorescence findings among subjects known to have had a variety of non-glomerular diseases.

Discussion

We investigated the occurrence of glomerular deposits of immunoglobulins and complement by immunofluorescence in material selected from the population on the basis of sudden death. There were more men, with a predominance of middle-aged men and old

women (fig 1). In our study there was a delay between death and necropsy. We have already shown, however, that immunofluorescence can reliably show the presence of glomerular immunoglobulins and complement up to 17 days after death.⁴ At light microscopy, autolysis was evident but it did not affect the interpretation of glomerular morphology. In fact, the two cases with biopsy confirmed glomerulonephritis (IgA glomerulonephritis, membranous glomerulonephritis) were also diagnosable after death on the basis of typical immunofluorescence and light microscopic findings.

Sinniah has described immunofluorescence results in tissue from subjects who had met with a violent death,¹ and so has Waldherr *et al* in patients at necropsy.² Sinniah found mesangial IgA to be present in 4%; Waldherr and coworkers detected IgA in 4.8% of the necropsied patients. Curschellas *et al* found glomerular IgA in 6.9% of biopsy specimens from cadaveric grafts studied by immunofluorescence.³ These results agree with those of our study. The occurrence of glomerular IgA seems to be high in various populations. Deposition is probably not inevitably associated with any clinically apparent condition. If the cases with glomerular IgA, together with those that had morphological or clinical laboratory findings of renal disease, are regarded as IgA glomerulonephritis, a prevalence of 1.3% for IgA glomerulonephritis can be estimated.

In IgA glomerulonephritis IgA is almost invariably accompanied by complement deposition,⁵ which has been considered to be of central importance in the pathogenesis of both human and experimentally induced IgA glomerulonephritis.^{6,7} On the other hand, the quality of antigen is thought to have a critical role in the development of glomerulonephritis associated with IgA immune complexes.⁸ The IgA deposits found in this study were mostly solitary. Isolated IgA deposits have also been shown to occur in coeliac disease,⁹ liver diseases,¹⁰ and neoplastic diseases.¹¹ In the study by Curschellas *et al* complement was detected along with IgA in only one out of seven cases with glomerular IgA.³ In our electron microscopic studies electron-dense deposits were found in mesangial areas and they were morphologically similar to those seen in IgA glomerulonephritis.

These results suggest that IgA can be deposited mesangially without leading to overt glomerular damage. The deposits probably do not do any damage until some as yet undefined circumstance induces clinically apparent glomerulonephritis. Complement seems to be one of the linking factors between subclinical IgA deposition and IgA glomerulonephritis.

Sinniah found IgM deposits in 1% of his material.¹ The corresponding figure in our study was 2.5%. Although mesangial IgM deposition is a common non-specific phenomenon found in immunofluorescence studies, two cases with microscopic haematuria diagnosed before death were regarded as IgM

glomerulonephritis.¹² In our study IgA glomerulonephritis occurred five times as often as IgM glomerulonephritis. This finding agrees with that of Mustonen *et al*,¹³ who found that the ratio of IgA glomerulonephritis to IgM glomerulonephritis was 3:4.

As to the other immunofluorescence findings, they seemed to be randomly distributed among various clinical conditions. One case with capillary IgG deposits proved to be membranous glomerulonephritis. On the other hand, the presence of other immune reactants remained unexplained on the basis of the available background data.

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