Renal failure in a patient with chronic lymphocytic leukaemia treated with fludarabine

M P Macheta, L A Parapia, D R Gouldesbrough

Abstract
The case history of a man with refractory chronic lymphocytic leukaemia who developed acute renal failure following treatment with fludarabine is presented. A renal biopsy specimen showed features of mesangiocapillary glomerulonephritis, a rare occurrence in chronic lymphocytic leukaemia. The rapid cytotoxic action of fludarabine may result in the development of renal glomerular lesions when used to treat a well differentiated B cell malignancy.

(Keywords: Fludarabine, renal failure, chronic lymphocytic leukaemia)

Case Report
In 1987, a 60 year old man presented with lymphadenopathy and lymphocytosis. A diagnosis of chronic lymphocytic leukaemia (CLL) was reached following examination of bone marrow, and peripheral blood lymphocyte marker studies. Three years later, the patient developed increasing superficial lymphadenopathy and enlarged tonsils. This resolved completely after three months of treatment with chlorambucil and prednisolone. In 1992, lymphadenopathy recurred with increasing lymphocytosis (white blood cell count $10^4 \times 10^7\text{l}$). Lymphadenopathy progressed despite treatment with chlorambucil and prednisolone for three months. Treatment with fludarabine was then started at a dose of 25 mg/m$^2$ daily (43 mg/day) for five days every four weeks. The patient had a history of ischaemic heart disease and was taking frusemide, nifedipine and warfarin.

On admission for the fifth course of fludarabine, the patient complained of exertional dyspnoea, orthopnoea and oliguria. On examination, there were signs of congestive cardiac failure. There was no lymphadenopathy or hepatosplenomegaly. Urinalysis revealed proteinuria and haematuria. Standard laboratory analyses revealed the following: haemoglobin 11.6 g/dl; white blood cell count $6.4 \times 10^3\text{l}$; platelets $133 \times 10^3\text{l}$; sodium 139 mmol/l; potassium 4.1 mmol/l; urea 9.0 mmol/l; creatinine 145 $\mu$mol/l; urate 610 $\mu$mol/l (normal range 250–500 $\mu$mol/l); albumin 34 g/l; and globulin 28 g/l. A chest x ray film revealed cardiomegaly and interstitial pulmonary oedema. No casts were observed on urine microscopy. C3 $0.39\text{g/l}$ (normal range 0.75–1.65 g/l); C4 $0.08\text{g/l}$ (normal range 0.2–0.65 g/l); antibodies to glomerular basement membrane and double stranded DNA, antineutrophil cytoplasm antibody, and cryoglobulins were negative; no paraproteins were detected on serum electrophoresis; bone marrow examination revealed no evidence of CLL or lymphoma; and ultrasonography showed both kidneys and urinary tracts to be normal. Treatment with diuretics, fluid restriction and allopurinol was instituted. Renal function continued to deteriorate rapidly and 10 days later the patient's creatinine concentration was raised further: 652 $\mu$mol/l. Percutaneous renal biopsy was performed. There was no improvement following daily pulses of intravenous methylprednisolone (1 g) and oral cyclophosphamide (75 mg). Peritoneal dialysis was started and the patient is currently undergoing regular haemodialysis.

On light microscopy, the renal biopsy showed focal accentuation of glomerular lobular architecture, a focal and segmented increase in mesangial cellularity and focal thickening of glomerular basement membranes. Tubules and blood vessels appeared normal. Immunofluorescence microscopy revealed coarse granular deposits of C3 within the basement membrane in some glomeruli and mesangial deposition in others. Irregular granular deposits of IgM were also present in the basement membrane. On electron microscopy, there were patchy subendothelial electron dense deposits and elongated deposits in some areas of thickened basement membrane (figure). No mesangial cell interposition was identified and there was no evidence of interstitial cellular infiltrate.

Discussion
There have been 44 published reports of renal glomerular disease associated with CLL and well differentiated B cell non-Hodgkin's lymphoma. Mesangiocapillary glomerulonephritis (MCGN) is the most common glomerular lesion reported, occurring in eight of the 11 patients with CLL and nephrotic syndrome described by Moulin et al. However, the spectrum of glomerular lesions described is wide and includes membranous and minimal change glomerulonephritis, focal segmental glomerulosclerosis and amyloidosis. Although the pathological changes in this case were atypical they most closely resemble those present in type I MCGN. In this case complete remission of CLL had been achieved when renal failure developed. However, Korzets et al noted that the stage of CLL and onset of glomerular disease were not necessarily related. Immune complex deposition is thought to

Annette Fox
Haematology Unit and Department of Histopathology, Bradford Royal Infirmary, Duckworth Lane, Bradford BD26 6RP
M P Macheta
L A Parapia
D R Gouldesbrough

Correspondence to: Dr M P Macheta.
Accepted for publication 7 June 1994


181
be the initiating event in MCGN.\textsuperscript{6} CLL and well differentiated B cell non-Hodgkin’s lymphoma are associated with abnormalities of immunoglobulin production and autoimmunity. In those cases where glomerular disease develops there is often evidence of M-protein production by the proliferating B cell clone. In a series of patients with CLL and MCGN one third had circulating M-protein compared with 5–10% of all patients with CLL. Cryoglobulinaemia and evidence of complement consumption are also common.\textsuperscript{1} Monotypic immunoglobulin or light chain deposition has been demonstrated within glomeruli by immunofluorescence even in the absence of circulating M-protein, strengthening the aetiological link between monoclonal lymphoid proliferation and glomerular disease.\textsuperscript{2}

In the series of patients with CLL/well differentiated B cell non-Hodgkin’s lymphoma and glomerular disease reported by Moulin et al.\textsuperscript{10} of the 13 patients were treated with chlorambucil and prednisolone. In six of seven patients with MCGN, treatment induced complete remission of glomerular disease (chlorambucil alone was used in five patients). In seven of eight patients with renal failure there was a substantial improvement in renal function even though five of these patients had MCGN, a condition which normally has a poor prognosis. In one case where repeat renal biopsy was performed following treatment with chlorambucil there was improvement in the histological features of MCGN. All three patients with MCGN, or atypical membranous glomerulonephritis, and CLL described by Touchard et al.\textsuperscript{12} improved, attaining remission after treatment with chlorambucil and prednisolone. Spontaneous remission of glomerular disease associated with CLL may also occur\textsuperscript{4} and successful treatment with interferon has been described.\textsuperscript{5}

Fludarabine is a purine analogue. It is an effective, rapidly cytotoxic agent for the treatment of refractory CLL and well differentiated B cell non-Hodgkin’s lymphoma.\textsuperscript{7,8} The major reported side effects are myelosuppression and demyelination. Although renal failure caused by tumour lysis following treatment with fludarabine is well documented,\textsuperscript{9} direct nephrotoxicity or glomerular disease have not been described. The suppliers have received reports of five other patients with CLL/well differentiated B cell non-Hodgkin’s lymphoma in whom renal failure developed after treatment with fludarabine. None of these patients underwent renal biopsy. Two were similar to this case in that acute renal failure developed three weeks after the second course of treatment. The release of a large amount of leukaemic cell antigen as a result of the rapid cytotoxic action of fludarabine may predispose patients to the formation of the immune complexes responsible for glomerular injury. Acute renal failure is an unusual symptom of MCGN and in the case presented here cardiac and vascular insufficiency may also have contributed to the rapid decline in renal function.

Haematologists should be aware of the rare association of well differentiated B cell lymphoproliferative disorders with renal glomerular disease because, with increasing use of fludarabine, such cases may occur more frequently. Furthermore, the onset of renal glomerular disease may be a further indication for treatment in CLL and well differentiated B cell non-Hodgkin’s lymphoma given the excellent response observed in many cases.

Renal failure in a patient with chronic lymphocytic leukaemia treated with fludarabine.
M P Macheta, L A Parapia and D R Gouldesbrough

doi: 10.1136/jcp.48.2.181

Updated information and services can be found at:
http://jcp.bmj.com/content/48/2/181

These include:

**Email alerting service**
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/