Azathioprine versus mycophenolate mofetil for long-term immunosuppression in lupus nephritis: results from the MAINTAIN Nephritis Trial

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ABSTRACT

Background Long-term immunosuppressive treatment does not efficiently prevent relapses of lupus nephritis (LN). This investigator-initiated randomised trial tested whether mycophenolate mofetil (MMF) was superior to azathioprine (AZA) as maintenance treatment.

Methods A total of 105 patients with lupus with proliferative LN were included. All received three daily intravenous pulses of 750 mg methylprednisolone, followed by oral glucocorticoids and six fortnightly cyclophosphamide intravenous pulses of 500 mg. Based on randomisation performed at baseline, AZA (target dose: 2 mg/kg/day) or MMF (target dose: 2 g/day) was given at week 12. Analyses were by intent to treat. Time to renal flare was the primary end point. Mean (SD) follow-up of the intent-to-treat population was 48 (14) months.

Results The baseline clinical, biological and pathological characteristics of patients allocated to AZA or MMF did not differ. Renal flares were observed in 13 (25%) AZA-treated and 10 (19%) MMF-treated patients. Time to renal flare, to severe systemic flare, to benign flare and to renal remission did not statistically differ. Over a 3-year period, 24 h proteinuria, serum creatinine, serum albumin, serum C3, haemoglobin and global disease activity scores improved similarly in both groups. Doubling of serum creatinine occurred in four AZA-treated and three MMF-treated patients. Adverse events did not differ between the groups except for haematological cytopenias, which were statistically more frequent in the AZA group (p=0.03) but led only one patient to drop out.

Conclusions Fewer renal flares were observed in patients receiving MMF but the difference did not reach statistical significance.

INTRODUCTION

Lupus nephritis (LN) occurs in up to 60% of patients with systemic lupus erythematosus (SLE) and significantly impacts their survival.2 Randomised trials performed at the National Institutes of Health (NIH) indicated that long-term use of a combination of steroids and high-dose intravenous cyclophosphamide (CY) pulses was superior to steroids alone to prevent renal impairment.3–5 Based on these studies, the so-called ‘NIH regimen’ became the standard of care for LN for three decades, despite its many side effects such as a high rate of severe infection and premature ovarian failure.

Two different therapeutic approaches have been recently proposed. First, mycophenolate mofetil (MMF) was shown to be at least as efficacious as oral/intravenous CY to induce a good renal response at 6 months in several pivotal randomised studies.6–8 Although long-term data are still unavailable, MMF is now widely used to treat LN. A second approach, the ‘Euro-Lupus regimen’, consists of prescribing lower doses of intravenous CY for a short period of time followed by long-term immunosuppression with azathioprine (AZA). In a randomised trial, this regimen was shown to achieve results comparable to a high-dose intravenous CY treatment protocol9 10 with a very low rate of end-stage renal disease at 10 years.11 Nevertheless, even on long-term AZA many renal relapses were observed, as in other series from the literature.12 We therefore designed a randomised superiority trial (the MAINTAIN Nephritis Trial) comparing AZA and MMF as long-term immunosuppressive treatment of LN, after a short course of low-dose intravenous CY, in order to test the hypothesis that MMF would reduce renal relapses.

PATIENTS AND METHODS

Inclusion/exclusion criteria

Between July 2002 and March 2006, 105 patients were included in the MAINTAIN Nephritis Trial by 27 European centres. All the following inclusion criteria were to be met: age ≥14 years, SLE according to the American College of Rheumatology (ACR) classification criteria,13 24 h proteinuria ≥500 mg, biopsy-proven proliferative WHO class III, IV, Vc or Vd lupus glomerulonephritis (biopsy performed less than 1 month before entry in the protocol), contraception (or sexual abstinence for paediatric patients) and signed informed consent. None of the following exclusion criteria could be met: non-lupus related renal disease (such as microthrombotic disease associated with antiphospholipid syndrome), treatment with glucocorticoids (GCs) (>15 mg equivalent prednisolone/day) in the last month before entry into the study (except a very short-course high-dose oral GC treatment before referral), treatment with CY, AZA, MMF or ciclosporin A in the previous year, pre-existing...
Outcomes

First choice treatment in all patients with hypertension (diabetic-range proteinuria (≥3 g/day). ACEI were prescribed as the
inhibitors (ACEI) were prescribed in all patients with neph-
renal response at 3 months. Angiotensin-converting enzyme
MMF (target dose 2 g/day), according to randomisation per-
mg (fixed dose) within a 10-week period and were then given,
patients received six fortnightly intravenous CY pulses of 500
weeks, GCs were tapered by 2.5 mg prednisolone/day every
0.5 mg equivalent prednisolone/kg/day for 4 weeks. After 4
by oral GC treatment given on day 4 at an initial dose of
A detailed treatment protocol is provided in the supplemen-
tary material. Briefly, all patients received three daily 750 mg
A total of 105 patients with biopsy-proven proliferative LN
were randomised into the trial (intent-to-treat population). We
anticipated a renal flare rate of 35% at 5 years in the
AZA group. We defined the clinically meaningful difference
as a 10% renal flare rate in the MMF group. To detect such
a difference, 51 patients needed to be randomised in each arm to
obtain a power of 0.80 with an α level of 0.05.

Statistical analyses

Survival curves were computed using the Kaplan–Meier method
and were statistically tested with the log rank test. We calculated
the HR and their 95% CIs using the univariate Cox proportional
hazards model. Unpaired t tests, χ2 tests and Fisher’s exact tests
were used as appropriate. Serial data were compared within and
between groups by repeated measures analysis of variance, with
a ‘between groups’ and a ‘repeated measures’ comparison. All
analyses were by intent to treat.

RESULTS

Patient disposition, follow-up and drug exposure

A total of 105 patients with biopsy-proven proliferative LN
were randomised into the trial (intent-to-treat population). Their
demographics and baseline characteristics did not significa-
cantly differ between the two groups, as described in table 1. Of
note, most patients were Caucasians and had recent-onset
kidney disease. In all, 10% of the patients had renal impair-
ment (serum creatinine ≥1.4 mg/dl) and 39% had nephrotic-
proteinuria (≥3 g/day). As depicted in figure 1, 39% had nephrotic-
proteinuria (≥3 g/day). As depicted in figure 1, 3
The primary end point of the trial was time to renal flare, analysed by survival curves computed, after the Kaplan–Meier
method, on the intent-to-treat population. A renal flare was
declared as (i) the recurrence or the development of nephrotic
syndrome (serum albumin ≤3.5 g/dl and 24 h proteinuria ≥3 g;
this type of renal flare is further referred to as ‘nephrotic
syndrome’), (ii) renal impairment (≥33% increase of serum
creatinine within a 1-month period directly attributed to
lupus and confirmed 1 week later; flare referred to as ‘renal
impairment’) or (iii) a threefold increase of 24 h proteinuria
within a 3-month period accompanied by microscopic haem-
maturia (defined as a number of red blood cells (RBC) per
high power field superior to upper normal limit for the local
laboratory) and ≥35% reduction of serum C3 level within a
3-month period (this definition of renal flare was only appli-
cable to those patients with low-grade baseline 24 h proteinu-
ria (≥0.5 g and <1 g); this type of renal flare is further referred
to as ‘proteinuria increase’).

The secondary end points of the trial were the number of
severe systemic and benign flares (both defined as described
in the supplementary material), the number of patients with-
drawing GCs and achieving renal remission. Renal remis-
sion was defined as a serum creatinine ≤1.4 mg/dl and a 24 h proteinuria <1 g and a urinary RBC count <10/high power
field, according to clinical practice in 2001, when this trial was
designed. Of note, renal remission was not a prerequisite to
receive maintenance treatment with AZA or MMF and could
therefore be reached at any time during follow-up. Disease
activity was measured by the validated ECLAM14 and Systemic
Lupus Erythematosus Disease Activity Index (SLEDAI)15 scor-
ing systems. Patients could be dropped from the trial for any of
the following reasons: death, drug toxicity, pregnancy or
pregnancy wish, doctor decision, patient decision (consent
drawn).

Power calculation

The trial was designed as a superiority trial of MMF over AZA.
The primary end point (time to renal flare) was used for power
calculation. We anticipated a renal flare rate of 35% at 5 years in
the AZA group. We defined the clinically meaningful difference
as a 10% renal flare rate in the MMF group. To detect such
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between groups by repeated measures analysis of variance, with
a ‘between groups’ and a ‘repeated measures’ comparison. All
analyses were by intent to treat.
Mean (SD) follow-up of the intent-to-treat population was 48 (14) months.

All patients received 3 g of intravenous CY. To evaluate exposure to AZA/MMF, the maximal and minimal doses taken by each patient throughout the study were identified. Patients from the AZA and MMF group received a mean (SD) maximal daily dose of 124 (33) mg and 2.0 (0.2) g, respectively. The mean (SD) minimal daily doses of AZA and MMF were 100 (38) mg and 1.6 (0.6) g, respectively. Only four and five patients never reached the target AZA or MMF dose, respectively.

Efficacy

Time to renal flare, the primary end point of the trial, was compared by Kaplan–Meier survival curves computed for the intent-to-treat population. As shown in figure 2, time to renal flare was not statistically different in the MMF group (10 patients/53; 19%) compared to the AZA group (13 patients/52; 25%). Eight patients from the AZA group and six patients from the MMF group qualified for a renal flare based on the recurrence/development of nephrotic syndrome, two from the AZA group and four from the MMF group based on renal impairment and three from the AZA group based on proteinuria increase (see patients and methods for renal flare definitions). One patient allocated to AZA had two consecutive episodes of renal flare (nephrotic syndrome).

Time to severe systemic flares (p=0.99; 4 and 3 episodes in the AZA and MMF group, respectively), to benign flares (p=0.60; 23 and 26 episodes in the AZA and MMF group, respectively), to GC withdrawal (p=0.88; 21 and 20 patients in the AZA and MMF group, respectively), to GC/intravenous CY/MMF (p=0.99; 4 and 3 episodes in the AZA and MMF group, respectively), to renal remission (p=0.11; 47 patients in both groups) did not statistically differ.

A subset analysis was performed on patients who, at any time during follow-up, reached a significant renal response defined as a ≥50% reduction of 24 h proteinuria (n=98; 48 in the AZA group and 50 in the MMF group). Importantly, a similar number of patients had a renal flare in the AZA (n=7) and the MMF group (n=7) after having achieved at least a 50% drop in 24 h proteinuria. Time to renal flare, computed from the time a ≥50% reduction of 24 h proteinuria was achieved, did not differ by Kaplan–Meier analyses (p=0.97; data not shown).

Table 1  Demographics and baseline disease characteristics*

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>AZA (n=52)</th>
<th>MMF (n=53)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Race, n (C/A/B)</td>
<td>41/4/7</td>
<td>42/5/6</td>
</tr>
<tr>
<td>Gender, n (F/M)</td>
<td>48/4</td>
<td>48/5</td>
</tr>
<tr>
<td>Age in years, mean±SD</td>
<td>33±11</td>
<td>33±10</td>
</tr>
<tr>
<td>Past SLE renal disease, %</td>
<td>13</td>
<td>18</td>
</tr>
<tr>
<td>Previous GC use, %</td>
<td>53</td>
<td>47</td>
</tr>
<tr>
<td>Previous IS use, %</td>
<td>27</td>
<td>29</td>
</tr>
<tr>
<td>Serum creatinine, mg/dl (mean ± SD)</td>
<td>1.02±0.47</td>
<td>1.01±0.33</td>
</tr>
<tr>
<td>24 h proteinuria, g (mean±SD)</td>
<td>2.94±2.42</td>
<td>3.63±2.80</td>
</tr>
<tr>
<td>Serum albumin, g/dl (mean±SD)</td>
<td>3.01±0.75</td>
<td>2.97±0.66</td>
</tr>
<tr>
<td>Haemoglobin, g/dl (mean±SD)</td>
<td>10.96±1.98</td>
<td>10.93±1.63</td>
</tr>
<tr>
<td>Serum C3, mg/dl (mean±SD)</td>
<td>55±29</td>
<td>49±26</td>
</tr>
<tr>
<td>ECLAM score, mean±SD</td>
<td>6.95±1.83</td>
<td>6.41±1.86</td>
</tr>
<tr>
<td>WHO class, n (III/IV/Vc/Vd)</td>
<td>17/6</td>
<td>19/6</td>
</tr>
</tbody>
</table>

*None of the 14 baseline variables were statistically different between the 2 groups (x^2 or unpaired t tests).

Figure 1  Trial profile and patient disposition. AZA, azathioprine; GCs, glucocorticoids; ITT, intent to treat.

Figure 2  Kaplan–Meier probability analysis of renal flare. Patients were allocated to the ‘GC/intravenous CY/AZA’ group (circles) or the ‘GC/intravenous CY/MMF’ group (squares) by randomisation. Survival curves were statistically tested with the log rank test. Data are HR (95% CI). Numbers shown along the abscissa are the number of patients at risk in each group. Analyses were by intent to treat. Time point of reference for follow-up is from baseline. AZA, azathioprine; CY, cyclophosphamide; GCs, glucocorticoids; MMF, mycophenolate mofetil.
Adverse events

Seven patients (four from the AZA and three from the MMF group) doubled their serum creatinine at last follow-up. Two of them (one from each group) developed end-stage renal failure and required renal replacement treatment. Two patients (both from the MMF group) died, the first from *Legionella pneumophilia* sepsis at month 36, after a renal flare at month 3 and doubling of serum creatinine at month 24; the other died from SLE at month 45, after a severe systemic flare at month 44.

All other adverse events are listed in table 2. None was statistically more frequent in one group compared to the other, with the exception of haematological cytopenias, which were more frequent in the AZA (14 episodes: 11 leucopenia, 2 anaemia and 1 combination of leucopenia and anaemia) compared...
Table 2  Adverse events*

<table>
<thead>
<tr>
<th>Event</th>
<th>AZA</th>
<th>MMF</th>
</tr>
</thead>
<tbody>
<tr>
<td>Death</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Due to SLE</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Renal</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Doubling of serum creatinine</td>
<td>4</td>
<td>3</td>
</tr>
<tr>
<td>End-stage renal failure</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Infection</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Benign infection</td>
<td>14</td>
<td>21</td>
</tr>
<tr>
<td>Herpes zoster</td>
<td>5</td>
<td>4</td>
</tr>
<tr>
<td>Herpes simplex</td>
<td>0</td>
<td>2</td>
</tr>
<tr>
<td>Cytomegalovirus</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td>Chickenpox</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>Salmonella sepsis</td>
<td>2</td>
<td>0</td>
</tr>
<tr>
<td>Upper urinary tract infection</td>
<td>0</td>
<td>2</td>
</tr>
<tr>
<td>Sepsis of unknown origin</td>
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<td>0</td>
</tr>
<tr>
<td>Streptococcus pneumonia</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Gynaecological</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Leucopenia</td>
<td>11</td>
<td>2</td>
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<tr>
<td>Leucopenia and anaemia</td>
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<td>0</td>
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<tr>
<td>Anaemia</td>
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</tr>
<tr>
<td>Renal haematoma</td>
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<td>1</td>
</tr>
<tr>
<td>Psas bleeding</td>
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</tr>
<tr>
<td>Gastrointestinal</td>
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<td></td>
</tr>
<tr>
<td>Nausea/diarrhoea</td>
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<td>8</td>
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<tr>
<td>Hepatitis</td>
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<td>1</td>
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<tr>
<td>Central nervous system</td>
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<tr>
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<tr>
<td>Headaches</td>
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<tr>
<td>Antimalarial retinopathy</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>Skin</td>
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<tr>
<td>Drug-induced rash</td>
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<td>Alopecia</td>
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<tr>
<td>Gynaecological</td>
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<tr>
<td>Transient amenorrhoe</td>
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<tr>
<td>Gynaecological bleeding</td>
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<tr>
<td>Metabolic</td>
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<tr>
<td>Cushing</td>
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<tr>
<td>Diabetes mellitus</td>
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<tr>
<td>Cardiovascular</td>
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<tr>
<td>Angina pectoris</td>
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<tr>
<td>Cerebrovascular accident</td>
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</tr>
<tr>
<td>Renal vein thrombosis</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>Subclavian vein thrombosis</td>
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<tr>
<td>Bone</td>
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<tr>
<td>Avascular osteonecrosis</td>
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<tr>
<td>Osteopenia</td>
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</tr>
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<td>Rib fractures</td>
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<td>0</td>
</tr>
<tr>
<td>Cancer</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cervix carcinoma</td>
<td>2</td>
<td>0</td>
</tr>
<tr>
<td>Total</td>
<td>73</td>
<td>60</td>
</tr>
</tbody>
</table>

*Figures are number of episodes. Adverse events other than death, doubling of serum creatinine or end-stage renal failure were not recorded after drop or after the primary end point of the trial was met.

DISCUSSION

In this investigator-initiated multicentre randomised long-term maintenance trial, MMF was not significantly superior to AZA in preventing renal flares in patients with proliferative LN, with a renal relapse rate of 19 and 25% in the MMF and AZA group, respectively. Time to severe systemic flare, benign flare, GC withdrawal and renal remission did not differ between the two groups. Drug-related toxicities were similar except for cytopenias, which were more common in the AZA group but were readily controlled by dose adjustments and led to only one patient dropping out.

Several caveats can be raised. First, MAINTAIN is a relatively small size European-based investigator-led clinical trial with mainly Caucasian patients rather than a large international multinational study. Second, it was an open label and not a double-blinded trial. It should be stressed, however, that the primary outcome (that is, renal relapse) was strictly defined on biological grounds and therefore unlikely to have been influenced by knowledge of patient allocation. Third, the trial design can be criticised as we did not request patients to have a significant renal response before starting maintenance treatment with AZA or MMF. This was a deliberate choice as we felt that such a design more closely paralleled clinical practice, including patients even if they did not satisfactorily respond after 5 months of induction treatment. Strictly speaking, MAINTAIN is a comparison of a ‘GC/intravenous CY/AZA protocol’ versus a ‘GC/intravenous CY/MMF protocol’. Importantly, a subset analysis performed on patients who, at any time during follow-up, achieved a significant renal response defined as a ≥50% reduction of 24 h proteinuria, showed similar renal flare rates in the two groups, thereby ruling out a major design bias.

Fourth, serum measures of the active metabolites of AZA (ie, 6-thiaauguanines (6-TGN)) or of MMF (ie, mycophenolic acid (MPA)) were not routinely performed, leaving open the possibility that patients who failed on one or the other drug were actually underdosed or non-adherent to the medication. This hypothesis might not be too far fetched based on the recent finding that patients who have undergone kidney transplant had a lower rejection rate if MMF doses were titrated according to serum MPA titres instead of fixed. Finally, the decision to design a superiority instead of an equivalence trial placed MMF in a more challenging position.

MMF and AZA were also compared for maintenance treatment of LN after induction treatment with pulse intravenous CY by Contreras et al. Patients included in this trial were almost entirely non-Caucasians and had severe LN. Six AZA-treated and three MMF-treated patients had a renal relapse, but this difference was not statistically significant due to the small number of patients in each group. During the review process of this manuscript, we became aware that the maintenance phase of the Aspreva Lupus Management Study (ALMS), also comparing AZA and MMF for maintenance immunosuppression of LN, met its primary end point: MMF was found to be superior to AZA in delaying the time to treatment failure, a composite index comprising death, serious renal damage and relapse of LN. Although more data are eagerly awaited before conclusions can be drawn, several reasons for the discrepancy between the results of MAINTAIN and ALMS can already be hypothesised, such as inclusion of patients from different ethnic groups in ALMS, a different trial design (only patients having achieved a renal response at 6 months were included in the maintenance phase of ALMS) or a different primary outcome measure (a composite end point in ALMS). However, the major reason for the difference may well stem from the numbers of
patients randomised in the 2 trials (105 and 227 in MAINTAIN and ALMS, respectively).

At the bedside, other arguments than primary renal efficacy will be balanced to choose between MMF and AZA. First, MMF has distinct effects in vitro suggesting that the drug might be cardioprotective in vivo, a potential advantage in LN given the high rate of cardiovascular disease associated with SLE. 19 Thus, MMF inhibits the proliferation of fibroblasts and vascular smooth muscle cells 20 and its use was shown to be associated with less cardiac allograft vasculopathy compared to AZA in patients who have undergone heart transplant. 21 Data indicating a cardioprotective effect of MMF in SLE are, however, inconclusive. 22 Second, the mode of action of MMF that is, inhibition of inosine monophosphate dehydrogenase, does not theoretically lead to mutagenic events. Conversely, at least part of the pharmacological effect of AZA involves incorporation of 6-TGNs into DNA, thereby raising concerns regarding mutagenesis. In this respect, an increased frequency of somatic mutations at the hypoxanthine phosphoribosyltransferase locus, related to total dose and treatment duration, was recently found in peripheral T lymphocytes of patients with inflammatory bowel disease treated with AZA, thereby stressing the potential carcinogenicity of the drug. 23

However, two arguments can be advocated in favour of AZA. First, AZA can be safely used during pregnancy whereas MMF is absolutely contraindicated. MMF is likely a human teratogen based on the unusual distribution of malformations observed among reported exposed offspring (microtia or anotia, external auditory canal atresia, orofacial clefts, hypertelorism, coloboma, microganthia and cardiovascular malformations). 24 When long-term immunosuppression is being considered in young women planning pregnancy, this issue needs to be appreciated, as indeed suggested by the trial reported here where patients dropping out for pregnancy or pregnancy planning were more frequently observed in the MMF arm compared to the AZA arm. Second, the cost of MMF is currently 10 times higher than that of AZA. Although the next availability of MMF generics might reduce these costs, it could be part of the decision at the bedside.

In conclusion, there were fewer renal flares seen in the MMF-treated patients but this did not reach statistical significance and we were not able to demonstrate the superiority of MMF over AZA. However, the data from the MAINTAIN Nephritis Trial suggest that at least two drugs are available for long-term use in patients with LN, with a reasonable efficacy and an excellent toxicity profile. The possibility that some patients who fail on one immunosuppressant could be successfully rescued by switching to the other was not addressed in this study, but is in a very reasonable consideration. A welcome step forward would consist of a priori identification of those patients who will respond to one or to the other drug, as well as the search for even more effective maintenance therapies. In this respect, the place of targeted treatment with biologicals clearly needs further investigation.

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REFERENCES


