Ciclosporin compared with prednisolone therapy for patients with pyoderma gangrenosum

Citation for published version:

Digital Object Identifier (DOI):
10.1111/bjd.15561

Link:
Link to publication record in Edinburgh Research Explorer

Document Version:
Publisher's PDF, also known as Version of record

Published in:
British journal of dermatology

Publisher Rights Statement:
This is an open access article under the terms of the Creative Commons Attribution-NonCommercial License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited and is not used for commercial purposes.

General rights
Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy
The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact openaccess@ed.ac.uk providing details, and we will remove access to the work immediately and investigate your claim.
Ciclosporin compared with prednisolone therapy for patients with pyoderma gangrenosum: cost-effectiveness analysis of the STOP GAP trial

J.M. Mason,1 K.S. Thomas,2 A.D. Ormerod,3 F.E. Craig,4 E. Mitchell,5 J. Norrie,6 and H.C. Williams2 on behalf of the U.K. Dermatology Clinical Trials Network's STOP GAP team

1Warwick Medical School, University of Warwick, Coventry, CV4 7AL, U.K.
2Centre of Evidence Based Dermatology, University of Nottingham, NG7 2NR, U.K.
3Division of Applied Medicine, Aberdeen University, Aberdeen, AB24 2ZD, U.K.
4Department of Dermatology, NHS Forth Valley, Stirling, FK8 2AU, U.K.
5Nottingham Clinical Trials Unit, University of Nottingham, Nottingham, NG7 2UH, U.K.
6Edinburgh Clinical Trials Unit, Usher Institute, University of Edinburgh, EH16 4TU, U.K.

Summary

Background Pyoderma gangrenosum (PG) is a painful, ulcerating skin disease with poor evidence for management. Prednisolone and ciclosporin are the most commonly used treatments, although not previously compared within a randomized controlled trial (RCT).

Objectives To compare the cost-effectiveness of ciclosporin and prednisolone-initiated treatment for patients with PG.

Methods Quality of life (QoL, EuroQoL five dimensions three level questionnaire, EQ-5D-3L) and resource data were collected as part of the STOP GAP trial: a multicentre, parallel-group, observer-blind RCT. Within-trial analysis used bivariate regression of costs and quality-adjusted life years (QALYs), with multiple imputation of missing data, informing a probabilistic assessment of incremental treatment cost-effectiveness from a health service perspective.

Results In the base case analysis, when compared with prednisolone, ciclosporin was cost-effective due to a reduction in costs [net cost: £1160; 95% confidence interval (CI) −2991 to 672] and improvement in QoL (net QALYs: 0.055; 95% CI 0.018–0.093). However, this finding appears driven by a minority of patients with large lesions (≥20 cm²) (net cost: £5310; 95% CI −9729 to 891; net QALYs: 0.077; 95% CI 0.004–0.151). The incremental cost-effectiveness of ciclosporin for the majority of patients with smaller lesions was £23 374/QALY, although the estimate is imprecise: the probability of being cost-effective at a willingness-to-pay of £20 000/QALY was 43%.

Conclusions Consistent with the clinical findings of the STOP GAP trial, patients with small lesions should receive treatment guided by the side-effect profiles of the drugs and patient preference – neither strategy is clearly a preferred use of National Health Service resources. However, ciclosporin-initiated treatment may be more cost-effective for patients with large lesions.

What’s already known about this topic?

- Pyoderma gangrenosum is characterized by severe, painful skin ulcers.
- Although prednisolone has been the main systemic treatment, ciclosporin has been used increasingly because of its perceived greater effectiveness and fewer side-effects.
Pyoderma gangrenosum (PG) is a rare, inflammatory skin disease characterized by progressive and painful necrotizing ulcers. Typically, PG presents as a tender erythematous nodule or pustule, quickly progressing to a large, demarcated ulcer with purplish, undermined edges. PG is associated with underlying systemic disease, and! in particular with inflammatory bowel disease, arthritis and haematological malignancies. Additionally it may develop following incidental or iatrogenic trauma. PG is associated with a three fold increased risk of death; its ulcers are characterized by debilitating pain and may require narcotic analgesia.

PG is diagnosed clinically after excluding other diagnoses because there are no adequate diagnostic tests and histological findings are relatively nonspecific. No national or international guidelines address PG management, which currently includes a range of poorly evidenced topical and systemic treatment options including antibiotics, steroids, calcineurin inhibitors and immunosuppressants. Only one randomized controlled trial (RCT) of treatments has previously been reported in patients with PG: a study of 30 patients compared infliximab against placebo and showed benefit for infliximab at 2 weeks. However, infliximab is not a first-line treatment for this condition.

Given the absence of high-quality evidence for the management of PG, the STOP GAP trial was designed to test whether treatment with ciclosporin was superior to prednisolone. In brief, STOP GAP was a multicentre, parallel-group, assessor-blind RCT, recruiting 112 adult patients, with outcomes assessed at baseline, 6 weeks and when the ulcer had healed (if within 6 months). Groups were balanced at baseline. The primary end point of velocity of healing at 6 weeks was similar between groups [adjusted mean difference 0·003 cm² daily, 95% confidence interval (CI) −0·20 to 0·21; P = 0·97; healing within 6 months was similar (ciclosporin 47·5%, prednisolone 47·2%; P = 0·84)]. Adverse reactions were similar (ciclosporin 67·8%, prednisolone 66·0%; P = 0·84), but serious adverse reactions may have been more common in the prednisolone group (ciclosporin 3%, prednisolone 13%; P = 0·082), in particular due to five serious infections that required hospitalization for parenteral antibiotics. Having found no difference for a range of objective and patient-reported outcomes, the trialists concluded that treatment decisions for individual patients should be guided by the different side-effect profiles of the two drugs and patient preference.

Economic analysis is intended to inform decision-makers about the value-for-money of treatment alternatives in a context where healthcare resources are limited and prioritization is informed (at least in part) by the efficient use of resources. An economic analysis was designed integrally within the STOP GAP trial, following a prospective analysis plan, to provide robust evidence of cost-effectiveness to inform health service decision-making.

Patients and methods

A within-trial patient-level cost-effectiveness analysis was undertaken using data from the STOP GAP trial. The analysis was from the National Health Service (NHS) perspective; individual patient data collected within the STOP GAP trial included NHS treatment costs and health status, estimated as quality-adjusted life years (QALYs). Cost-effectiveness analysis captures the effect of treatment as changes in cost and QALYs. Because follow-up was limited to 24 weeks, no discounting of costs and benefits was applied. The analysis followed intention-to-treat principles, in which patients were included in the analysis according to the treatment allocated by randomization and irrespective of subsequent care.

Outcomes

Generic health-related quality of life (QoL) was assessed using the EuroQol (EQ) questionnaire: a patient-completed two-page questionnaire consisting of the EQ five dimensions three level questionnaire (EQ-5D-3L) descriptive system and the EQ visual analogue scale (EQ-VAS). The EQ-5D-3L includes

What does this study add?

- For patients with small lesions (< 20 cm²), neither treatment is clearly more cost-effective than the other.
- However, ciclosporin-initiated treatment may be the more cost-effective option in patients with large (≥ 20 cm²) lesions.
- Decisions about treatment will continue to be informed primarily by patient preference, underlying comorbidities, and drug side-effect profiles (e.g. serious infections with prednisolone, hypertension and renal dysfunction with ciclosporin).

STOP GAP was a pragmatic randomized controlled trial comparing ciclosporin and prednisolone: clinical effectiveness was similar, but only 50% of ulcers had healed by 6 months on either drug and adverse events were common with both drugs.
five questions addressing mobility, self-care, usual activities, pain/discomfort and anxiety/depression), with each dimension assessed at three levels: no problems, some problems and extreme problems. EQ-5D scores were converted to health status scores using the U.K. time-trade-off value set recommended by the EuroQol group, providing a single health-related index including 0 (death) and 1 (perfect health), where negative scores are possible for some health states. Patients who died during the study were subsequently scored 0 at later scheduled follow-up visits. The EQ-VAS reports self-rated health on a vertical VAS where 100 denotes ‘best imaginable health state’ and 0 denotes ‘worst imaginable health state’. Additionally, the Dermatology Life Quality Index (DLQI) was recorded as a disease-specific measure: DLQI asks patients 10 questions about how their skin condition has affected their life over the past week providing an aggregate score of range 0–30. QoL measures were captured at baseline, 8 weeks and up to 24 weeks (unless healing had occurred).

Using the trapezoidal rule, the ‘area under the curve’ (AUC) of health status scores was calculated, providing patient-level QALY estimates for the cost-effectiveness analysis. Similarly, EQ-VAS and DLQI scores were integrated discretely over time. Because AUC estimates were predicted to correlate with baseline scores (and thus potential baseline imbalances), AUC estimates were adjusted for baseline scores.

**Resource use and cost**

Resource assessments occurred at 8 and 24 weeks (or when healed if earlier), during mandatory clinical visits, augmented by telephone calls with patients when clinics were missed. Use by patients of study drugs was recorded in the trial drug log. PG-related health service contacts were recorded during clinic visits using patient diaries as an aide memoire.

Patient costs were initially estimated in U.K. pounds sterling (2012) as the sum of resources used weighted by their reference costs. Study drugs were prescribed at varying doses and durations. Using national Prescribing Cost Analysis (PCA) data, average costs per unit weight of therapeutic were determined and applied to patient drug-use records: ciclosporin £0.0242 mg⁻¹ and prednisolone £0.0237 mg⁻¹. Costs for inpatient stays (in days) and outpatient visits were estimated using Hospital Episodes Statistics (HES) and the National Schedule of Reference Costs (NSRC). National HES data were explored for inpatient episodes with a primary diagnosis of L88 Pyoderma gangrenosum: the five most common admission codes associated with that diagnosis were included, accounting for 83% of admissions. (HES admission codes are similar to U.S. Diagnosis-Related Group codes).

Per diem costs for each code were estimated from the NSRC, and a volume-weighted average cost per admission for PG was estimated as the cost per day. Inpatient stays were costed at £387 per day and outpatient visits at £130 per visit. General practitioner (GP) clinic and home visits, and practice and district nurse visits were costed using unit costs provided by the Personal Social Services Research Unit at the University of Kent; community care contacts: GP (clinic) £43, GP (home) £110, practice nurse £14 and district nurse £39. Patient costs were subsequently updated to 2015 U.K. pounds sterling using the Hospital and Community Health Services index.

**Analysis**

Follow-up of patients with PG within trials is problematic and some incompleteness of data was anticipated. Consequently, a base case analysis was constructed where missing data were imputed using multiple imputation. The base case analysis included the imputed within-trial incremental cost and QALYs gained, adjusted for trial baseline covariates. Supportive sensitivity analyses included only patients with complete data, thus exploring the impact of imputation.

The base case analysis used multiple imputation, conducted according to good practice guidance. Multiple imputation provides unbiased estimates of treatment effect if data are missing at random: this assumption was explored in the data, for example by using logistic regression for missingness of costs and QALYs against baseline variables. A regression model was used to generate multiple imputed datasets (or ‘draws’) for individual treatment groups, where missing values were predicted drawing on predictive covariates: these included age, sex, target lesion size (< 20 cm², ≥ 20 cm²), presence or absence of underlying systemic disease. Outcome measures (at each time point) and costs contributed as both predictors and imputed variables. Each draw provided a complete dataset, which reflected the distributions and correlations between variables. Predictive mean matching was used to enhance the plausibility and robustness of imputed values, as normality could not be assumed. The imputation model used fully conditional (Markov chain Monte Carlo, MCMC) methods (multiple imputation by chained equations), which are appropriate when missing and correlated data occur in more than one variable. Each draw was analysed independently using bivariate regression (see below) and the estimates obtained were pooled to generate mean and variance estimates of costs and QALYs using Rubin’s rule – a method that captures within and between variances for imputed samples. To minimize the information loss of finite imputation sampling, 50 draws were taken, resulting in a loss of efficiency relative to infinite sampling of less than 0.5% in all imputed values. The distribution of imputed and observed values was compared visually and statistically to establish that imputation did not introduce bias into subsequent estimation.

Bivariate regression using seemingly unrelated regression equations was used to model incremental changes in costs and QALYs. This method respects the correlation of costs and outcomes within the data, and allows adjustment for a set of covariates, which can be explored and which improve precision. Baseline QoL scores were included within all models to allow for potential baseline imbalances. Joint distributions of costs and outcomes were generated using the
(nonparametric) bootstrap method, with replicates used to populate a cost-effectiveness plane. Bootstrapping jointly resamples costs and outcomes from the original data with replacement (maintaining the sample correlation structure) to create a new bootstrap sample from which a change in costs and QALYs are estimated. Using bias-corrected nonparametric bootstrapping, 5000 bootstraps were taken per model or draw evaluated. Mean estimates are reported with 95% credible intervals.

The incremental cost-effectiveness ratio (ICER) was estimated as the difference between treatments in mean total costs divided by the difference in mean total QALYs. Value-for-money is determined by comparing the ICER with a threshold value, typically the National Institute for Health and Care Excellence threshold for U.K. studies, of £20K–30K/QALY.31 This represents the willingness-to-pay (WTP) for an additional QALY, and lower values than the threshold could be considered cost-effective for use in the NHS. Base case assumptions are explored using a range of supportive sensitivity analyses. The net monetary benefit (NMB) of changing treatment was reported as a recalculation of the ICER at a range of thresholds of WTP for an additional QALY. The NMB succinctly describes the resource gain (or loss) when investing in a new treatment when resources can be used elsewhere at the same threshold. NMB estimates were used to generate cost-effectiveness acceptability curves (CEACs). The CEAC compares the likelihood that treatments are cost-effective as the WTP threshold varies.30

The expected value of perfect information (EVPI) is the upper limit of the value to a healthcare system of further research to eliminate uncertainty.32 Findings from cost-effectiveness analyses remain uncertain because of the imperfect information they use. If a wrong adoption decision (to make a treatment available) is made this will bring with it costs in terms of health benefit forgone: the NMB framework allows this expected cost of uncertainty to be determined and guide whether further research should be conducted to eliminate uncertainty.

Analyses and modelling were undertaken in Stata 14 SE (StataCorp LLC; College Station, TX, U.S.A.). Reporting follows the Consolidated Health Economic Evaluation Reporting Standards (CHEERS) statement.33

### Results

#### Completeness of data

All 112 patients included within the trial primary analysis of effectiveness were included in the economic analysis (Table 1). Patients with complete EQ-5D assessments for all periods numbered 67 in total (60%). One patient died during the study and was subsequently scored 0 on visits that followed for both cost and EQ-5D score and is included in the analysis. There was a pattern of decreasing completeness as follow-up proceeded. Resource data was complete for 67 patients (60%). When considering both utilities and resource use, complete information was available for 55 patients (49%). Completeness of data was similar when comparing treatment arms. Missing values were imputed to provide a base case analysis including all 112 patients.

#### Complete case estimates

Mean EQ-5D scores, resource use and cost data are reported by treatment in Table 2. Over the 24-week follow-up period there were no significant differences in QALYs when comparing treatments. Differences in resource use comparing groups were not statistically significant at any time point, although there is a suggestion of greater inpatient usage by patients on prednisolone. Time as an inpatient was recorded for only 10 patients: total durations were 14 and 5 days for ciclosporin and 54, 48, 46, 38, 16, 7, 6 and 1 for prednisolone. Patterns of resource use were costed using national reference values (see Patients and methods: Resource use and cost). Although costs for patients receiving ciclosporin were less over 24 weeks, the decrease was not statistically significant: −£1046 (95% CI: −£3534 to £1341) (see Table 3).

#### Cost-effectiveness analysis

The joint distribution of incremental cost and outcome for the base case analysis is shown graphically in Figure 1 (see also Table 3). Patients allocated to ciclosporin (compared with prednisolone) experienced a modest average increase in QoL (0·055 QALYs; 95% CI 0·018–0·093) over 24 weeks. Health costs were lower for patients receiving ciclosporin but the difference was imprecise: −£1160; 95% CI: −£2991 to 672); cost differences were predominantly driven by differences in hospitalization (Table 2). The joint distribution of cost and outcome is summarized within the NMB metric. Using a WTP criterion of less than £20 000 per QALY gained, the NMB associated with ciclosporin-initiated therapy was positive £2263 (95% CI: 216–4311). Thus, the base case analysis

### Table 1 Completeness of data by follow-up visit

<table>
<thead>
<tr>
<th>Health status</th>
<th>Ciclosporin (n = 59)</th>
<th>Prednisolone (n = 53)</th>
<th>Total (n = 112)</th>
</tr>
</thead>
<tbody>
<tr>
<td>EQ-5D baseline</td>
<td>56 (95)</td>
<td>52 (98)</td>
<td>108 (96)</td>
</tr>
<tr>
<td>EQ-5D 8 weeks</td>
<td>46 (78)</td>
<td>41 (77)</td>
<td>87 (78)</td>
</tr>
<tr>
<td>EQ-5D 24 weeks</td>
<td>41 (69)</td>
<td>29 (55)</td>
<td>70 (63)</td>
</tr>
<tr>
<td>Complete cases</td>
<td>39 (66)</td>
<td>28 (53)</td>
<td>67 (60)</td>
</tr>
<tr>
<td>Resource use</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Drug use</td>
<td>59 (100)</td>
<td>53 (100)</td>
<td>112 (100)</td>
</tr>
<tr>
<td>Health service 8 weeks</td>
<td>47 (80)</td>
<td>38 (72)</td>
<td>85 (76)</td>
</tr>
<tr>
<td>Health service 24 weeks</td>
<td>41 (69)</td>
<td>34 (64)</td>
<td>75 (67)</td>
</tr>
<tr>
<td>Complete cases</td>
<td>38 (64)</td>
<td>29 (55)</td>
<td>67 (60)</td>
</tr>
<tr>
<td>Health status and resource use</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Complete cases</td>
<td>32 (54)</td>
<td>23 (43)</td>
<td>55 (49)</td>
</tr>
</tbody>
</table>

EQ-5D, EuroQol five dimensions questionnaire.
suggests NHS resources would be better directed to ciclosporin- than prednisolone-initiated therapy in terms of cost-effectiveness. This finding is echoed in the cost-effectiveness acceptability curve, which expresses the NMB finding as a probability (Fig. 2: all patients). The likelihood that ciclosporin- than prednisolone-initiated therapy in terms of cost-effectiveness analysis of the STOP GAP trial, J.M. Mason et al.

### Sensitivity analyses

Comparing mean costs and QoL estimates using different modelling assumptions supported the base case finding (Table 3). The qualitative similarity of imputed and complete case estimates supports the validity of the imputation process and assumptions.

### Subgroup analyses

There was no interaction between treatment effect and baseline covariates except in the case of index lesion size. The 40 patients (36%) recruited with large lesions (≥ 20 cm²) experienced a different pattern of costs from the patients with smaller lesions (see Fig. 3 and Table 3).

Ciclosporin-initiated treatment markedly lowered costs for patients presenting with large lesions (−£5310, 95% CI $–9729$ to $–891$), but not patients with smaller lesions (£1007, 95% CI −269 to 2283). These differences were driven by the pattern of hospitalization, which predominantly occurred in patients receiving prednisolone and may be linked to the occurrence of serious adverse events.

For patients presenting with large lesions, ciclosporin-initiated treatment appears to be a cost-effective strategy (Figs 2 and 3: index lesion ≥ 20 cm²). However, for patients presenting with smaller lesions, for ciclosporin-initiated treatment, cost-effectiveness (£23 374/QALY) is uncertain with the 95% confidence region including preference for either treatment; consequently neither strategy is clearly a preferred use of NHS resources for patients with smaller lesions (Figs 2 and 3: index lesion < 20 cm²).

### Value of further research

An EVPI analysis was conducted to explore the value of reducing uncertainty about the cost-effectiveness of ciclosporin- or prednisolone-initiated therapy. EVPI analysis at the patient level was conducted treating the two trial strata for index lesion size as independent trials. There is considerable certainty about the findings for the trial as a whole as well as for patients with large lesions: the remaining value of obtaining perfect information is low (Fig. 4). However, there remains

---

Table 2 Health status, resource use and cost (complete cases)

<table>
<thead>
<tr>
<th>Health status</th>
<th>Ciclosporin (C) Mean (SD)</th>
<th>Prednisolone (Pr) Mean (SD)</th>
<th>(C)–(Pr) Mean (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>EQ-SD baseline</td>
<td>0.51 (0.35)</td>
<td>0.44 (0.38)</td>
<td>0.07 (−0.06 to 0.22)</td>
</tr>
<tr>
<td>EQ-SD 8 weeks</td>
<td>0.65 (0.30)</td>
<td>0.53 (0.39)</td>
<td>0.12 (−0.03 to 0.27)</td>
</tr>
<tr>
<td>EQ-SD 24 weeks</td>
<td>0.80 (0.22)</td>
<td>0.66 (0.38)</td>
<td>0.15 (−0.01 to 0.30)</td>
</tr>
<tr>
<td>EQ-SD AUC</td>
<td>0.33 (0.08)</td>
<td>0.29 (0.15)</td>
<td>0.04 (−0.02 to 0.10)</td>
</tr>
</tbody>
</table>

AUC, area under the curve; CI, confidence interval; EQ-SD, EuroQol five dimensions questionnaire; GP, general practitioner; NHS, National Health Service. *Ordinary least squares regression-estimated means and 95% CIs. †Resource use has different missing values in the two periods: overall resource use is not a simple sum of these items. ‡Average (mean) weight of allocated study drug.
### Table 3: Cost-effectiveness, cost/QALY (£, 2015): ciclosporin compared with prednisolone

<table>
<thead>
<tr>
<th>Incremental cost (£, 2015)</th>
<th>Incremental QALY</th>
<th>ICERb (95% CI)</th>
<th>NMBc (£)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Base case</td>
<td>1160 (2991 to 672)</td>
<td>0.055 (0.018 to 0.093)</td>
<td>Dominant</td>
</tr>
<tr>
<td>Sensitivity analyses</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1. Imputed attributable costs and QALYs, baseline EQ-5D adjusted</td>
<td>1046 (3534 to 1341)</td>
<td>0.047 (0.001 to 0.090)</td>
<td>Dominant</td>
</tr>
<tr>
<td>2. Complete case attributable costs and QALYs, covariate adjusted</td>
<td>1007 (&lt; 100)</td>
<td>0.013 (0.0001 to 0.085)</td>
<td>Dominant</td>
</tr>
</tbody>
</table>

CI, confidence interval; EQ-5D, EuroQol five dimensions questionnaire; ICER, incremental cost-effectiveness ratio; NMB, net monetary benefit; QALY, quality-adjusted life year.aICER: dominance indicates average costs were less and average benefit greater for ciclosporin-initiated therapy. bProbability cost-effective or NMB if willing to pay £20 000 per QALY gained. cProbability cost-effective or NMB if willing to pay £30 000/QALY.

### Other end-points

The three QoL measures used in the STOP GAP trial are reported in Table 4. EQ-VAS, like EQ-5D, provides 6-month approximations of quality-adjusted survival for each treatment group. EQ-VAS is scored 1–100: equivalent QALY scores are obtained by dividing by 100, although EQ-VAS is not recommended for QALY estimation within trials, as values are preference-rated rather than societal. DLQI is scored 0–30: the average score over the 6-month follow-up period is reported, using the AUC between the three time points to calculate the average. Being a disease-specific QoL measure, the DLQI is potentially more sensitive to change than a generic measure.

Between-group differences for all three imputed QoL measures are shown in Table 4, including unadjusted, baseline score-adjusted and full covariate-adjusted estimates. For each measure, there is a trend favouring ciclosporin.

### Discussion

The STOP GAP trial featured a pragmatic multicentre design reflecting real-world clinical practice; thus, cost and outcome profiles are likely to reflect routine care in NHS settings. Patient-level data from the STOP GAP trial provide the most robust evidence to date on whether ciclosporin or prednisolone is cost-effective as first-line treatment for patients with PG. The base case analysis (using multiple imputation) found ciclosporin-initiated treatment to be cost-effective compared with prednisolone, primarily due to a modest net cost savings and improvement in QoL. However, this finding was driven by the performance of the subgroup of patients with large lesions. In the majority of patients with smaller lesions (< 20 cm²) the estimated cost-effectiveness was too imprecise to differentiate between treatments. These findings are consistent with the results of the clinical trial, which found no difference between treatments in speed of healing, 6-month healing rates or recurrence, but a (near-significant) difference in the EQ-5D based on complete cases. Further, the trial reported a (near-significant) difference in more serious
Fig 1. Cost-effectiveness plane: ciclosporin compared with prednisolone, base case analysis (cost per QALY, £, 2015). QALY, quality-adjusted life year.

Fig 2. Cost-effectiveness acceptability curve: ciclosporin compared with prednisolone, base case and subgroup analyses.

Fig 3. Cost-effectiveness plane: ciclosporin compared with prednisolone, base case and subgroup analyses (cost per QALY, £, 2015). QALY, quality-adjusted life year.
adverse reactions with prednisolone, particularly for infections, which might increase costs.

There are several caveats to these findings. PG is a rare disease and recruitment is problematic; although the largest trial of its kind, STOP GAP recruited only 112 patients from 39 U.K. hospitals over 3.5 years. Lesion size was a stratification variable within the trial randomization making the strata subgroups nested randomized controlled comparisons within the overall trial. Reflecting the subgroup patient numbers, differentiation of cost-effectiveness by lesion size would be strengthened by further evidence before prioritizing ciclosporin routinely for patients with large lesions.

Another weakness is the incompleteness of the data contributing to the economic analysis, a consequence of trying to maintain data quality over so many sites and time, and when the energy of trialists might focus on the clinical data. Exploring the consequences of imputation, the findings appear robust within a range of sensitivity analyses.

A final issue concerns the profile of costs and EQ-5D scores over time (Table 2). In the case of QoL, differences seem to be present and continuing beyond 24 weeks while costs have not clearly converged (accepting the different time periods involved). Thus, there might be a case to model extrapolated costs and outcomes beyond 24 weeks. In essence, modelling an extrapolated time horizon is appropriate when it (i) permits better characterization of the decision problem; (ii) allows evidence synthesis (e.g. from multiple trials); or, (iii) improves characterization of uncertainty. While the within-trial analysis presented provides findings relevant to health service decision-makers, evidence is lacking on which to model plausible longer-term treatment and prognosis of patients with PG. Although the assumptions involved and quality of available trial data further limit the value of such modelling, any attempt would be likely to further emphasize the value of ciclosporin in preference to prednisolone in large lesions. The trial also captured relapses of symptoms beyond 24 weeks: these were infrequent and balanced between groups; thus, their inclusion would not influence the findings.

For patients presenting with smaller lesions the economic and clinical findings align in the sense that clinical outcomes are similar and the cost-effectiveness analysis is too imprecise to differentiate between these strategies. Uncertainty about the cost-effectiveness of ciclosporin- or prednisolone-initiated therapy for patients with small lesions is unlikely to be

Table 4 EQ-5D, EQ-VAS and DLQI estimates: ciclosporin compared with prednisolone

<table>
<thead>
<tr>
<th>AUC estimates</th>
<th>Mean difference</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>EQ-5D imputed, unadjusted</td>
<td>0.061</td>
<td>0.016 to 0.105</td>
</tr>
<tr>
<td>EQ-5D imputed, baseline adjusted</td>
<td>0.046</td>
<td>0.010 to 0.083</td>
</tr>
<tr>
<td>EQ-5D imputed, covariate adjusted</td>
<td>0.055</td>
<td>0.018 to 0.093</td>
</tr>
<tr>
<td>EQ-VAS imputed, unadjusted</td>
<td>0.043</td>
<td>0.001 to 0.085</td>
</tr>
<tr>
<td>EQ-VAS imputed, baseline adjusted</td>
<td>0.077</td>
<td>0.004 to 0.151</td>
</tr>
<tr>
<td>EQ-VAS imputed, covariate adjusted</td>
<td>2.214</td>
<td>-0.799 to 5.227</td>
</tr>
<tr>
<td>EQ-VAS imputed, covariate adjusted</td>
<td>2.051</td>
<td>-0.501 to 4.603</td>
</tr>
<tr>
<td>EQ-VAS imputed, covariate adjusted</td>
<td>2.556</td>
<td>-0.117 to 5.229</td>
</tr>
<tr>
<td>DLQI imputed, baseline adjusted</td>
<td>-2.646</td>
<td>-4.796 to -0.497</td>
</tr>
<tr>
<td>DLQI imputed, covariate adjusted</td>
<td>-1.214</td>
<td>-2.685 to 0.258</td>
</tr>
<tr>
<td>DLQI imputed, covariate adjusted</td>
<td>-1.202</td>
<td>-2.719 to -0.316</td>
</tr>
<tr>
<td>DLQI imputed, baseline adjusted</td>
<td>-1.005</td>
<td>-2.795 to 0.785</td>
</tr>
<tr>
<td>DLQI imputed, covariate adjusted</td>
<td>-1.566</td>
<td>-4.350 to 1.218</td>
</tr>
</tbody>
</table>

AUC, area under the curve; CI, confidence interval; DLQI, Dermatology Life Quality Index; EQ-5D, EuroQol five dimensions questionnaire; EQ-VAS, EQ visual analogue scale.
resolved, at least within the NHS jurisdiction, given the challenges in conducting a further definitive trial, although EVPI suggests there might be value in doing so. It is likely in the health service setting that uncertainty about cost-effectiveness will be a secondary concern, with the clinical findings of similar effectiveness permitting continued use of either ciclosporin- or prednisolone-initiated therapy as the clinical context dictates. The subgroup analysis indicates ciclosporin may be preferred on cost-effectiveness grounds, particularly in patients with large lesions.

References
