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EXTENDED REPORT

Treatment outcome in early diffuse cutaneous systemic sclerosis: the European Scleroderma Observational Study (ESOS)


Handling editor Tore K Kvien


ABSTRACT

Objectives The rarity of early diffuse cutaneous systemic sclerosis (dcSSc) makes randomised controlled trials very difficult. We aimed to use an observational approach to compare effectiveness of currently used treatment approaches.

Methods This was a prospective, observational cohort study of early dcSSc (within three years of onset of skin thickening). Clinicians selected one of four protocols for each patient: methotrexate, mycophenolate mofetil (MMF), cyclophosphamide or ‘no immunosuppressant’. Patients were assessed three-monthly for up to 24 months. The primary outcome was the change in modified Rodnan skin score (mRSS). Confounding by indication at baseline was accounted for using inverse probability of treatment (IPT) weights. As a secondary outcome, an IPT-weighted Cox model was used to test for differences in survival.

Results Of 326 patients recruited from 50 centres, 65 were prescribed methotrexate, 118 MMF, 87 cyclophosphamide and 56 no immunosuppressant. 276 (84.7%) patients completed 12 and 234 (71.7%) 24 months follow-up (or reached last visit date). There were statistically significant reductions in mRSS at 12 months in all groups: −4.0 (−5.2 to −2.7) units for methotrexate, −4.1 (−5.3 to −2.9) for MMF, −3.3 (−4.9 to −1.7) for cyclophosphamide and −2.2 (−4.0 to −0.3) for no immunosuppressant (p value for between-group differences=0.346). There were no statistically significant differences in survival between protocols before (p=0.389) or after weighting (p=0.440), but survival was poorest in the no immunosuppressant group (84.0%) at 24 months.

Conclusions These findings may support using immunosuppressants for early dcSSc but suggest that overall benefit is modest over 12 months and that better treatments are needed.

Trial registration number NCT02339441.

INTRODUCTION

The diffuse cutaneous subtype of systemic sclerosis (dcSSc) is rare (SSc incidence is around 10–20/ million/year,1 of whom approximately 25% will have diffuse disease) but carries high morbidity and mortality due to early internal organ involvement and rapidly progressive, painful skin thickening. Also, 5-year and 10-year survival rates, although improving, are in the order of 68% and 50%, respectively.2,3

At present, there is no drug known to favourably influence disease course. Randomised controlled trials (RCTs) have historically been confounded by disease rarity (only small numbers of patients are recruited, often over long periods) and strict entry criteria meaning that severe cases are often excluded.4 These strict criteria further restrict sample sizes and limit generalisability. Therefore, although RCTs represent a gold standard for assessing drug efficacy, results may not be applicable to
real-life clinical settings. Small trials run the risk of being underpowered, thus potentially yielding false-negative results. The past three decades have seen a number of promising treatments for early dcSSc failing to meet efficacy end points in RCTs: examples include methotrexate (multinational, 71 patients) and anti-transforming growth factor β1 antibody therapy (multinational, 45 patients).

A further difficulty in recruiting into RCTs of early dcSSc is that many clinicians have reservations about placebo therapy in a potentially life-threatening disease and favour immunosuppression, consistent with the European League Against Rheumatism (EULAR) recommendations, which advocate methotrexate for skin manifestations in early dcSSc, although this agent has been shown to be of only limited efficacy. Immunosuppressants are potentially hazardous, especially in patients prone to internal organ disease and infection.

Against this background, our aim was to compare, using an observational approach, the effectiveness of standard treatment approaches (mainly immunosuppressant treatments but including a ‘no immunosuppressant’ option to reflect that some patients or clinicians may choose this approach) in the early management of patients with dcSSc, capturing entry and outcome data in a systematic way. Modern statistical approaches allow robust interrogations of prospective observational studies, as an adjunct to, or even substitute for, RCTs in rare diseases, although the potential of these novel approaches has not yet been realised.

METHODS

Study design

The European Scleroderma Observational Study (ESOS) was a prospective, observational cohort study (ClinicalTrials.gov identifier: NCT02339441), in which standardised data were collected at study entry and at follow-up visits, and entered electronically by investigators at each centre into an electronic case record form. All data were checked by the project coordinator and any inconsistencies were discussed with the chief investigator and (if appropriate) the local principal investigator. The main inclusion criteria were early dcSSc (skin involvement proximal to elbow, knee, face, neck and within three years of the onset of skin thickening) and age >18 years. Exclusion criteria were previous stem cell transplantation, previous immunosuppressant treatment for >4 months or use of any immunosuppressant drug other than methotrexate, mycophenolate mofetil (MMF) or cyclophosphamide within the month prior to study entry.

Clinicians selected the protocol of their choice for each patient. The recommended treatment protocols, as decided by the Steering Committee to reflect international best clinical practice, were

1. Methotrexate (oral or subcutaneous with a target dose of 20–25 mg weekly).
2. MMF (500 mg twice daily for 2 weeks increasing to 1 g twice daily).
3. Cyclophosphamide. Possible regimens included:
   i. Intravenous. Minimum monthly dose 500 mg/m² with a recommended duration of 6–12 months.
   ii. Oral. 1–2 mg/kg/day with a recommended duration of 12 months. Patients treated with cyclophosphamide were then usually ‘transferred’ to a maintenance immunosuppressive drug (methotrexate, MMF or azathioprine) as per the treating clinician’s choice.
4. No immunosuppressant treatment, to give the option of including patients in whom immunosuppression was not felt indicated or appropriate (or declined by the patient).

Patients were assessed at baseline, with subsequent visits scheduled three-monthly for 24 months (or between 12 and 24 months for those patients recruited after September 2013). To have 80% power to detect a difference between two treatment arms of five modified Rodnan skin score (mRSS) units at 12 months would require 63 patients per protocol. Allowing 20% loss to follow-up, and varying numbers recruited to the different protocols, recruitment target was 316 patients.

Patients

Patients were recruited between July 2010 and September 2014. Demographic characteristics including age, gender, smoking habit, ethnicity, antibody status (anti-topoisomerase-1 (anti-Scl70), anti-RNA III polymerase, anticientromere) and presence of visceral organ involvement were recorded for all patients. The algorithms to determine the presence of different types of organ involvement are summarised in online supplementary table S1.

Outcome measures

The primary outcome measure, assessed at each visit, was the change in mRSS over time. All mRSS assessments were performed by those experienced in skin scoring. The mRSS is assessed clinically at 17 body sites on a 0–3 scale (maximum score 51) and measures the extent of skin thickening. It is the most commonly used primary outcome measure in RCTs of dcSSc, reflecting disease severity and predicting mortality. All other outcomes/recorded variables were mainly part of routine clinical practice and are summarised in online supplementary table S2. Secondary end points included pulmonary function (forced vital capacity (FVC; % predicted) and carbon monoxide diffusing capacity (DLCO: % predicted)), quality of life15–18 (including the Health Assessment Questionnaire Disability Index (HAQ-DI)15 and Cochin Hand Function Scale18), occurrence of side effects and survival.

Statistical analysis

In an observational study, patient characteristics differ between groups and any differences in outcomes might be driven by those characteristics rather than the treatments (confounding by indication). In each of the analyses (for the different outcome measures), all variables associated with the outcome were considered as confounders.

Differences between protocols at baseline

Kruskal-Wallis test was applied for continuous variables and Fisher’s test for categorical variables.

Influence of baseline characteristics on mRSS at baseline and over time

The association between baseline variables and mRSS was assessed by simple linear regressions, entering each characteristic separately as a predictor of mRSS. To examine how each variable affected the progression of mRSS, the regression equation was modified by adding a term for time and its interaction with the baseline predictor value.

Differences in the changes between groups for all outcomes

Inverse probability of treatment (IPT) weights equalise the distributions of confounders between the treatment groups, thus removing confounding by indication.21 Treatment probabilities


Clinical and epidemiological research
were computed using multinomial logistic regressions, with the baseline values of the selected confounders as predictors. Censoring weights rebalance the data such that the distributions of confounders remain unchanged throughout the study. For each observation, the probability of remaining uncensored given the baseline values of the confounders, the initial protocol and a cubic spline for time was calculated using a pooled logistic regression model. Multiplying both weights yielded the IPT and inverse probability of censoring (IPTC) weights. Weights >20 were truncated at that value.

Treatment effects were assessed using IPTC-weighted linear regression models, which include an intercept, a time term, indicator variables for treatment groups and interactions between time and treatments. The model followed an intention-to-treat approach. Differences in the interaction terms reflected differences in the evolution of outcome.

Cochin hand function data were log-transformed (after adding one to each value) to correct for a highly left-skewed distribution. CIs for the difference of logs were back-transformed, adding one to each value) to correct for a highly left-skewed distribution. CIs for the difference of logs were back-transformed, yielding a percentage difference between predicted baseline and 12-month levels.

Because of missing data at baseline for confounders, multiple imputation by chained equations was applied with STATA V13.1. Imputations were performed separately for each different outcome model. Moreover, each analysis was restricted to the subset of patients with available outcome data at baseline.

Survival analysis
Kaplan-Meier curves, adjusted using IPT weights, provide estimates of the cumulative probability of surviving in each of the protocols. An IPTC-weighted Cox regression, including indicator variables for the protocols, was used to test for differences in survival between protocols. Both overall and adverse event-free survival were examined.

RESULTS
In total, 326 patients from 50 centres (19 countries) were recruited into the study (figure 1): 160 from mainland Europe and the Middle East, 134 from the UK, 15 from Australia and 17 from North America (six centres from Australia and North America joined after the initial recruitment wave). Not being a randomised study, the number of patients starting on each protocol differed: 65 (19.9%) methotrexate, 118 (36.2%) MMF, 87 (26.7%) cyclophosphamide and 56 (17.2%) no immunosuppressant treatment. Median (IQR) doses are shown in online supplementary table S3.

Baseline characteristics of patients
The median mRSS (21, IQR 16–27) and its distribution did not differ across all four treatment groups (p=0.306) (table 1). There were significant differences between treatment groups in gender (patients in the cyclophosphamide group less likely to be female, p=0.003) and duration of skin thickening (the ‘no immunosuppressant’ group had the longest, p=0.001). Also, patients in the cyclophosphamide group were more likely to have had previous immunosuppression (p=0.007) or steroid treatment (p=0.001). At baseline, 94 (28.8%) patients were taking oral corticosteroids, with a median dose of 10 mg/day (range 2.5–60 mg/day).

Organ involvement
There were significant differences between groups for presence of pulmonary fibrosis, cardiac, renal and muscle involvement. Patients on cyclophosphamide were more likely to have pulmonary fibrosis (p=0.036 across groups) or cardiac involvement (p=0.009 across groups). Patients in the ‘no immunosuppressant’ group were more likely to have renal involvement (p=0.039), and the methotrexate group had more frequent muscle involvement (p=0.002).

Functional ability
Scores for the HAQ-DI, Functional Assessment of Chronic Illness Therapy (FACT) fatigue and Short-Form 36 (SF36) physical and mental indexes did not differ significantly between groups. However, there were significant differences across groups in the cochin hand function scale (CHFS), which was poorest in the cyclophosphamide group (p=0.025).

Concomitant medications
As anticipated in a study of patients with early dcSSc, there was substantial use of concomitant medications (see online supplementary table S4).

Progression through the study
Figure 1 shows how patients progressed through the study. Overall, 276 patients (84.7%) remained in the study at 12 months of follow-up and 234 (71.7%) completed 24 months (or reached the last study visit date of 30 September 2015).

Changes in protocol
A total of 60 (18.4%), 12 (3.7%) and 1 (0.3%) patients changed protocol one, two or three times during the study. Among patients still in the study, adherence to initial protocol at 24 months for the different cohorts was 76.2% (methotrexate), 79.7% (MMF), 79.2% (cyclophosphamide) and 73.3% (no immunosuppressant) (see online supplementary figure S1). In the no immunosuppressant cohort, 10 out of 56 patients commenced an immunosuppressant (figure 1).

Withdrawals and deaths
In total, 35 patients (10.7%) died and 42 (12.9%) withdrew from the study (including lost to follow-up). Of the 35 deceased patients, 31 cases were primarily attributed to SSc-related causes (26 most likely primarily cardiorespiratory, 2 renal crises, 2 gastrointestinal (one aspiration) and 1 peritonitis (on peritoneal dialysis following renal crisis)), 3 died of cancer (1 nasopharyngeal, 1 rectal, 1 colorectal) and in 1 case the cause was unknown.

Influence of baseline variables on the initial skin score and on skin score trajectory
Table 2 summarises the effect of different characteristics on the initial mRSS and its subsequent trajectory, as analysed with linear regression.

Using the associations described by table 2, the confounders identified for the skin score were age, duration of skin thickening, current or previous steroid use, anti-topoisomerase, anti-RNA polymerase III, pulmonary fibrosis, pulmonary hypertension, cardiac, renal and muscle involvement, as well as HAQ-DI, Cochin hand function and FACT fatigue scores (see online supplementary table S5 for lists of confounders and online supplementary tables S6–S13 for each model’s confounder selection process).

Changes in skin score over time in the different treatment groups
The mean change in mRSS after 12 and 24 months was −2.9 and −6.7 units. Based on a weighted regression model, there were
Figure 1  Progression of patients through the study.

Withdrawals include patients who left the study for the following reasons: unable to travel, changed hospital, enrolled in a randomized clinical trial, no longer willing to participate or lost to follow-up with no given reason. Among the 8 patients who were lost to follow-up in the first year, 5 patients were due for their 24-month visit after the study deadline.

Protocol violations brought about a premature exit from the study. There were 20 protocol violations mid-study: 9 who started rituximab (as their main treatment), 5 who received a stem cell transplantation, 4 who started azathioprine (not as part of a post-cyclophosphamide transfer), 1 who started tocilizumab and 1 who entered a randomized clinical trial.

Protocol changes designate patients who changed immunosuppressants, started taking or stopped receiving one (in non-observance to their initial assigned protocol). Changes that were a part of a post-cyclophosphamide transfer were not considered protocol changes. If a patient deviated from her/his initial protocol more than once, only the initial change (along with its reason) was recorded in the flow-chat.

There were 58 post-cyclophosphamide transfers: 40 patients went on to mycophenolate mofetil, 9 on to azathioprine, 5 on to methotrexate and 4 on to no immunosuppressant.

In the no immunosuppressant cohort, 10 out of 56 patients deviated from their protocol at any point: 2 switched to methotrexate, 5 to mycophenolate mofetil and 3 to cyclophosphamide. 7 of those 10 cases were linked to pulmonary complications, one to associated inflammatory arthritis, one to skin thickening and one because of preference.
### Table 1  Baseline characteristics and differences between protocols

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Protocol 1 Methotrexate n=65</th>
<th>Protocol 2 Mycophenolate n=118</th>
<th>Protocol 3 Cyclophosphamide n=87</th>
<th>Protocol 4 No immunosuppressant n=56</th>
<th>P Value*</th>
<th>Total n=326</th>
<th>Missing at baseline, no. (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, years</td>
<td>52.5 (41.5–61.1)</td>
<td>50 (40.7–61.1)</td>
<td>54.1 (45.7–60.5)</td>
<td>52.3 (42.3–60.3)</td>
<td>0.324</td>
<td>52.3 (43–60.8)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Female, no. (%)</td>
<td>50 (76.9%)</td>
<td>94 (79.7%)</td>
<td>49 (56.3%)</td>
<td>40 (71.4%)</td>
<td>0.003</td>
<td>233 (71.5%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Months since onset of skin thickening</td>
<td>10.3 (5.5–20.8)</td>
<td>12.6 (7.8–21.8)</td>
<td>10.2 (5.9–14.5)</td>
<td>16.5 (8.7–27)</td>
<td>0.001</td>
<td>11.9 (7–21)</td>
<td>18 (5.5%)</td>
</tr>
<tr>
<td>Years since onset of first non-Raynaud’s manifestation</td>
<td>1.3 (0.7–2.1)</td>
<td>1.4 (0.9–2)</td>
<td>1.0 (0.6–1.4)</td>
<td>1.7 (0.9–2.5)</td>
<td>0.001</td>
<td>1.2 (0.8–2)</td>
<td>6 (1.8%)</td>
</tr>
<tr>
<td>Years since onset of Raynaud’s phenomenon</td>
<td>1.3 (0.7–2.5)</td>
<td>1.9 (1.1–3.1)</td>
<td>1.4 (0.9–2.4)</td>
<td>2.2 (1.1–3.7)</td>
<td>0.014</td>
<td>1.7 (1–2.9)</td>
<td>22 (6.7%)</td>
</tr>
<tr>
<td>Previous immunosuppressant use, no. (%)†</td>
<td>3 (4.6%)</td>
<td>5 (4.2%)</td>
<td>15 (17.2%)</td>
<td>3 (5.4%)</td>
<td>0.007</td>
<td>26 (8%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Current or previous steroid use, no. (%)</td>
<td>33 (50.8%)</td>
<td>39 (34.2%)</td>
<td>51 (58.6%)</td>
<td>18 (32.1%)</td>
<td>0.001</td>
<td>141 (43.8%)</td>
<td>4 (1.2%)</td>
</tr>
<tr>
<td>Current smoker (%)</td>
<td>11 (16.9%)</td>
<td>17 (15.3%)</td>
<td>18 (21.7%)</td>
<td>11 (20.8%)</td>
<td>0.646</td>
<td>57 (18.3%)</td>
<td>14 (4.3%)</td>
</tr>
<tr>
<td>History of cancer, no. (%)</td>
<td>5 (7.8%)</td>
<td>4 (2.4%)</td>
<td>4 (4.7%)</td>
<td>7 (12.5%)</td>
<td>0.121</td>
<td>20 (6.2%)</td>
<td>4 (1.2%)</td>
</tr>
<tr>
<td>Caucasian, no. (%)</td>
<td>54 (83.1%)</td>
<td>92 (78.0%)</td>
<td>77 (88.5%)</td>
<td>49 (87.5%)</td>
<td>0.201</td>
<td>272 (83.4%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>mRSS (0–51)</td>
<td>21 (17–24)</td>
<td>21 (16–27)</td>
<td>22 (17–29)</td>
<td>20 (15.5–26)</td>
<td>0.306</td>
<td>21 (16–27)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Haemoglobin (g/L)</td>
<td>127 (118–136)</td>
<td>126 (118–137)</td>
<td>130 (116–140)</td>
<td>130 (118–139)</td>
<td>0.721</td>
<td>128 (118–137)</td>
<td>13 (4%)</td>
</tr>
<tr>
<td>White blood count ($\times 10^3$/L)</td>
<td>7.7 (6.7–9.4)</td>
<td>7.9 (6.4–9.5)</td>
<td>8.8 (7.3–10.6)</td>
<td>8.0 (6.7–9.4)</td>
<td>0.029</td>
<td>8 (6.8–9.9)</td>
<td>14 (4.3%)</td>
</tr>
<tr>
<td>Platelets ($\times 10^9$/L)</td>
<td>300 (254–337)</td>
<td>298 (246–370)</td>
<td>309 (259–359)</td>
<td>281 (250–337)</td>
<td>0.459</td>
<td>298 (253–356)</td>
<td>15 (4.6%)</td>
</tr>
<tr>
<td>ESR (mm/hour)</td>
<td>17 (10–29)</td>
<td>18 (8–30)</td>
<td>21 (9–48)</td>
<td>20 (10–35)</td>
<td>0.341</td>
<td>18 (8–34)</td>
<td>77 (23.6%)</td>
</tr>
<tr>
<td>CRP (mg/L)</td>
<td>4.0 (2.2–7.5)</td>
<td>5.0 (1.1–11)</td>
<td>5.9 (3.2–20.0)</td>
<td>4.8 (3.0–11.7)</td>
<td>0.026</td>
<td>5 (2.1–11.8)</td>
<td>90 (27.6%)</td>
</tr>
<tr>
<td>Anti-topoisomerase (anti-ScI70, no. %)</td>
<td>20 (31.3%)</td>
<td>49 (42.6%)</td>
<td>39 (45.3%)</td>
<td>18 (33.3%)</td>
<td>0.228</td>
<td>126 (39.5%)</td>
<td>7 (2.1%)</td>
</tr>
<tr>
<td>Anti-RNA polymerase III, no. %</td>
<td>9 (18.8%)</td>
<td>23 (23.2%)</td>
<td>9 (13.0%)</td>
<td>9 (19.6%)</td>
<td>0.433</td>
<td>50 (19.1%)</td>
<td>64 (19.6%)</td>
</tr>
<tr>
<td>Anticentromere, no. %</td>
<td>9 (14.1%)</td>
<td>6 (5.4%)</td>
<td>4 (4.7%)</td>
<td>3 (5.7%)</td>
<td>0.147</td>
<td>22 (7%)</td>
<td>12 (3.7%)</td>
</tr>
<tr>
<td>Organ involvement</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pulmonary fibrosis, no. (%)</td>
<td>8 (12.3%)</td>
<td>13 (11%)</td>
<td>21 (24.1%)</td>
<td>5 (8.9%)</td>
<td>0.036</td>
<td>47 (14.4%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>FVC (% predicted)</td>
<td>93.7 (81–106)</td>
<td>90 (75–102)</td>
<td>82.5 (68.5–96.5)</td>
<td>90 (75–100)</td>
<td>0.026</td>
<td>89 (75–102)</td>
<td>19 (5.8%)</td>
</tr>
<tr>
<td>DLCO (% predicted)</td>
<td>73 (62–83)</td>
<td>64.5 (48–77)</td>
<td>57 (41–73)</td>
<td>64 (52–75)</td>
<td>&lt;0.0005</td>
<td>64 (50–78)</td>
<td>35 (10.7%)</td>
</tr>
<tr>
<td>Pulmonary hypertension, no. %</td>
<td>4 (21.8%)</td>
<td>7 (5.9%)</td>
<td>10 (11.5%)</td>
<td>5 (8.9%)</td>
<td>0.488</td>
<td>26 (8%)</td>
<td>1 (0.3%)</td>
</tr>
<tr>
<td>sPAP or RVSP (mm Hg)†</td>
<td>27 (23–32)</td>
<td>27 (20–33)</td>
<td>30 (23–37)</td>
<td>27 (23–35)</td>
<td>0.472</td>
<td>29 (21–34)</td>
<td>124 (38%)</td>
</tr>
<tr>
<td>Cardiac involvement, no. %</td>
<td>5 (7.7%)</td>
<td>8 (6.8%)</td>
<td>19 (21.8%)</td>
<td>7 (13.0%)</td>
<td>0.009</td>
<td>39 (12%)</td>
<td>2 (0.6%)</td>
</tr>
<tr>
<td>Renal involvement, no. %</td>
<td>1 (1.5%)</td>
<td>13 (11%)</td>
<td>10 (11.5%)</td>
<td>8 (14.3%)</td>
<td>0.039</td>
<td>32 (9.8%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>eGFR (mL/min)</td>
<td>90 (63–90)</td>
<td>84.5 (60–90)</td>
<td>90 (60–90)</td>
<td>80 (60–90)</td>
<td>0.339</td>
<td>85 (60–90)</td>
<td>92 (28.2%)</td>
</tr>
<tr>
<td>Renal crisis</td>
<td>0 (0%)</td>
<td>8 (6.8%)</td>
<td>4 (4.6%)</td>
<td>4 (7.1%)</td>
<td>0.110</td>
<td>16 (4.9%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Plasma creatinine (\mu mol/L)</td>
<td>63.5 (55–72)</td>
<td>69 (55–83)</td>
<td>64.5 (53–86)</td>
<td>64 (56–77)</td>
<td>0.422</td>
<td>65 (55–79.5)</td>
<td>70 (21.5%)</td>
</tr>
</tbody>
</table>

Continued...
Comparison of survival between treatment protocols

Survival was lowest in the no immunosuppressant group at both 12 and 24 months but differences between protocols were not statistically significant either before (p=0.389) or after weighting (p=0.440). In the adjusted model, at 24 months, those in the no immunosuppressant group had a predicted survival rate of 84.0% compared with 94.1% for methotrexate, 88.8% for MMF and 90.1% for cyclophosphamide (figure 3). Patients with lung involvement (pulmonary fibrosis and/or hypertension) at baseline had significantly poorer survival than those without: at 24 months, their predicted survival rate was 74.6% versus 91.7% (p<0.0005) and similarly for cardiac involvement, 71.6% versus 90.7% (p<0.0005).

Adverse effects

Of the 75, 182 and 101 patients who were ever on methotrexate, MMF or cyclophosphamide, respectively, 29 (38.7%), 40 (22.0%) and 23 (22.8%) were reported to have had side effects, necessitating drug discontinuation in 9 (12.0%), 14 (7.7%) and 5 (4.5%) patients, respectively. A survival analysis on protocol exits due to adverse effects showed no differences in the tolerability of the three treatments (p=0.212) (see online supplementary figure S3).

DISCUSSION

Our main findings were, first, that there were no significant differences in outcome between the four treatment protocols (methotrexate, MMF, cyclophosphamide, no immunosuppression), although there may be a signal in favour of immunosuppression for early dcSSc. Although skin score improved in all four treatment protocols, this was least in the no immunosuppressant group. Second, ESOS statistically significant reductions in mRSS in all four treatment groups at 12 months (−4.0 (−5.2 to −2.7) units for methotrexate, −4.1 (−5.3 to −2.9) for MMF, −3.3 (−4.9 to −1.7) for cyclophosphamide and −2.2 (−4.0 to −0.3) for the no immunosuppressant group), but the differences between treatments were not significant (p=0.346) (table 3 and figure 2).

Changes in secondary outcomes over time in the different treatment groups

Lung function

After adjusting for potential confounders, the change rates of FVC and DLCO were not significantly different in the four treatment groups (p=0.460 and 0.505) (table 3). However, in a subset of patients with pulmonary fibrosis or suspected pulmonary fibrosis (cases confirmed on high-resolution CT (HRCT) irrespective of FVC or DLCO, or with one of the following if HRCT not performed: FVC or DLCO under 55% predicted or definite bibasal shadowing on X-ray), there was a significant difference in the change rate of FVC over time (p=0.035). Patients initially prescribed cyclophosphamide demonstrated 7.4% absolute increase in FVC (% predicted) compared with 2.0% decrease for methotrexate, 3.2% increase for MMF and 4.0% increase for the ‘no immunosuppressant’ group (table 3).

Functional ability and hand function

Changes over time for the HAQ-DI and CHFS did not differ between protocols (p=0.130 and 0.073), regardless of adjusting (table 3).

Development of internal organ involvement

This is described in online supplementary figure S2.
confirms the relative effectiveness of cyclophosphamide in patients with pulmonary fibrosis.23 26
An important point when interpreting our findings (and therefore a note of caution) is that the ‘no immunosuppressant’
group was not a control group. Patients in this group had a longer disease duration than the other three groups and were
more likely to have renal involvement.
Our findings lend support to two recently published studies (the Autologous Stem Cell Transplantation International
Scleroderma trial (ASTIS) trial of autologous stem cell transplantation27 and the Scleroderma Lung Study (SLS) II (comparing
MMF and cyclophosphamide),26 which suggest benefit, including in mRSS, from immunosuppression (as did SLS I28). In
ASTIS, those patients randomised to cyclophosphamide had an 8.8 unit fall in mRSS (from 25.8) at 24 months (compared
with 3.3 in ESOS 12 months), but the cyclophosphamide protocol was more intense, and the patients had more severe
disease (patients with the highest mRSS at baseline tend to improve more quickly14 as also demonstrated by our own
 findings (table 2)). mRSS fell by 19.9 units in those patients
randomised to stem cell transplantation27 (and therefore intensive
immunosuppression). In SLS I,25 patients with dcSSc randomised to cyclophosphamide experienced a 5.3 unit fall in
mRSS at 12 months (compared with 3.3 in ESOS), whereas mRSS fell by 1.7 on placebo (compared with 2.2 units in the
ESOS ‘no immunosuppressant’ group). In SLS II,26 mRSS at 24 months fell 4.9 units on MMF (compared with 4.1 units in
ESOS at 12 months) and by 5.4 after 12 months treatment with cyclophosphamide, although these values are not directly com-
parable because they relate to patients with limited cutaneous and dcSSc combined.
The methodological strength of ESOS, which built upon experience gained in a previous, smaller observational study,28
was its design: its standardised protocols emulated the conditions of a clinical trial, and although not randomised, patients
were enrolled into four homogenous treatment arms with well-defined interventions and a systematic record of protocol changes and exits. Entry criteria were deliberately inclusive: RCTs often exclude patients with internal organ involvement and for whom immunosuppression is most likely to be

Table 3  Predicted yearly changes in outcomes and survival rates according to initial protocol, with and without adjusting (95% CI)

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Model specification</th>
<th>p Value</th>
<th>Methotrexate</th>
<th>Mycophenolate mofetil</th>
<th>Cyclophosphamide</th>
<th>No immunosuppressant</th>
</tr>
</thead>
<tbody>
<tr>
<td>mRSS (0–51)</td>
<td>No adjusting, n=326</td>
<td>0.252</td>
<td>−4.4 (−5.7 to −3.2)</td>
<td>−3.8 (−4.9 to −2.8)</td>
<td>−3.5 (−5.0 to −2.0)</td>
<td>−2.4 (−3.9 to −1)</td>
</tr>
<tr>
<td></td>
<td>Adjusting for confounding (weighted model), n=326</td>
<td>0.346</td>
<td>−4.0 (−5.2 to −2.7)</td>
<td>−4.1 (−5.3 to −2.9)</td>
<td>−3.3 (−4.9 to −1.7)</td>
<td>−2.2 (−4.0 to −0.3)</td>
</tr>
<tr>
<td>FVC (% predicted)</td>
<td>No adjusting, n=307</td>
<td>0.045</td>
<td>−1.7 (−4.4 to 1.0)</td>
<td>2.1 (−0.1 to 4.3)</td>
<td>4.0 (0.5 to 7.5)</td>
<td>2.7 (−0.8 to 6.2)</td>
</tr>
<tr>
<td></td>
<td>Adjusting for confounding (weighted model), n=307</td>
<td>0.460</td>
<td>−0.5 (−3.7 to 2.6)</td>
<td>2.0 (−0.7 to 4.6)</td>
<td>3.3 (−0.6 to 7.2)</td>
<td>2.0 (−1.6 to 5.6)</td>
</tr>
<tr>
<td></td>
<td>Subset with PF on HRCT, weighted, n=1291</td>
<td>0.035</td>
<td>−2.0 (−5.9 to 2.0)</td>
<td>3.2 (−0.6 to 7.0)</td>
<td>7.4 (2.2 to 12.7)</td>
<td>4.0 (−1 to 9.0)</td>
</tr>
<tr>
<td>DLCO (% predicted)</td>
<td>No adjusting, n=291</td>
<td>0.703</td>
<td>−1.2 (−3.5 to 1.0)</td>
<td>0.3 (−1.7 to 2.3)</td>
<td>0.6 (−2.0 to 3.3)</td>
<td>−0.3 (−4.2 to 3.7)</td>
</tr>
<tr>
<td></td>
<td>Adjusting for confounding (weighted model), n=291</td>
<td>0.505</td>
<td>−1.6 (−3.8 to 0.6)</td>
<td>0.8 (−1.5 to 3.1)</td>
<td>0.1 (−3.1 to 3.2)</td>
<td>−0.1 (−3.7 to 3.4)</td>
</tr>
<tr>
<td></td>
<td>Subset with PF on HRCT, weighted, n=116</td>
<td>0.809</td>
<td>−0.8 (−5.4 to 3.8)</td>
<td>1.9 (−1.8 to 5.6)</td>
<td>1.8 (−1.9 to 5.4)</td>
<td>1.6 (−2.6 to 5.7)</td>
</tr>
<tr>
<td>HAQ-DI (0–3)</td>
<td>No adjusting, n=307</td>
<td>0.070</td>
<td>−0.1 (−0.2 to 0.1)</td>
<td>0 (−0.1 to 0.1)</td>
<td>−0.2 (−0.3 to −0.1)</td>
<td>0.1 (−0.1 to 0.2)</td>
</tr>
<tr>
<td></td>
<td>Adjusting for confounding (weighted model), n=307</td>
<td>0.130</td>
<td>−0.1 (−0.3 to 0)</td>
<td>0 (−0.1 to 0.1)</td>
<td>−0.1 (−0.3 to 0)</td>
<td>0.1 (−0.1 to 0.2)</td>
</tr>
<tr>
<td></td>
<td>Subset with PF on HRCT, weighted, n=116†</td>
<td>0.072</td>
<td>−1.1 (−3.5 to 2.2)</td>
<td>−0.3 (−1.6 to 1.2)</td>
<td>−3.1 (−5.1 to −0.5)</td>
<td>1.1 (−0.3 to 3.0)</td>
</tr>
<tr>
<td>Cochin hand function (0–90)†</td>
<td>No adjusting, n=230</td>
<td>0.073</td>
<td>−1.4 (−3.5 to 1.4)</td>
<td>−0.6 (−2.0 to 1.1)</td>
<td>−2.4 (−4.7 to 0.6)</td>
<td>1.7 (−1.1 to 4.0)</td>
</tr>
<tr>
<td></td>
<td>Adjusting for confounding (weighted model), n=230</td>
<td>0.440</td>
<td>(12 months) 93.5%</td>
<td>(12 months) 96.5%</td>
<td>(12 months) 88.1%</td>
<td>(12 months) 88.9%</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>(24 months) 93.5%</td>
<td>(24 months) 89.3%</td>
<td>(24 months) 85.4%</td>
<td>(24 months) 85.1%</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>(12 months) 94.1%</td>
<td>(12 months) 91.7%</td>
<td>(12 months) 88.6%</td>
<td>(24 months) 80.1%</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>(24 months) 94.1%</td>
<td>(24 months) 90.1%</td>
<td></td>
<td>(24 months) 84.0%</td>
</tr>
</tbody>
</table>

Significance p: Fisher’s test for equality of change rates between protocols, for each outcome variable.

*Results are reported in terms of changes after 12 months. However, all study data (from baseline to the 24-month end point) were used in estimation. To obtain 24-month changes, multiply results above by 2.

†For the subanalysis involving the subset of patients with pulmonary fibrosis at baseline, patients with definite bibasal pulmonary fibrosis confirmed on HRCT were included, irrespective of FVC value. If no HRCT scan was performed at baseline, an FVC<55%, DLCO<55% predicted or definite bibasal shadowing on X-ray was also a basis for inclusion.

‡Changes expressed in units for the Cochin regression are an approximation derived from the 95% CI of percentage changes between baseline and 12 months (on a scale shifted by one unit), applied to the predicted baseline values for each group in the original scale.

DLCO, carbon monoxide diffusing capacity; FVC, forced vital capacity; HAQ-DI, Health Assessment Questionnaire Disability Index; HRCT, high-resolution CT; mRSS, modified Rodnan skin score (17 sites); PF, pulmonary fibrosis.
Figure 2  Modified Rodnan skin score (mRSS) during baseline and follow-up visits, by initial protocol. For each group of patients, according to their initial protocol, the distribution of the skin score is illustrated on the left-hand side by box and whisker plots (indicating the median and IQR) at baseline, 12 and 24 months. On the right-hand side, the distribution of individual 1-year changes in the skin score is described by histograms and a kernel density estimate. In addition, a vertical green line indicates the value of the average 1-year change in the skin score, irrespective of treatment choice. The bottom panel in the figure describes the estimated changes in mRSS (with 95% CI) according to initial protocol, based on the results from the adjusted model (described in table 3).
beneficial. By recruiting 326 patients from 50 centres, ESOS represents a large cohort of patients with very early dcSSc (median duration of skin thickening 11.9 months): its data will serve as a benchmark when designing and interpreting future clinical trials. This is especially relevant with a number of novel treatment approaches currently being explored including biological agents. For example, in a recent RCT of tocilizumab, mRSS fell over 24 weeks by 3.9 units from 26 in the 43 tocilizumab-treated patients and by 1.2 units from 26 in the 44 placebo-treated patients, this latter fall comparable to the ESOS ‘no immunosuppressant’ response. In comparing between these studies, the higher baseline mRSS in the tocilizumab study should be borne in mind.

The main weakness of observational studies is that each patient’s outcome on her/his treatment arm cannot be completely disentangled from her/his initial characteristics. For instance, ESOS has verified that patients with lung and cardiac involvement tend to be prescribed cyclophosphamide. However, adjusting using IPT weights minimises the problem of confounding by indication.

In conclusion, observational studies offer a rich population-wide perspective assessing treatment effects in a real-world setting. ESOS achieved its aim of following a large international cohort of patients with early dcSSc over 2 years, each of whom was treated according to one of four protocols. The message for clinicians is that there is a weak signal to support using immunosuppressants for early dcSSc (and in particular cyclophosphamide for patients with pulmonary fibrosis). However, it is clear that there remains a pressing need for the development of more effective and targeted treatments.

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Figure 3 Kaplan-Meier estimated survival curves by treatment group.
Clinical and epidemiological research


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Contributors

ALH, ML, LM, AS, EB, LC, JHWD, OD, KE, WJG, RO, MV and CPD were principal investigators at the different sites and recruited patients. GT and CPD were principal investigators at the different sites and reviewed the report, provided comments and approved the final report.

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Competing interests

ALH has done consultancy work for Actelion, served on a Data Safety Monitoring Board for Apricus, received research funding and speaker’s fees from Actelion, and speaker’s fees from GSK. JHWD has consultancy relationships and/or has received research funding from Actelion, BMS, Celgene, Bayer Pharma, Boehringer Ingelheim, IBD Therapeutics, Sanofi-Aventis, Novartis, UCB, GSK, Arias Biopharma, Active Biotech, Galapagos, Inventiva, Medac, Pfizer, Anamara and Ruli and is stock owner of 4D Science GmbH. OD has received consultancy fees from 4D Science, Actelion, Active Biotech, Bayer, Biogenidec, BMS, Boehringer Ingelheim, EpiPharm, Ergose, espefional Foundation, Genentech/Roche, GSK, Inventiva, Lilly, Medac, Medimmune, Pharmaceuticals, Pfizer, Serodapharm, and Sinoxa and received research grants from Actelion, Bayer, Boehringer Ingelheim, Ergose, Pfizer and Sanofi, and has a patent mir-29 for the treatment of systemic sclerosis licenced. WG has received teaching fees from Pfizer. FH has received research funding from Actelion. MEA has undertaken advisory board work and received honoraria from Actelion, and received speaker’s fees from Bristol-Myers Squibb. LC has done advisory board work for Gilead and served Data Safety Monitoring Boards for Cytori and Reata. HG has done consultancy work and received honoraria from Actelion. UM-L is funded in part by EUSTAR/EULAR. JMD has received honoraria from Eli Lilly, Pfizer, Roche, MSD and BMS. AR receives funding from AstaZeneca. CPD has done consultancy for GSK, Actelion, Bayer, Invernessa and Merck-Serono, received research grant funding from GSK, Actelion, CSL Behring and Inventiva, received speaker’s fees from Bayer and given trial advice to Merck-Serono.

Patient consent Obtained.

Ethics approval

The Ethics Committee of each centre approved the study.

Provenance and peer review

Not commissioned; externally peer reviewed.

Data sharing statement

At present, unpublished data from the study are not available for sharing. This position may change in 6–12 months time.

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