ORIGINAL ARTICLE

Therapeutic and diagnostic outcomes of a standardised, comprehensive care pathway for patients with systemic sclerosis

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ABSTRACT

Objectives: To determine the outcomes, including number of medical interventions and initiation of immunosuppressive treatment of a standardised, comprehensive, diagnostic care pathway for patients with systemic sclerosis (SSc). Patient characteristics associated with need for medical interventions and with need for immunosuppressive treatment were determined.

Methods: Data were routinely gathered in connection with a 2-day care pathway combining multidisciplinary care and complete diagnostic work-up of organ involvement in SSc. The number of patients in whom the pathway resulted in medical interventions, and/or initiation of immunosuppressives was recorded. Patient characteristics and diagnostic tests results were compared between patients with and without medical interventions, and patients with and without initiation of immunosuppressives by means of multivariable logistic regression analyses.

Results: During a period of 44 months, 226 patients with SSc were referred to the care pathway. They included 186 (82%) women with mean age of 54 (SD 14.5) years, and median disease duration of 4 years (range 1–11); 73 (32%) of them had diffuse cutaneous SSc. Medical interventions were initiated in 191 (85%) patients, including initiation of immunosuppressive treatment in 49 (22%). Presence of telangiectasias and higher erythrocyte sedimentation rate were associated with any medical intervention. Of commonly available variables, lower age, higher skin score and absence of anticentromere antibody were associated with initiation of immunosuppressives.

Conclusions: A standardised comprehensive 2-day care pathway for patients with SSc resulted in additional diagnostic or therapeutic interventions in 85% of the patients, regardless of SSc subtype and disease duration. In 22% of the patients, immunosuppressive treatment was initiated.

INTRODUCTION

Systemic sclerosis (SSc) is a rare, multisystem connective tissue disorder which is highly heterogeneous in presentation and severity, ranging from limited forms to diffuse disease involving vital visceral organs.1,2 The frequency and diverse nature of the organ-based complications make systematic assessment and long-term follow-up essential for good management of SSc.

Recent studies have demonstrated an improvement in the overall survival among patients with SSc over the last few decades.3,4 Apart from improved survival of the population in general,1 different other factors, including greater awareness for organ involvement and better management of complications are thought to contribute to this improvement.1 As mortality in SSc from 5-year follow-up onwards is mainly determined by presence of organ complications,4 regular follow-up of patients with SSc, including screening for organ involvement seems justified. Indeed, current guidelines advocate a screening, including assessment of organ involvement in patients with SSc, to be
performed at least annually. As risk for different organ-based complications varies between subsets of SSc, the most extensive annual screening is advocated for patients at high risk for organ involvement. For these patients, and for patients newly presenting with SSc, annual screening, including evaluation of pulmonary, renal, cardiac, gastrointestinal and vascular complications is generally advocated.

In addition, a range of SSc-specific manifestations, including, for example, Raynaud’s phenomenon (RP) atrophy, joint contractures and digital ulcers, result in specific limitations in everyday life, for which additional counselling and non-pharmacological interventions can be relevant. Although evidence for efficacy of non-pharmacological interventions is limited, potential importance of psychosocial and rehabilitation interventions for a chronic disease are clear.

To improve healthcare for patients with SSc treated in the Leiden University Medical Center (LUMC), we aimed to develop a care pathway combining comprehensive medical assessment of organ complications with multidisciplinary care specifically targeting SSc-related manifestations.

The execution of a care pathway combining a wide range of diagnostic procedures with multidisciplinary team care in a patient-friendly manner is an organisational challenge. Patients were found to prefer a yearly standardised, multidisciplinary assessment in a day-care setting rather than multiple visits to various outpatient clinics. In line with this preference, a yearly, 2-day standardised diagnostic and multidisciplinary care pathway was developed. It is meant to serve both patients from the university hospital as well as those referred by rheumatologists in non-academic hospitals in the region. The execution of this complex diagnostic and multidisciplinary care pathway in one centre is in line with international management guidelines, which advocate referral of patients with SSc to a specialised centre.

Regarding the therapeutic management resulting from the diagnostic care pathway, the 2009 EULAR guidelines on treatment of SSc, listing evidence and consensus-based pharmacological treatments for various manifestations of SSc, constitute a firm base. However, no clear definitions on required severity of manifestations of SSc with regard to start of treatments are available, which is due to the heterogeneity and complexity of the disease. In the absence of such definitions, all management decisions in the context of the above mentioned care pathway were taken at a multidisciplinary team conference, based on the individual patient characteristics, and all available scientific and clinical knowledge among the team members.

The aim of the current report is to determine the outcomes of this standardised, comprehensive, care pathway in terms of medical and paramedical interventions, and initiation of immunosuppressive treatment.

**MATERIALS AND METHODS**

**Study design**

Data for the present study were prospectively and routinely gathered as part of the standardised, annual 2-day care pathway. The present analysis only concerned the short-term outcomes of patients taking part in the care pathway for the first time between April 2009 and December 2012. Ethical approval was obtained from the Institutional Review Board of the LUMC. All participants gave written informed consent.

**Patients**

Admission criteria for the care pathway include a diagnosis of SSc according to the referring rheumatologist, or a strong suspicion for SSc, and a request for a complete diagnostic work-up to confirm the diagnosis. Patients can be referred by rheumatologists from the outpatient clinic of the LUMC or from any other hospital in the Netherlands.

**The 2-day care pathway**

The care pathway was started in 2009 (and is still operational now, in 2016). It comprises a visit to the rheumatologist, pulmonologist and cardiologist. In addition to the extensive medical screening, patients are routinely seen by a physical therapist, and a specialised nurse. Additionally, consultations with a social worker and/or occupational therapist are scheduled by indication. The specialised nurse acts as the personal manager and coordinator for the patients during the 2-day programme, explores specific information and healthcare needs, and provides individual counselling and instructions.

For every patient, the healthcare pathway is performed on 2 consecutive days between 8:00 and approximately 16:00. Patients not living in the area of Leiden are offered the opportunity to stay overnight in a hotel room inside the hospital. At initiation of the care pathway, the maximum capacity was 2 patients per week; based on the number of referrals, the capacity was increased to 4 patients weekly in 2012. Two weeks after completion of the care pathway, findings for each particular patient are thoroughly evaluated during a multidisciplinary team conference, attended by at least two rheumatologists, one pulmonologist and one specialised nurse, who is also informed about findings of the occupational therapist, the physical therapist, and the social worker.

**Assessments**

All assessments are performed as part of routine care. They include questionnaires to be completed by patients, as well as interviews and physical performance tests, laboratory tests, nailfold videocapillaroscopy (NVC) and imaging.
Sociodemographic characteristics
For all patients, the following social and demographic characteristics are recorded: age, origin, smoking habit, use of alcohol and caffeine and educational level ((1) primary education (0–8 years; low education level), (2) secondary education (9–16 years; medium education level), or (3) higher vocational education/university (postsecondary; high education level)).

SSc classification and severity
Patients included for the current analysis all fulfil diagnostic criteria for SSc as defined by the American College of Rheumatology or LeRoy and Medsger criteria for (early) SSc. Duration of RP, duration of non-RP (time since first symptom other than RP), disease duration (time since diagnosis SSc was confirmed by a physician), physical complaints and modified Rodnan Skin Score (mRSS), are determined by an experienced rheumatologist.

NVC is routinely performed in all patients starting from 2011 with qualitative assessment of capillary patterns: normal/aspecific pattern, SSc pattern (early/active/late) or borderline changes.

Physical functioning
Patients are asked to fill in the Scleroderma Health Assessment Questionnaire; a 20-item questionnaire comprising eight domains of activities of daily living, with the final score ranging from 0 (no disability) to 5 (severe disability).

In addition, the 6 min walk distance (6MWD) is determined. The 6MWD evaluates the global and integrated responses of all the systems involved during exercise, including pulmonary and cardiovascular systems, systemic and peripheral circulation, blood, neuromuscular units and muscle metabolism. It reflects daily exercise, and has good construct validity, as demonstrated in patients with SSc with pulmonary arterial hypertension (PAH).

Quality of life
Quality of life is measured with the SF-36, which contains eight subscales. The scoring range of the SF-36 subscales is (0–100), with higher scores indicating better quality of life. The subscales are converted into two summary scales: the physical component summary, and the mental component summary scale, standardised to a score with a mean of 50 and a SD of 10 in the general Dutch population frequency table and factor score coefficients are used.

Laboratory investigations
- Measurement of haemoglobin, erythrocyte sedimentation rate (ESR), C reactive protein (CRP), creatine phosphokinase (CPK), NT-pro brain natriuretic peptide, creatinine and estimated glomerular filtration rate.
- Measurement of autoantibodies, including antinuclear antibody (indirect immunofluorescence on HEP-2000 cells (Biomedical Diagnostics, Antwerpen, Belgium)), antitopoisomerase I (anti-Scl-70) and anticientromere (ACA), both measured with Enzyme-Linked ImmunoAssay technique (Immunocap 250, ThermoFisher Scientific, Nieuwegein, The Netherlands). Anti-RNA polymerase III was measured externally (ELISA; Sanquin, Amsterdam, The Netherlands); urine screening for erythrocyturia and proteinuria; 24 h urine collection.

Cardiopulmonary investigations
- High-resolution CT (HRCT) of the thorax, pulmonary function tests (PFTs; including analyses of vital capacity (VC), and diffusing capacity for carbon monoxide (DLCO)), cardiopulmonary exercise test (CPET; including analyses of maximum heart rate, maximum wattage, maximum oxygen consumption (VO2) and maximum ventilation).
- Echocardiography, ECG and 24 h rhythm (Holter) registration.

Diagnosis of interstitial lung disease (ILD) is determined based on the presence of a non-specific interstitial pneumonia (NSIP) pattern, or usual interstitial pneumonia (UIP) pattern on HRCT thorax as reported by the radiologist.

The systolic pulmonary artery pressure (SPAP) is estimated using echocardiography by an experienced cardiologist, and elevated pulmonary pressure is defined using a cut-off value of 35 mm Hg. Left ventricular end-diastolic and end-systolic volumes are measured using the biplane modified Simpson’s rule. Left ventricular ejection fraction is calculated as left ventricular end-diastolic—IV end-systolic volume/left ventricular end-diastolic volume. Furthermore, the presence of pericardial effusion is noted.

Presence of arrhythmias is defined as presence of multiform ventricular extrasystoles >100/day, couplets or runs of ventricular tachycardia or supraventricular tachycardia of at least 30 s on 24 h Holter monitoring. Conduction abnormalities are defined as complete left bundle branch block or right bundle branch block, atrioventricular block (first second or third degree), or pacemaker rhythm for sinus node dysfunction.

Diagnostic and/or therapeutic management strategies resulting from the care pathway
The multidisciplinary team conference results in specific advices on (1) diagnostic follow-up, (2) referral to medical specialists, (3) referral to other healthcare professionals or (4) initiation of treatment, or immunosuppressive treatment, and change of supportive medication, including treatment for peripheral vascular complications, for each individual patient. Advice on diagnostic follow-up comprises all patients in whom...
additional diagnostic procedures are advocated during the next 12 months, including right heart catheterisation for possible PAH, gastroscopy, HRCT thorax or PFT for follow-up on pulmonary involvement. Patients with suspicion for possible PAH based on echocardiography, are additionally discussed in the multidisciplinary PAH meeting attended by the rheumatologist, cardiologist, pulmonologist and internist in order to discuss the indication for a right heart catheterisation (RHC).

Referral to medical specialists includes all patients in whom follow-up and treatment by medical specialists additional to follow-up by a rheumatologist is advocated. The decision to initiate immunosuppressive medication is based on consensus between experts during the multidisciplinary team conference in absence of a gold standard precisely guiding treatment decisions in SSC. Initiation of new immunosuppressive treatment is mainly considered in case of extensive and/or progressive skin or organ involvement, for example, in case of relevant decline in VC and/or DLCO (without using an absolute threshold) in combination with the presence of NSIP and/or UIP on HRCT. Autologous haematopoietic stem cell transplantation (HSCT) is applied according to inclusion criteria and treatment regimen, as described in the ASTIS trial. Azathioprine (AZA) is prescribed in case of active arthritis or pulmonary involvement, with contraindication for cyclophosphamide or mycophenolate mofetil; hydroxychloroquine (HCQ) is prescribed in case of SSC overlap syndrome with rheumatoid arthritis (RA). Rituximab is given either in patients with SSC overlap syndrome with RA, or as part of a randomised placebo-controlled clinical trial (RITIS), registered at http://www.clinicaltrialsregister.eu/ EudraCT Number: 2008-007180-16. Treatment for peripheral vascular complications included: bosentan, sildenafil, intravenous iloprost, calcium channel blockers and antibiotics (the latter in case of active digital ulcers with superimposed bacterial infection).

On the basis of the individual patient’s need, the specialised nurse discussed specific non-pharmacological treatment advices regarding smoking habits, Raynaud’s phenomenon (including prescription of therapeutic gloves), care for digital ulcers, and dietary advices.

In addition, patients with specific problems regarding activities or work participation, and/or psychosocial problems, which could not be solved by a single consultation or advice during the care pathway, were referred to the appropriate health professionals, according to the nature and severity of their problems, as well as their personal preferences.

**Patient satisfaction**

As part of quality management of the day care department, patients who participated in the care pathway were requested to complete a questionnaire concerning different aspects of the care pathway and the healthcare providers involved (see online supplementary file). All patients received this questionnaire on the second day of the care pathway, including a return envelope. For the current analyses, questions concerning patients’ satisfaction with the care pathway and healthcare providers as such were evaluated.

**Statistical analysis**

According to their distribution, continuous variables are either presented as mean and SD or medians with IQR (p25–75). Categorical variables are presented as frequencies with percentages.

To determine which patients benefit most from the care pathway in terms of specific medical interventions (defined as: diagnostic follow-up, referral or change of treatment) and initiation of immunosuppressive treatment following the care pathway, patients with and without medical intervention, and patients with and without initiation of immunosuppressives were compared. Univariable and multivariable regression analyses were used to determine characteristics associated with the initiation of any medical intervention and with the initiation of immunosuppressive medication. To identify patient characteristics commonly available from history taking, physical examination and laboratory testing, and thus help to identify those patients who benefit most from the care pathway, multivariate regression analyses were also performed excluding results generated by diagnostic tests performed as part of the care programme. Comparisons between the characteristics of patients who did and who did not return the satisfaction questionnaire were done by means of unpaired t tests, Mann-Whitney U tests, or χ² tests, where appropriate.

For all data, an available case analysis was performed. Data entry was performed using Microsoft Office Access 2003. All statistical analyses were executed using SPSS V20.0 software (SPSS Inc, Chicago, USA).

**RESULTS**

**Patient population**

Two hundred and forty patients were referred to the healthcare programme between 2009 and 2012. The diagnosis SSC was confirmed in 226 patients. These patients were mostly women (n=186, 82%), Caucasian (n=151, 67%) and, on average, 54 years old (SD 14.5). Patients had a median disease duration of 4 years (range 1–11 years) and 75 of them (32%) had DCSSc (table 1).

**Adherence with the care pathway**

All patients filled in the questionnaires and underwent laboratory investigations, ECG, 6MWD, HRCT, PFT and echocardiography.

Evaluation by CPET was performed in 209 patients (92%), since 17 patients were not able to cycle, mainly caused by musculoskeletal disability. NVC was performed in 86 patients (38%), of whom 78 patients (91%) had a SSC pattern. An early pattern was seen in 10 patients, an active pattern in 37 patients and a late pattern in 31 patients.
One hundred and nineteen (53%) patients visited an occupational therapist, and 80 (35%) patients visited a social worker.

Medical interventions resulting from the care pathway
The frequency of specific interventions resulting from the care pathway is shown in Figure 1. In total, one or...
more additional diagnostic or therapeutic interventions were initiated in 191 (85%) of the patients. In 95 patients (42%), additional diagnostic tests were advocated. Among the patients with elevated SPAP (range 36–80 mm Hg; n=29; >35 mm Hg–≤40 mm Hg; n=10;>40 mm Hg–≤45 mm Hg; n=17;>45 mm Hg), six underwent RHC, which confirmed the diagnosis of PAH in four patients, resulting in initiation of bosentan. Two patients were classified as having pulmonary hypertension based on SPAP ≥50 mm Hg by echocardiography alone (n=1: SPAP 61 mm Hg, n=1 SPAP 54 mm Hg) and started with bosentan without confirmation of actual pulmonary pressures by RHC, as the general condition of the patient did not allow such a procedure. The remaining patients with elevated SPAP were considered as either borderline pulmonary hypertension, and/or pulmonary hypertension due to left heart disease, or chronic lung disease. As no direct therapeutic intervention was expected following RHC, the multidisciplinary PAH group advised to repeat echocardiography within 6–12 months in these cases.

Stringent pulmonary follow-up of ILD was advised by additional PFT in 33 patients, and by additional HRCT thorax in 17 patients.

Additional examinations to obtain more detailed information about specific organ involvement was considered mandatory only in a minority of patients: colonoscopy (n=7) and DXA scan (n=6) were the investigations most frequently advised.

Eighty-eight patients (39%) were referred to medical specialists. Patients were most frequently referred to a cardiologist (n=17), a gastroenterologist (n=16), a pulmonologist (n=14) or a dermatologist (n=14).

Change of treatment
At presentation, 77 patients (34%) were using immunosuppressive medication. Based on the findings during the care pathway, new immunosuppressive treatment was started in 49 patients (22%). Newly prescribed treatment included autologous HSCT, cyclophosphamide, mycophenolate mofetil (MMF), corticosteroids, methotrexate (MTX), AZA, HCQ, rituximab and rituximab/placebo. Two patients started with a combination of drugs: the first with cyclophosphamide and prednisone, and second started with MTX and was included in the RITIS trial. In one patient, treatment with cyclophosphamide was started, and it was advised to switch to MMF maintenance therapy after 6 monthly pulses (female, 27 years of age, wish to conceive in the future). Initiation of immunosuppressive therapy was advocated in presence of: ILD (n=24), high or progressive skin score (n=18), very early SSC (≤6 months); puffy fingers (n=3), arthritis (n=5), renal involvement (n=2), myositis (n=1) and symptomatic pericardial effusion (n=1). In none of the patients with previous HSCT, new immunosuppressive medication was started.

Advices on supportive medication most frequently consisted of changes (prescription, increase, decrease) in calcium channel blockers (n=35) and ACE inhibitors (n=33), or prescription of eye drops (n=33). In 58 patients (26%), medication for peripheral vascular complications was started, most often calcium channel blockers (n=32). In n=9 patients, a combination of different vasoactive drugs and/or antibiotics was prescribed. Fifteen patients started with bosentan, 9 patients started with intravenous iloprost, and in n=12 patients antibiotic treatment for superimposed bacterial infections was prescribed.
Determinants of medical interventions

In total, one or more medical interventions (defined as: (1) need for additional diagnostic follow-up, (2) referral to medical specialists, (3) referral to other healthcare professionals or (4) initiation of immunosuppressive treatment and change of supportive medications), were initiated in 191 (85%) of patients as a result of the care pathway.

Multivariable logistic regression analyses showed that patients with medical interventions had a higher ESR, and more often had telangiectasias than patients without any medical intervention (table 2).

Determinants of initiation of immunosuppressive therapy

Univariate logistic regression analysis of disease characteristics, and results of diagnostic investigations showed that younger age, DcSSc, shorter duration of disease and (non-)RP, proximal muscular weakness, dyspnoea, anti-Scl-70, pulmonary crackles, NSIP, higher mRSS, ESR, CRP and CPK, and lower VC, wattage and VO₂ max at CPET, were significantly associated with the start of new immunosuppressive therapy. Presence of telangiectasias, and ACA decreased the chance for start of new immunosuppressive therapy (table 3).

Multivariable analysis showed that next to NSIP according to HRCT, CPK, mRSS and age were associated with start of immunosuppressive treatment, with NSIP according to HRCT having the largest impact (table 3). Repeated multivariate testing including only variables resulting from history taking, physical examination and laboratory testing identified age (OR 0.96; 95% CI (0.93 to 0.99)), skin score (mRSS; OR 1.12; 95% CI (1.04 to 1.21)), and presence of ACA (OR 0.19; 95% CI 0.04 to 0.85)) as most significantly associated with initiation of immunosuppressive therapy. Higher age and presence of ACA decreased the change of start of immunosuppressives.

Non-medical interventions resulting from the care pathway

Forty-three patients (19%) were referred to other healthcare professionals, most frequently to a physical therapist in primary setting (n=26) for an exercise programme, and/or improvement of physical functioning, an occupational therapist (n=7) for assistive devices and/or adjustments at home, or a dietician (n=7).

Non-pharmacological treatment advices provided by the specialised nurse included therapeutic gloves (9% silver; n=16), stopping smoking (n=13), and other lifestyle advices including dietary advices (n=5).

Patient satisfaction

In total, 96 patients (42%) returned the questionnaire concerning different aspects of the care pathway.

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Table 2 Univariable and multivariable logistic regression analysis for patients with and without any medical intervention

<table>
<thead>
<tr>
<th>Variable</th>
<th>Univariable OR (95% CI)</th>
<th>p Value</th>
<th>Multivariable OR (95% CI)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sociodemographics</td>
<td></td>
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</tr>
<tr>
<td>Age, years, mean (SD)</td>
<td>1.02 (0.99 to 1.04)</td>
<td>0.241</td>
<td>1.00 (0.98 to 1.03)</td>
<td>0.823</td>
</tr>
<tr>
<td>Female, N (%)</td>
<td>1.05 (0.40 to 2.72)</td>
<td>0.925</td>
<td>1.36 (0.50 to 3.70)</td>
<td>0.554</td>
</tr>
<tr>
<td>Disease duration, years, median (IQR)</td>
<td>1.01 (0.96–1.06)</td>
<td>0.699</td>
<td></td>
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<tr>
<td>Onset of RP, years, median (IQR)</td>
<td>1.01 (0.98–1.04)</td>
<td>0.623</td>
<td></td>
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<tr>
<td>Onset of non-RP, years, median (IQR)</td>
<td>1.02 (0.97–1.07)</td>
<td>0.471</td>
<td></td>
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<tr>
<td>Disease characteristics, N (%)</td>
<td></td>
<td></td>
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<tr>
<td>DcSSc</td>
<td>1.05 (0.48 to 2.28)</td>
<td>0.904</td>
<td></td>
<td></td>
</tr>
<tr>
<td>DU</td>
<td>2.59 (0.96 to 7.02)</td>
<td>0.061</td>
<td></td>
<td></td>
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<tr>
<td>Telangiectasia</td>
<td>2.19 (1.05 to 4.54)</td>
<td>0.036</td>
<td>2.27 (1.08 to 4.80)</td>
<td>0.031</td>
</tr>
<tr>
<td>Synovitis</td>
<td>2.04 (0.46 to 9.11)</td>
<td>0.351</td>
<td></td>
<td></td>
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<tr>
<td>Friction rubs</td>
<td>0.54 (0.10 to 2.77)</td>
<td>0.456</td>
<td></td>
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<tr>
<td>Pulmonary crackles</td>
<td>2.50 (0.32 to 19.73)</td>
<td>0.385</td>
<td></td>
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<tr>
<td>MRSS, median (IQR)</td>
<td>1.73 (0.71 to 4.18)</td>
<td>0.226</td>
<td></td>
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<tr>
<td>Anti-Scl-70 antibody</td>
<td>1.44 (0.59 to 3.56)</td>
<td>0.426</td>
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<tr>
<td>Anticentromere antibody</td>
<td>0.65 (0.30 to 1.40)</td>
<td>0.271</td>
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<tr>
<td>Diagnostic investigations</td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>ESR, mm, median (IQR)</td>
<td>1.04 (1.01–1.07)</td>
<td>0.013</td>
<td>1.04 (1.01–1.07)</td>
<td>0.012</td>
</tr>
<tr>
<td>CRP, mg/dL, median (IQR)</td>
<td>1.06 (0.96–1.16)</td>
<td>0.246</td>
<td></td>
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<tr>
<td>CPK, mg/dL, median (IQR)</td>
<td>1.00 (1.00–1.00)</td>
<td>0.743</td>
<td></td>
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<tr>
<td>Vital capacity % of predicted, mean (SD)</td>
<td>0.99 (0.97 to 1.01)</td>
<td>0.266</td>
<td></td>
<td></td>
</tr>
<tr>
<td>NSIP according HRCT, N (%)</td>
<td>1.80 (0.74 to 4.37)</td>
<td>0.195</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CPET; maximum VO₂% of predicted, mean (SD)</td>
<td>0.99 (0.97 to 1.00)</td>
<td>0.061</td>
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</table>

CPET, cardiopulmonary exercise test; CPK, creatine phosphokinase; CRP, C reactive protein; DcSSc, diffuse cutaneous SSc; DU, digital ulcers; ESR, erythrocyte sedimentation rate; MRSS, modified Rodnan Skin Score; NSIP, non-specific interstitial pneumonia; VO₂ max, maximum volume oxygen.
Table 3  Univariable and multivariable logistic regression analysis for patients who started and did not start with immunosuppressive therapy as a result from the care pathway

<table>
<thead>
<tr>
<th>Sociodemographics</th>
<th>Start N=49</th>
<th>Not started N=177</th>
<th>Univariable OR (95% CI) p Value</th>
<th>Multivariable OR (95% CI) p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, years, mean (SD)</td>
<td>48 (15)</td>
<td>56 (14)</td>
<td>0.96 (0.94 to 0.98) &lt;0.001</td>
<td>0.94 (0.91 to 0.98) 0.005</td>
</tr>
<tr>
<td>Female, N (%)</td>
<td>38 (78)</td>
<td>140 (86)</td>
<td>1.76 (0.79 to 3.93) 0.167</td>
<td>1.76 (0.79 to 3.93) 0.167</td>
</tr>
<tr>
<td>Disease duration, years, median (IQR)</td>
<td>1 (0–4)</td>
<td>4 (1–12)</td>
<td>0.93 (0.88 to 0.98) 0.012</td>
<td>0.93 (0.88 to 0.98) 0.012</td>
</tr>
<tr>
<td>Onset of RP, years, median (IQR)</td>
<td>6 (2–16)</td>
<td>13 (6–21)</td>
<td>0.96 (0.93 to 0.99) 0.020</td>
<td>0.96 (0.93 to 0.99) 0.020</td>
</tr>
<tr>
<td>Onset of non-RP, years, median (IQR)</td>
<td>3 (1–5)</td>
<td>7 (3–13)</td>
<td>0.94 (0.89 to 0.99) 0.016</td>
<td>0.94 (0.89 to 0.99) 0.016</td>
</tr>
<tr>
<td>Disease characteristics, N (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>DcSSc</td>
<td>26 (53)</td>
<td>33 (56)</td>
<td>4.45 (2.26 to 8.78) &lt;0.001</td>
<td>1.33 (0.35 to 5.05) 0.676</td>
</tr>
<tr>
<td>Digital ulcers</td>
<td>11 (22)</td>
<td>49 (30)</td>
<td>0.66 (0.31 to 1.40) 0.281</td>
<td>0.66 (0.31 to 1.40) 0.281</td>
</tr>
<tr>
<td>Telangiectasia</td>
<td>20 (41)</td>
<td>107 (66)</td>
<td>0.35 (0.18 to 0.68) 0.002</td>
<td>0.46 (0.16 to 1.30) 0.155</td>
</tr>
<tr>
<td>Synovitis</td>
<td>8 (16)</td>
<td>15 (9)</td>
<td>1.93 (0.76 to 4.86) 0.165</td>
<td>1.93 (0.76 to 4.86) 0.165</td>
</tr>
<tr>
<td>Friction rubs</td>
<td>4 (8)</td>
<td>4 (3)</td>
<td>3.53 (0.85 to 14.69) 0.083</td>
<td>3.53 (0.85 to 14.69) 0.083</td>
</tr>
<tr>
<td>Proximal muscular weakness</td>
<td>8 (16)</td>
<td>6 (4)</td>
<td>5.07 (1.67 to 15.44) 0.004</td>
<td>5.06 (0.93 to 27.68) 0.061</td>
</tr>
<tr>
<td>Pulmonary crackles</td>
<td>21 (44)</td>
<td>36 (22)</td>
<td>2.72 (1.38 to 5.37) 0.004</td>
<td>2.44 (0.74 to 8.02) 0.142</td>
</tr>
<tr>
<td>mRSS, median (IQR)</td>
<td>5 (2–20)</td>
<td>2 (0–5)</td>
<td>1.13 (1.07 to 1.18) &lt;0.001</td>
<td>1.09 (1.00 to 1.01) 0.042</td>
</tr>
<tr>
<td>Anti-Scl-70 antibody</td>
<td>18 (37)</td>
<td>28 (17)</td>
<td>4.07 (1.88 to 8.82) &lt;0.001</td>
<td>4.07 (1.88 to 8.82) &lt;0.001</td>
</tr>
<tr>
<td>Anticentromere antibody</td>
<td>4 (8)</td>
<td>72 (44)</td>
<td>0.11 (0.04 to 0.33) &lt;0.001</td>
<td>0.11 (0.04 to 0.33) &lt;0.001</td>
</tr>
<tr>
<td>Diagnostic investigations</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ESR, mm, median (IQR)</td>
<td>19 (11–41)</td>
<td>14 (6–28)</td>
<td>1.02 (1.00 to 1.03) 0.033</td>
<td>1.02 (1.00 to 1.03) 0.033</td>
</tr>
<tr>
<td>CRP, mg/dL, median (IQR)</td>
<td>3 (3–7)</td>
<td>3 (3–4)</td>
<td>1.04 (1.01 to 1.07) 0.020</td>
<td>1.04 (1.01 to 1.07) 0.020</td>
</tr>
<tr>
<td>CPK, mg/dL, median (IQR)</td>
<td>95 (54–174)</td>
<td>85 (61–114)</td>
<td>1.01 (1.00 to 1.01) 0.009</td>
<td>1.01 (1.00 to 1.01) 0.009</td>
</tr>
<tr>
<td>Vital capacity % of predicted, mean (SD)</td>
<td>86 (19)</td>
<td>98.6 (21)</td>
<td>0.97 (0.95 to 0.99) &lt;0.001</td>
<td>0.97 (0.95 to 0.99) &lt;0.001</td>
</tr>
<tr>
<td>NSIP according HRCT, N (%)</td>
<td>27 (55)</td>
<td>36 (22)</td>
<td>4.13 (2.10 to 8.10) &lt;0.001</td>
<td>4.89 (1.561 to 15.30) 0.006</td>
</tr>
<tr>
<td>CPET; maximum VO2% of predicted, mean (SD)</td>
<td>79 (20)</td>
<td>91 (26)</td>
<td>0.98 (0.97 to 0.99) 0.006</td>
<td>0.98 (0.97 to 0.99) 0.006</td>
</tr>
</tbody>
</table>

DISCUSSION

The present study describes the medical management of 226 patients with SSc referred to a 2-day comprehensive diagnostic care pathway. In 191 (85%) patients, the care pathway resulted in specific advices on additional diagnostic follow-up, referral or change of treatment. Of patient characteristics only telangiectasias and ESR were associated with any medical intervention. Commonly available patient characteristics associated with change of immunosuppressives included lower age, absence of ACA and higher mRSS. These data indicate that a standardised comprehensive care pathway is of benefit for the far majority of patients with SSc, regardless of SSc subtype and disease duration. Systematic organ screening has been shown to contribute to survival of patients with SSc over the past decades, mainly in patients with DcSSc. This observation, in combination with preference advices on additional diagnostic care pathway. In 191(85%) patients, the care pathway resulted in specific advices on additional diagnostic follow-up, referral or change of treatment. Of patient characteristics only telangiectasias and ESR were associated with any medical intervention. Commonly available patient characteristics associated with change of immunosuppressives included lower age, absence of ACA and higher mRSS. These data indicate that a standardised comprehensive care pathway is of benefit for the far majority of patients with SSc, regardless of SSc subtype and disease duration. Systematic organ screening has been shown to contribute to survival of patients with SSc over the past decades, mainly in patients with DcSSc. This observation, in combination with preference advices on additional diagnostic care pathway.
was high and patient satisfaction rates were very high. This latter finding probably reflects the successful match between patients’ needs and care provision, as the care pathway was developed based on the questionnaire on patients’ needs and preferences. Whether implementation of the described care pathway will contribute to diminishing the number of visits to the hospital in the long run, remains to be determined. Most likely, benefits in terms of medical interventions will be numerically different after following annual visits. Future evaluations will have to demonstrate in which subgroups of patients repeated extensive annual screening is most useful and whether standardised and multidisciplinary care as applied in the care pathway will contribute to improved outcome.

As one or more medical interventions took place in 191 of 226 patients, the care pathway did not result in any specific medical intervention in 1 of 6–7 patients. To evaluate in which patients referral to the care pathway was most useful, multivariable regression analyses were performed. Presence of telangiectasias was associated with need for any medical intervention, while it was protective for starting immunosuppressive treatment. It is known that presence of telangiectasias is associated with vascular complications including digital ulcers and pulmonary hypertension. Indeed, subgroup comparisons within our own cohort between patients with telangiectasias and patients without telangiectasias showed that patients with telangiectasias had more often calcinosis, more often had a change in vasoactive medication, and they were discussed more frequently in the PAH meeting. Patients without telangiectasias needed more stringent follow-up on pulmonary fibrosis reflected by need for additional PFTs (see online supplementary table S1). This observation is thus in line with previous studies showing the association between vascular complications and telangiectasias in SSc. Consequently, the number and type of interventions resulting from the care pathway is most likely different for the different subgroups of SSc, as reflected by the differential association of telangiectasias with the different interventions.

Our study confirms previous findings about clinically important investigations in evaluation of severity and activity of SSc. Multivariable analysis showed that age, mRSS, proximal muscle weakness, CPK and NSIP, according to HRCT-thorax were significantly associated with the start of new immunosuppressive therapy. These findings are explained by the fact that skin involvement and pulmonary involvement were the most frequent indications for immunosuppressive treatment, and thus, reflect a certain circularity. In addition, muscle involvement was identified as a relevant parameter, reflected by both CPK and proximal muscle weakness. SSc overlap with myositis is found in approximately 15% of the patients with SSc, and is associated with more extensive disease. Although these parameters seem to be rather non-specific, two important recent studies also identified CPK level and myopathy as important predictors for worse outcome. As no circularity can be involved specifically for these two parameters, we do believe that our data confirm previous data on CPK and myopathy being important predictors for worse prognosis, and thus, should probably be taken into account in therapeutic decision-making.

By contrast with previous studies, no attributive value for start of immunosuppressive therapy of type of SSc, PFTs and cardiac parameters was found. Several factors can account for a lack of association between cardiac parameters and of immunosuppressive treatment. First, only in case of active myocarditis, immunosuppressive treatment is advocated. In case of diastolic dysfunction, or significant arrhythmias, primary interventions consist of ACE inhibitors, antiarrhythmics and placement of implantable cardioverter defibrillator where appropriate. Finally, the included diagnostic procedures might not have revealed subclinical myocardial involvement, as plain echocardiography is rather insensitive for this goal, and the majority of patients already received treatment with calcium channel blockers and/or ACE inhibitors for different indications. Another interesting finding was that, apart from consultations with a nurse and physical therapist, more than half the patients took the opportunity to visit an occupational therapist and/or social worker. This observation shows that many patients encounter problems which can clearly not be solved by medical treatment alone. The relatively high proportion of patients in the present study expressing a need to see additional healthcare providers again underscores that more research into the most appropriate non-pharmacological management strategies in this patient group is warranted.

This study has a number of limitations which should be taken into account when interpreting the results. First, selection bias cannot be excluded, as patients referred to the care programme in a tertiary care setting might not represent the total population of patients with SSc. However, comparison with other cohorts of patients with SSc evaluated by tertiary care centres, including the EUSTAR centres, showed that social and demographic characteristics, disease severity and functional status of our cohort are quite similar. At admission, 34% of the patients were treated with immunosuppressive therapy, which is also comparable to reports from the EUSTAR cohort. Second, start of immunosuppressive therapy was based on consensus between experts during a multidisciplinary discussion, in the absence of a gold standard precisely guiding treatment decisions in SSc. Whether treatment decisions in the current cohort were adequate remains to be determined. Ideally, one should dispose of a control group receiving standard care to compare number of medical interventions, progression-free survival and quality of life. Finally, the current study describes results of the care pathway at short term. Long-term follow-up is needed to evaluate the impact on patients’ health status and healthcare usage. These results could guide further improvements of the contents and organisation of care.
In conclusion, our data demonstrate that a comprehensive diagnostic and multidisciplinary care pathway for patients with SSC in a day-care setting, combining as such annual screening on organ involvement as indicated according to (inter)national guidelines with multidisciplinary team care, is feasible. In general, adherence to the care pathway was very high, and resulted in medical interventions in the majority of patients, independent of SSC subtype and disease duration. Future evaluations are needed to evaluate impact on disease outcome.

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Contributors JM collected the data, was involved in statistical analysis, writing and critically reviewing the manuscript and approved the final version. AAS collected the data, was involved in critically reviewing the manuscript and approved the final version. NAM collected the data, was involved in critically reviewing the manuscript, and approved the final version. LJM was involved in CT data acquisition, critically reviewing the manuscript, and approved the final version. TS was involved in statistical analysis, and approved the final version.

Competing interests JM was supported by an unrestricted educational grant from Actelion Pharmaceuticals Nederland BV (Woerden, The Netherlands). JM was supported by an unrestricted educational grant and critically reviewing the manuscript, and approved final version. TPMVV was involved in the design of the study, was involved in writing and critically reviewing the manuscript, and approved the final version. TWJH reviewed the data statistics, and approved final version. MKN collected the data, was involved in critically reviewing the manuscript, and approved the final version.

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