

# Performance of Forced Vital Capacity and Lung Diffusion Cutpoints for Associated Radiographic Interstitial Lung Disease in Systemic Sclerosis

Kimberly Showalter , Aileen Hoffmann, Gerald Rouleau , David Aaby, Jungwha Lee, Carrie Richardson, Jane Dematte, Rishi Agrawal, Rowland W. Chang, and Monique Hinchcliff

**ABSTRACT.** *Objective.* Forced vital capacity (FVC) and DLCO are used for screening of systemic sclerosis–associated interstitial lung disease (SSc-ILD). The study purpose was to determine the sensitivity, specificity, and negative predictive value (NPV) (proportion of true negative screening tests) of FVC and DLCO thresholds for SSc-ILD on chest high-resolution computed tomography (HRCT) scans. *Methods.* Patients fulfilling American College of Rheumatology 2013 SSc criteria with a chest HRCT scan and pulmonary function tests (PFT) were studied. A thoracic radiologist quantified radiographic ILD. Optimal FVC and DLCO % predicted thresholds for ILD were identified using receiver-operating characteristic curves. The FVC and DLCO combinations with greatest sensitivity and specificity were also determined. Subanalysis was performed in patients with positive Scl-70 autoantibodies. *Results.* The study included 265 patients. Of 188 (71%) with radiographic ILD, 59 (31%) had “normal” FVC ( $\geq 80\%$  predicted), and 65 out of 151 (43%) had “normal” DLCO ( $\geq 60\%$  predicted). FVC  $< 80\%$  (sensitivity 0.69, specificity 0.73), and DLCO  $< 62\%$  (sensitivity 0.60, specificity 0.70) were optimal thresholds for radiographic SSc-ILD. All FVC and DLCO threshold combinations evaluated had NPV  $< 0.70$ . The NPV for radiographic ILD for FVC  $< 80\%$  was lower in patients with positive Scl-70 autoantibody (NPV = 0.05) compared to negative Scl-70 autoantibody (NPV = 0.57). *Conclusion.* Radiographic ILD is prevalent in SSc despite “normal” PFT. No % predicted FVC or DLCO threshold combinations yielded high NPV for SSc-ILD screening. “Normal” FVC and DLCO in patients with SSc, especially those with positive Scl-70 autoantibodies, should not obviate consideration of HRCT for ILD evaluation. (First Release October 1 2018; J Rheumatol 2018;45:1572–6; doi:10.3899/jrheum.171362)

## Key Indexing Terms:

SYSTEMIC SCLEROSIS    INTERSTITIAL LUNG DISEASE    FORCED VITAL CAPACITY

From the Department of Medicine, Division of Rheumatology, Division of Pulmonary and Critical Care Medicine, Department of Preventive Medicine, Institute for Public Health and Medicine, Department of Radiology, Northwestern University Feinberg School of Medicine, Chicago, Illinois, USA.

K. Showalter, MD, Department of Medicine, Northwestern University Feinberg School of Medicine; A. Hoffmann, MS, Department of Medicine, Division of Rheumatology, Northwestern University Feinberg School of Medicine; G. Rouleau, MS, Department of Preventive Medicine, Northwestern University Feinberg School of Medicine; D. Aaby, MS, Department of Preventive Medicine, Northwestern University Feinberg School of Medicine; J. Lee, PhD, MPH, Department of Preventive Medicine, and Institute for Public Health and Medicine, Northwestern University Feinberg School of Medicine; C. Richardson, MD, Department of Medicine, Northwestern University Feinberg School of Medicine; J. Dematte, MD, MBA, Department of Medicine, Division of Pulmonary and Critical Care Medicine, Northwestern University Feinberg School of Medicine; R. Agrawal, MD, Department of Radiology, Northwestern University Feinberg School of Medicine; R.W. Chang, MD, MPH, Department of Medicine, Division of Rheumatology, and Department of Preventive Medicine, and Institute for Public Health and Medicine, Northwestern University Feinberg School of Medicine; M. Hinchcliff, MD, MS, Department of Medicine, Division of Rheumatology, and Department of Preventive Medicine, and Institute for Public Health and Medicine, Northwestern University Feinberg School of Medicine.

Address correspondence to Dr. M. Hinchcliff, Associate Professor of Medicine, Northwestern University Feinberg School of Medicine,

Department of Medicine, Division of Rheumatology, McGaw Pavilion, 240 E. Huron St., Chicago, Illinois 60611, USA.

E-mail: m-hinchcliff@northwestern.edu

Accepted for publication June 1, 2018.

Interstitial lung disease (ILD) is a leading cause of death in patients with systemic sclerosis (SSc)<sup>1</sup>, and chest high-resolution computed tomography (HRCT) is the gold standard diagnostic test<sup>2</sup>. Pulmonary function tests (PFT) including forced vital capacity (FVC) and DLCO % predicted are frequently used to screen for ILD to avoid HRCT-associated radiation. However, evidence shows that a significant number of SSc patients with FVC values within the normal range ( $\geq 80\%$  predicted) demonstrate radiographic ILD<sup>3,4</sup>. We hypothesized that optimal individual FVC and DLCO % predicted thresholds, as well as optimal FVC and DLCO combinations, could be identified with acceptably high negative predictive value (NPV) for associated radiographic SSc-ILD. The prevalence of a disease in a population affects the NPV, and radiographic ILD is common in patients with SSc<sup>1,3,4,5</sup>. One HRCT study of 215 consec-

utive patients with SSc found limited ILD (< 10% of lung involvement on HRCT) in almost 50% of patients<sup>5</sup>. However, Suliman, *et al*<sup>3</sup> evaluated 102 consecutive patients with SSc at the University Hospital Zurich, and found that 40 out of 75 patients (53%) with an FVC > 80% predicted had significant radiographic ILD (> 20% lung involvement according to Goh, *et al*<sup>5</sup>). Using data from patients with SSc at our center, our aims were to (1) identify optimal individual FVC and DLCO % predicted cutpoints for associated ILD on HRCT, and (2) to determine the sensitivity, specificity, and NPV of varying individual and combined FVC and DLCO % predicted thresholds for associated ILD on HRCT.

## MATERIALS AND METHODS

Our present study was approved by the Northwestern University Institutional Review Board (STU00066807). A waiver of informed consent was obtained for this retrospective study because all patients had given consent for the Northwestern Scleroderma Patient Registry, and permission to review electronic health records is included in that consent (STU00002669).

Patients had sine, diffuse, or limited cutaneous SSc, fulfilled American College of Rheumatology 2013 SSc criteria, and had a PFT within 12 months of a chest HRCT scan<sup>6</sup>. Patients who had pulmonary procedures or diagnoses that could potentially affect PFT results including lung transplant, lobectomy, or lung malignancy were excluded. Patients with comorbid pulmonary hypertension (PH)-ILD were included in the analyses.

Clinical data [SSc subtype, smoking history (current, previous, or never), alcohol use (present or absent), and modified Rodnan skin score] were collected from rheumatology clinic notes within 1 year of HRCT date. SSc disease duration was defined as duration between first non-Raynaud SSc symptom and HRCT scan date. Mean  $\pm$  SD or n (%) was reported for baseline characteristics.

Chest HRCT is defined by the use of images at 1–2 mm slice thickness reconstructed using specialized computer algorithms to increase the sharpness of lung parenchyma. Ground glass opacities (GGO) and fine reticulation on HRCT are characteristic of SSc-ILD<sup>7</sup>. Although lung biopsies are seldom performed for diagnosis and survival is independent of histology, nonspecific interstitial pneumonia (NSIP) followed by usual interstitial pneumonia (UIP) are the most common histological patterns in patients with SSc<sup>1</sup>. An experienced thoracic radiologist (RA), blinded to clinical data, quantified ILD on HRCT according to the method published by Kazerooni, *et al* and manually reviewed each HRCT to confirm lung changes consistent with ILD as described<sup>8</sup>. Briefly, each lobe was scored for GGO and fibrosis involvement using a modified Likert scale (0 = no disease; 1 = < 5% involvement of the lobe, 2 = 6–25% involvement; 3 = 26–50% involvement; 4 = 51–75% involvement, and 5 = 76–100%). The total lung score was the sum of the GGO and fibrosis scores for each lobe. ILD was defined as present if the morphologic features of the disease matched a recognized pattern of disease (i.e., NSIP, UIP, etc.).

PFT data obtained at our center or at an outside institution were analyzed. All available FVC values were studied and correlation with total lung capacity (TLC) was determined using Pearson's correlation coefficient. However, to optimize DLCO data quality, only PFT with inspiratory vital capacity (IVC) and FVC within 0.85 of each other were included<sup>9</sup>. Patients with poor-quality DLCO data as indicated by an IVC:FVC ratio < 0.85 were excluded from the analyses of DLCO. However, all FVC data were analyzed. Percent predicted DLCO was corrected for hemoglobin values obtained from a complete blood count (CBC) performed within 6 months of the PFT date. National Health and Nutrition Examination Survey III (NHANES III) reference values were used<sup>10</sup>.

Two independent receiver-operating characteristic (ROC) curves were generated for radiographic SSc-ILD at varying FVC and DLCO % predicted values. For all analyses, R (version 3.4.0) ROCR package was used to

determine the optimal cutpoints, defined as the greatest combined sensitivity and specificity, for radiographic SSc-ILD. Various established PFT thresholds were evaluated. Subanalysis was performed according to Scl-70 autoantibody profile, because SSc-ILD is highly prevalent in patients with antitopoisomerase I (Scl-70) autoantibody positivity<sup>1</sup>.

## RESULTS

Of 729 patients enrolled in the Northwestern Scleroderma Program, 404 had interpretable HRCT images for manual review. One hundred thirty-nine patients were excluded: 30 with an overlap autoimmune diagnosis or a confounding pulmonary procedure, and 109 with PFT > 1 year from HRCT date. The remaining 265 studies were analyzed. The DLCO values from 29 PFT were excluded because the IVC to FVC ratio was  $\leq$  0.85, thereby rendering the DLCO potentially inaccurate<sup>9</sup>. Of the remaining 236 patients, 214 had hemoglobin assessed within 6 months of PFT for DLCO adjustment (Supplementary Figure 1, available with the online version of this article) and were included for DLCO and combined threshold analyses.

The majority of patients were women (81.5%) and had diffuse cutaneous SSc (49%; Table 1). Antitopoisomerase I serum autoantibodies were present in 78 (30%). The mean SSc disease duration (time between first non-Raynaud SSc symptom to HRCT scan) was 6.4 (range 0–42) years with a median of 2 years. Radiographic ILD was present in 188 subjects (71%). There was a strong linear correlation between FVC and TLC ( $r = 0.85$ ), thus FVC was used in the remainder of the analyses. Fifty-nine out of 188 (31%) had normal FVC ( $\geq 80$  % predicted). A total of 151 patients had available and

Table 1. Baseline study population characteristics (n = 265).

Characteristics	Mean $\pm$ SD or n (%)
Age at time of HRCT, yrs	50 $\pm$ 12
Sex, female	216 (82)
Body mass index, kg/m <sup>2</sup>	29 $\pm$ 6
Ethnicity, white	192 (73)
Smoker, current or former	97 (37)
Alcohol, current	96 (46)
Proton pump inhibitor use, current	171 (65)
SSc disease subtype	
Diffuse cutaneous	129 (49)
Limited cutaneous	131 (49)
Sine scleroderma	5 (2)
Years between first non-Raynaud SSc symptom and HRCT date	6 $\pm$ 11
SSc specific antibodies, positive	
Antitopoisomerase I (Scl-70)	78 (30)
Anticentromere antibodies	48 (18)
Anti-RNA polymerase III	59 (23)
Modified Rodnan skin score	13 $\pm$ 11
Radiographic ILD present	188 (71)
FVC % predicted	76 $\pm$ 18
DLCO % predicted	60 $\pm$ 20

SSc: systemic sclerosis; HRCT: high-resolution computed tomography; ILD: interstitial lung disease; FVC: forced vital capacity.

accurate DLCO measurement (IVC:FVC within 85% of each other) and radiographic ILD, and 65 of those 151 (43%) had normal DLCO ( $\geq 60\%$  predicted). Thirty-one out of 214 (14%) of these patients had both FVC and DLCO % predicted within normal range and radiographic ILD. To ensure that patients with poor PFT were not systematically excluded from DLCO analysis, we compared the FVC between the group included versus excluded for low IVC on the DLCO maneuver. Those with DLCO excluded owing to low IVC:FVC ratio did not have a significantly lower FVC than those with acceptable DLCO maneuvers.

For associated ILD, the data-derived optimal FVC and DLCO cutpoints were 80% predicted (sensitivity 0.69, specificity 0.73) for FVC, and 62% predicted (sensitivity 0.60, specificity 0.70) for DLCO (Figure 1, Table 2). The combination of DLCO  $< 62\%$  or FVC  $< 80\%$  predicted had a sensitivity and specificity for ILD of 0.80 and 0.56, respectively. The sensitivity and specificity for various combinations of traditional and data-derived optimal FVC and DLCO cutpoints for ILD are reported in Table 2. Using an ILD screening algorithm of meeting either FVC  $< 80\%$  or DLCO  $< 62\%$  predicted, 82

persons in our cohort (31%) would screen negative for ILD (NPV 0.53). A more liberal algorithm of either FVC  $< 80\%$  or DLCO  $< 70\%$  predicted would inappropriately classify 58 persons (22%; NPV 0.57), while an FVC  $< 80\%$  or DLCO  $< 80\%$  predicted would inappropriately classify 32 persons (12%) as lacking ILD (NPV 0.65; Table 2).

Subanalysis was performed based on Scl-70 autoantibody status. The NPV for radiographic ILD for FVC  $< 80\%$  was lower in patients with positive Scl-70 autoantibody (NPV 0.05) compared to those negative (NPV 0.57; Table 2).

## DISCUSSION

The study aim was to determine the sensitivity, specificity, and NPV of varying independent and combined FVC and DLCO % predicted thresholds for associated radiographic ILD in SSc. The clinical relevance of new FVC and DLCO threshold values is to inform the development of more rational and data-driven ILD screening protocols that include HRCT in the most at-risk patients. Our ROC curve results show FVC = 80% and DLCO = 62% predicted are the optimal (maximized combined sensitivity and specificity)

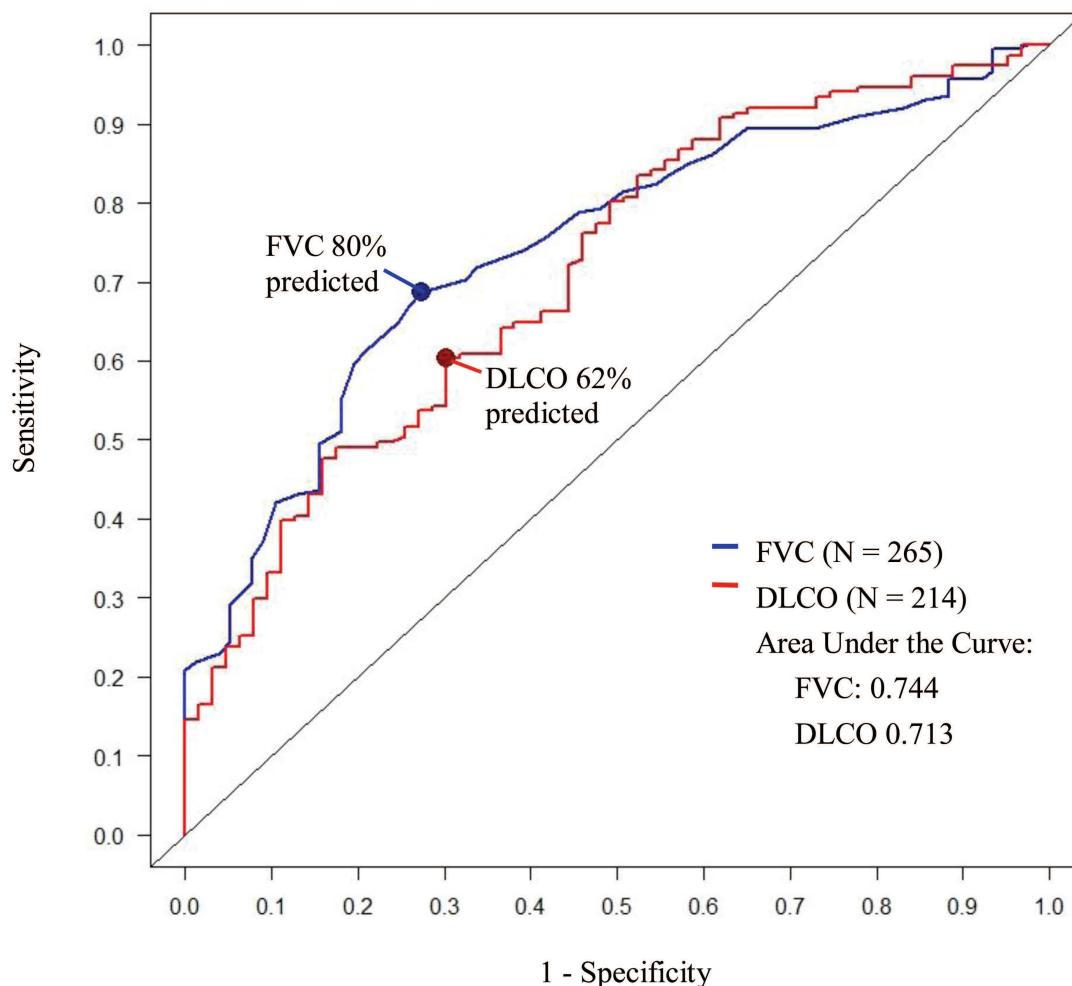


Figure 1. Receiver-operating characteristic curves for % predicted forced vital capacity (FVC) and DLCO, demonstrating the performance of varying FVC and DLCO % predicted cutpoints for associated radiographic interstitial lung disease in systemic sclerosis.

Table 2. Performance of pulmonary function test thresholds for prevalent radiographic interstitial lung disease on chest high-resolution computed tomography images in patients with systemic sclerosis.

Variables	Sensitivity	Specificity	Positive Predictive Value	Negative Predictive Value
Entire cohort				
FVC % predicted, n = 265				
< 80 (conventional and optimal)	0.69	0.73	0.86	0.49
DLCO % predicted, n = 214				
< 60 (conventional)	0.58	0.70	0.82	0.41
< 62 (optimal)	0.60	0.70	0.83	0.42
< 70 (alternative)	0.80	0.51	0.80	0.52
< 80 (alternative)	0.92	0.32	0.77	0.63
Combination of PFT thresholds % predicted, n = 214				
FVC < 80 and DLCO < 60	0.46	0.81	0.85	0.38
FVC < 80 or DLCO < 60	0.79	0.57	0.82	0.53
FVC < 80 and DLCO < 62	0.49	0.81	0.86	0.40
FVC < 80 or DLCO < 62	0.80	0.56	0.81	0.53
FVC < 80 and DLCO < 65	0.53	0.78	0.85	0.41
FVC < 80 or DLCO < 65	0.82	0.46	0.78	0.52
FVC < 80 and DLCO < 70	0.61	0.76	0.86	0.45
FVC < 80 or DLCO < 70	0.87	0.43	0.78	0.57
FVC < 80 and DLCO < 80	0.66	0.73	0.85	0.47
FVC < 80 or DLCO < 80	0.94	0.27	0.76	0.65
Autoantibody Scl-70–positive patients				
FVC < 80, n = 78				
DLCO < 60, n = 62	0.75	0.05	0.98	0.05
FVC < 80 or DLCO < 60, n = 62	0.70	1.00	1.00	0.10
FVC < 80 or DLCO < 60, n = 62	0.85	0.50	0.98	0.10
Autoantibody Scl-70–negative patients				
FVC < 80, n = 183				
DLCO < 60, n = 150	0.63	0.73	0.78	0.57
FVC < 80 or DLCO < 60, n = 150	0.50	0.68	0.70	0.48
FVC < 80 or DLCO < 60, n = 150	0.74	0.57	0.72	0.60

FVC  $\geq$  80% and DLCO  $\geq$  60% predicted represent traditional normal cutpoints based on 95% CI in healthy population. PFT: pulmonary function test; FVC: forced vital capacity.

thresholds for SSc-ILD. Similar area under the curve (AUC) for FVC (AUC = 0.74) and DLCO (AUC = 0.71) indicate that FVC and DLCO are both “moderately accurate” tests for radiographic ILD in SSc patients, with 0.70–0.90 considered “moderately accurate,” and  $>$  0.90 considered “highly accurate”<sup>11</sup>. We found no individual or combined FVC and DLCO % predicted algorithm that had high NPV for SSc-ILD screening.

Our study population consisted of predominantly white women, reflecting SSc demographics and the patient population who receive care at our center, and thus our results may not be generalizable to men and racially diverse SSc populations. Only 407 out of 729 registry patients had available HRCT imaging and were included in the analysis. This likely enriched our population for patients with pulmonary disease. However, the 71% prevalence of radiographic ILD in our cohort is comparable to reported prevalence rates in other tertiary care cohorts (36–84%)<sup>3,4,5</sup>. The 30% prevalence of Scl-70 autoantibody positivity in our study cohort is slightly higher than previous reports (20%)<sup>1</sup>. Yet, ILD is known to be more common in SSc patients with Scl-70 antibodies, rendering HRCT examination referral in this group more likely.

Individual and combined, traditional, and data-derived

FVC and DLCO % predicted thresholds had relatively low sensitivity, specificity, and NPV values for associated radiographic ILD. Our data-derived optimal cutpoints for radiographic ILD detection were identical (FVC 80% predicted) or similar (DLCO 62 vs 60% predicted), and did not outperform traditional thresholds of “normal” that are based upon 95% CI from the NHANES III reference population. We examined alternative FVC and DLCO thresholds of 65%, 70%, and 80% predicted and found low NPV for ILD.

Similar to prior study results, “normal” PFT did not discriminate between SSc patients with and without radiographic SSc-ILD. Of our patients with radiographic ILD, defined using the quantitative Kazerooni method, 31% had “normal” FVC ( $\geq$  80% predicted), 43% had “normal” DLCO ( $\geq$  60% predicted), and 14% had both FVC  $\geq$  80 and DLCO  $\geq$  60% predicted. Suliman, *et al* studied 102 patients with SSc and defined radiographic ILD using the more simplistic Goh method that classifies ILD as mild ( $<$  20%), intermediate, or severe ( $>$  20%)<sup>3,5</sup>. They found that 24 out of 102 patients (23%) had normal FVC ( $\geq$  80% predicted) and radiographic ILD and 40 out of 75 patients (53%) with FVC  $\geq$  80% predicted had severe radiographic ILD<sup>3</sup>. Steele, *et al* evaluated various SSc-ILD screening algorithms and defined



ILD as present if (1) crackles were noted on physical examination, (2) radiograph showed interstitial markings consistent with fibrosis, or (3) PFT measures were met: FVC < 70% and FEV1/FVC > 70%; or FVC < 80% and FEV1/FVC > 70%. Both FVC < 70 and < 80% predicted thresholds had low sensitivity for SSc-ILD (54% and 61%, respectively)<sup>4</sup>.

At traditional thresholds (FVC < 80% and DLCO < 60%), we found FVC compared to DLCO to be more specific for associated SSc-ILD (72% vs 69%). This is most likely due to SSc vascular disease, including concurrent PH, which also results in reduced DLCO<sup>12</sup>. In a multivariable analysis by Nihtyanova, *et al*, low DLCO was considered a significant predictor variable in models for both PH and pulmonary fibrosis. In contrast, low FVC was only a predictor for pulmonary fibrosis and not PH<sup>13</sup>. However, our ROC curve results show that these tests perform similarly for ILD detection.

The association between SSc-ILD and positive Scl-70 autoantibodies is well recognized. The NPV for radiographic ILD of 5% (+Scl-70) versus 57% (−Scl-70) for FVC < 80% predicted and of 10% (+Scl-70) versus 48% (−Scl-70) for DLCO < 60% predicted demonstrates how crucial it is to maintain a high index of suspicion for ILD in patients with +Scl-70 antibodies.

Study limitations include analysis of data from participants recruited at 1 site that specializes in SSc care, which limits generalizability. Also, DLCO was not reported for every patient and the absence of CBC within 6 months prevented correction for hemoglobin, leading to exclusion of that patient from analyses. ILD was dichotomized as present or absent, which prevents determination of the effect of ILD severity–varying PFT thresholds. Finally, we excluded patients with lung malignancy, transplant, and lung resection, and thus limited our evaluation of these patients, who may have severe ILD. Study strengths include our large sample size (265 vs 102 analyzed in Suliman, *et al*'s cohort<sup>3</sup>), and our use of stringent criteria to identify high-quality PFT (inclusion of DLCO with hemoglobin performed within 6 mos and exclusion of DLCO values from PFT with IVC:FVC ratio < 0.85), to ensure DLCO quality<sup>9</sup>. The ILD scoring system also differed between studies as described above. We also add to the literature by examining a variety of cutpoints to determine whether new PFT thresholds exist that could be used to screen for SSc-ILD. Our study is limited to a single center specializing in SSc care, which may limit generalizability. As a cross-sectional analysis, we are also unable to report longitudinal change in PFT.

We show that significant radiographic ILD is present in patients with SSc, especially in those with positive Scl-70 autoantibodies, despite a “normal” PFT. We conclude that clinicians should maintain a high index of suspicion for SSc-ILD despite normal PFT results, and should consider performing a screening chest HRCT examination. However, we recognize that the amount of radiation from 1 chest HRCT

(~3–27 mSv) can exceed the equivalent dose of over 100 chest radiographs (~0.06–0.25 mSv)<sup>14</sup>. Additional studies that examine longitudinal change in PFT over time and the presence of radiographic ILD may help to shed light on how clinicians may best monitor for SSc-ILD while avoiding unnecessary radiation exposure.

## ONLINE SUPPLEMENT

Supplementary material accompanies the online version of this article.

## REFERENCES

1. Wells AU. Interstitial lung disease in systemic sclerosis. *Presse Med* 2014;43:329-43.
2. Meyer K. Diagnosis and management of interstitial lung disease. *Transl Respir Med* 2014;2:4.
3. Suliman YA, Dobrota R, Huscher D, Nguyen-Kim TD, Maurer B, Jordan S, et al. Brief report: pulmonary function tests: high rate of false-negative results in the early detection and screening of scleroderma-related interstitial lung disease. *Arthritis Rheum* 2015;67:3256-61.
4. Steele R, Hudson M, Lo E, Baron M; Canadian Scleroderma Research Group. Clinical decision rule to predict the presence of interstitial lung disease in systemic sclerosis. *Arthritis Care Res* 2012;64:519-524.
5. Goh NS, Desai SR, Veeraraghavan S, Hansell DM, Copley SJ, Maher TM, et al. Interstitial lung disease in systemic sclerosis: a simple staging system. *Am J Respir Crit Care Med* 2008; 177:1248-54.
6. van den Hoogen F, Khanna D, Fransen J, Johnson SR, Baron M, Tyndall A, et al. 2013 classification criteria for systemic sclerosis: an American College of Rheumatology/European League against Rheumatism collaborative initiative. *Arthritis Rheum* 2013; 65:2737-47.
7. Lammi MR, Baughman RP, Birring SS, Russell AM, Ryu JH, Scholand M, et al. Outcome measures for clinical trials in interstitial lung diseases. *Curr Respir Med Rev* 2015;11:163-74.
8. Kazerooni EA, Martinez FJ, Flint A, Jamadar DA, Gross BH, Spitznagel DL, et al. Thin-section CT obtained at 10-mm increments versus limited three-level thin-section CT for idiopathic pulmonary fibrosis: correlation with pathologic scoring. *AJR Am J Roentgenol* 1997;169:977-83.
9. Macintyre N, Crapo RO, Viegi G, Johnson DC, van der Grinten CP, Brusasco V, et al. Standardisation of the single-breath determination of carbon monoxide uptake in the lung. *Eur Respir J* 2005; 26:720-35.
10. Hankinson JL, Odencrantz JR, Fedan KB. Spirometric reference values from a sample of the general U.S. population. *Am J Respir Crit Care Med* 1999;159:179-87.
11. Greiner M, Pfeiffer D, Smith RD. Principles and practical application of the receiver-operating characteristic analysis for diagnostic tests. *Prev Vet Med* 2000;45:23-41.
12. Steen V. Predictors of end stage lung disease in systemic sclerosis. *Ann Rheum Dis* 2003;62:97-9.
13. Nihtyanova SI, Schreiber BE, Ong VH, Rosenberg D, Moynadeh P, Coghlan JG, et al. Prediction of pulmonary complications and long-term survival in systemic sclerosis. *Arthritis Rheumatol* 2014;66:1625-35.
14. Kalra MK, Maher MM, Rizzo S, Kanarek D, Shepard JA. Radiation exposure from chest CT: issues and strategies. *J Korean Med Sci* 2004;19:159-66.