

## Stabilization of Microcirculation in Patients with Early Systemic Sclerosis with Diffuse Skin Involvement following Rituximab Treatment: An Open-label Study

To the Editor:

Systemic sclerosis (SSc) is a multisystemic autoimmune disease characterized by fibrosis of the skin and internal organs, generalized microvasculopathy, and antibody response against various cellular antigens. Severe organ involvement occurs early in the course of diffuse cutaneous SSc (dcSSc) and has a bad prognosis<sup>1</sup>. Survival of the first years of the disease is associated with improved outcome. Therapies that may help the patient overcome this early period seem warranted<sup>2</sup>. Rituximab (RTX) has been reported as optional therapy in SSc<sup>3,4,5</sup>. Our group reported stabilization of internal organ involvement during a 2-year followup in an open pilot study of a 2-treatment course (months 0 and 6) of RTX in patients with early dcSSc<sup>6,7</sup>. In our pilot studies, modified Rodnan skin score (mRSS) decreased significantly after RTX course. The percent of decrease in the open pilot studies was corroborated by a similar decrease in the percentage of collagen score in blindly assessed histopathological skin analyses.

Because SSc is characterized by a pronounced microangiopathy over time (i.e., more severe loss of capillaries over time), it may be worthwhile to investigate whether treatment with RTX could also stabilize microangiopathy in dcSSc<sup>8,9</sup>. Nailfold videocapillaroscopy (NVC) is a reliable technique to evaluate the microcirculation (capillaries) in SSc<sup>10</sup>. Subsequent to the former studies, our additional study described herein, using the same design as the already published studies, is the first (to our knowledge) to assess microangiopathic evolution after a 2-treatment course (months 0 and 6) with RTX in early dcSSc<sup>6,7</sup>. Six consecutive patients with “early” (see below) dcSSc received an infusion of 2 times 1000 mg RTX at months 0 and 6, together with 100 mg methylprednisolone. Low-dose prednisolone ( $\leq 10$  mg/day) was allowed, provided that the patients were taking a stable dose at least 12 weeks before inclusion in the trial. All disease-modifying antirheumatic drugs [except methotrexate (MTX)] were stopped 12 weeks before screening<sup>6,7</sup>. Patients (5 men and 1 woman) were receiving a stable dose of MTX (10–25 mg/week) as background therapy since at least 12 weeks. Their median age was 49 years (range 37–64 yrs) and their median SSc disease duration was 11 months (range 5–22 mos, from onset of the first non-Raynaud phenomenon). Four of the 6 patients were anti-scl70-positive and 1/6 was RNA-polymerase-positive.

Nailfold capillaries were imaged using a nailfold videocapillaroscope with high magnification (200 $\times$ ) in 8 fingers (f), f2–5 of both hands, at months 0, 3, 6, and 12. The number of capillaries in the distal row were recorded, as well as the number of giants and hemorrhages, and the number of abnormally shaped [(neo)angiogenetic] capillaries. Scores were calculated, as described previously<sup>9</sup>. The observers evaluating the capillaroscopic images (CP, VR) were blinded to the study design and subjects' identity. Next to capillaroscopic assessment, clinical readouts (mRSS, lung function, and echocardiography) and Disease Activity Score (DAS) were done at 0, 3, 6, and 12 months<sup>6,7</sup>. Mixed-model analysis (MMA) with random intercept for patient was used to evaluate changes in variables over time. A statistical significance level of 0.05 was used.

There was a clinical significant change in skin score with a mean (SD) mRSS of 24.8 (5.95) at baseline and 10.2 (1.17) at Month 12 (MMA  $p < 0.001$ , mean/median % improvement: 59%/60%) and a significant decrease in DAS, with a mean (SD) of 4.2 (1.69) at baseline and 0.6 (0.74) at Month 12 (MMA  $p < 0.001$ , mean/median % improvement: 86%/87%). Indices of internal organ involvement remained stable (Table 1).

Semiquantitatively scored NVC variables remained stable, showing no progression of the microvascular damage during the 12-month followup (Table 2)<sup>9</sup>. More specifically, whereas more pronounced loss of capillaries was expected over time, the number of capillaries remained stable<sup>8</sup>. In this way, the mean score (SD) of capillary loss at baseline/12 months was 2.170 (0.408)/1.830 (0.408, MMA  $p = 0.341$ ). There was also no significant change in the number of other scleroderma type morphological characteristics (giants, hemorrhages, and neoangiogenesis) over time (Table 2). There was 1 serious adverse event, a scleroderma renal crisis. It occurred before the first RTX infusion and was considered probably unrelated to the study medication. It reacted well to angiotensin-converting enzyme inhibitors without need for renal replacement therapy, further bouts, or deterioration during the 12-month followup.

To our knowledge, ours is the first open pilot study to suggest that 2 immunosuppressive treatment courses with RTX, while receiving background stable MTX, may not only have potential efficacy for skin and stabilization of internal organ involvement, but also additional stabilization of microangiopathic variables in early dcSSc. Larger, randomized controlled trials are needed to further investigate these findings.

Table 1. Changes in clinical variables in patients with early and severe diffuse cutaneous systemic sclerosis treated with rituximab (n = 6). Data are mean (SD), median (IQR).

	0M	3M	p*	6M	p*	12M	p*	p MMA
mRSS, 0–51 points	24.8 (5.95) 25.0 (19.3–30.0)	18.6 (8.68) 15.0 (12.5–26.5)	0.026	13.8 (5.19) 13.5 (8.8–19.3)	< 0.001	10.2 (1.17) 10.0 (9.0–11.3)	< 0.001	< 0.001
DLCO, % of normal	70.2 (12.98) 68.0 (59.8–85.0)	61.0 (12.21) 61.0 (51.0–71.0)	0.105	66.5 (15.98) 67.0 (57.5–75.3)	0.213	69.3 (18.13) 69.5 (59.0–83.8)	0.771	0.302
FVC, % of normal	99.7 (13.77) 99.0 (88.8–113.3)	94.4 (11.08) 97.0 (84.0–103.5)	0.590	100.5 (17.74) 100.5 (86.0–116.0)	0.808	101.2 (13.85) 105.5 (91.0–111.5)	0.646	0.781
TLC, % of normal	87.5 (9.40) 87.0 (78.0–94.8)	83.8 (9.52) 81.0 (76.5–92.5)	0.970	88.7 (15.74) 85.0 (78.3–103.8)	0.740	94.5 (14.6) 94.0 (83.8–104.5)	0.062	0.191
FEV, % of normal	92.5 (10.15) 90.0 (83.5–104.3)	91.0 (12.02) 95.0 (78.5–101.5)	0.447	91.8 (12.29) 95.0 (79.3–101.0)	0.796	97.3 (11.76) 98.0 (87.5–107.0)	0.077	0.088
DAS, 0–10 points	4.2 (1.69) 3.8 (3.1–5.5)	1.8 (1.15) 2.0 (0.8–2.8)	< 0.001	1.1 (0.58) 1.0 (0.5–1.6)	< 0.001	0.6 (0.74) 0.5 (0.0–0.9)	< 0.001	< 0.001
sPAP, mmHg	30.5 (2.65) 31.0 (27.8–32.8)	30.4 (6.19) 28.0 (25.5–36.5)	0.581	33.0 (8.25) 32.0 (25.5–41.0)	0.734	30.4 (4.45) 29.0 (26.5–35.0)	0.430	0.603
LVEF, % of normal	58.5 (8.10) 57.5 (51.3–66.8)	65.8 (7.82) 69.0 (58.0–72.0)	0.094	59.5 (4.93) 59.0 (55.0–63.3)	0.830	63.5 (5.72) 64.0 (58.8–69.0)	0.239	0.238

\* Significance of all values versus baseline. M: month; MMA: mixed-model analysis; mRSS: modified Rodnan skin score; FVC: forced vital capacity; TLC: lung total capacity; FEV: forced expiratory volume; DAS: Disease Activity Score; sPAP: systolic pulmonary artery pressure; LVEF: left ventricular ejection fraction; IQR: interquartile range.

Table 2. Microangiopathic evolution (SQ scores) in patients with early and severe diffuse cutaneous systemic sclerosis treated with rituximab (n = 6). Data are mean (SD), median (IQR).

	0M	3M	p*	6M	p*	12M	p*	p MMA
SQ capillary loss	2.17 (0.408) 2.0 (2.0–2.3)	2.20 (0.447) 2.0 (2.0–2.5)	0.915	2.17 (0.753) 2.0 (1.8–3.0)	1.000	1.83 (0.408) 2.0 (1.8–2.0)	0.147	0.341
SQ giants	0.67 (0.516) 1.0 (0.0–1.0)	1.00 (0.000) 1.0 (0.0–1.0)	0.122	1.00 (0.000) 1.0 (0.0–1.0)	0.090	1.17 (0.408) 1.0 (1.0–1.3)	0.016	0.093
SQ hemorrhages	0.67 (0.516) 1.0 (0.0–1.0)	0.80 (0.447) 1.0 (0.5–1.0)	0.590	0.83 (0.408) 1.0 (0.8–1.0)	0.463	1.00 (0.000) 1.0 (0.0–1.0)	0.154	0.529
SQ neoangiogenesis	0.83 (0.408) 1.0 (0.8–1.0)	1.20 (1.095) 1.00 (0.5–2.0)	0.132	1.00 (0.632) 1.00 (0.8–1.3)	0.573	0.83 (0.753) 1.00 (0.0–1.3)	1.000	0.383

\* P value versus baseline. SQ: semiquantitative; M: month; MMA: mixed-model analysis; IQR: interquartile range.

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