Conclusion: a significant improvement in endothelial function and microvascular involvement was achieved after three months of HCQ treatment. It is a novel finding in SSc which can represent a new therapeutic possibility for the prevention of microvascular complications of the disease.

REFERENCES:


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DIFFERENTIAL PERFORMANCE OF NAILFOLD VIDEO CAPILLAROSCOPIC PARAMETERS IN THE DIAGNOSIS AND PROGNOSIS OF SYSTEMIC SCLEROSIS
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Background: The role of Nailfold Video Capillaroscopy (NVC) in the identification of patients with Raynaud phenomenon (RP) at risk to develop systemic sclerosis (SSc) is well established. However, it is not clear if certain capillaroscopic indices perform better than others at predicting SSc development and which NVC parameters have a prognostic value in established SSc.

Objectives: To comparatively assess the performance of different NVC parameters in predicting development of SSc, very early diagnosis of SSc (VEDOSS), or mixed connective tissue disease (MCTD) in patients with RP. Also, to longitudinally examine the consistency of clinical correlations of NVC parameters in SSc patients at two different time points and evaluate the prognostic capacity of NVC in SSc.

Methods: Consecutive RP patients referred to our department for NVC (138 with SSc, 12 with VEDOSS, 6 with MCTD, 36 with primary RP, and 50 with non-SSc secondary RP) were evaluated at baseline, both clinically and capillaroscopically; 175 were reevaluated after a mean±SD of 3.3±1.48 years. Sixty-two healthy volunteers served as controls. Qualitative assessment of NVC images permitted categorization of patients to a normal, early, active or late capillaroscopic pattern. Capillary loss, dilated, giant or ramified capillaries and micro-hemorrhages were evaluated by a scoring system (range 0 to 4 points) based on the mean of the 1mm fields of each of the 2nd, 3rd, 4th and 5th finger of both hands. FVC and DLOC were recorded, if performed within 6 months of NVC. FVC and DLOC deterioration were considered clinically significant if >10% and >15%, respectively. Skin thickening was measured using the modified Rodman Skin Score (mRSS). MRSS deterioration was considered clinically significant if >3.5. First occurrence of digital ulcers in patients with no prior such history and vital status were also recorded at follow-up.

Results: Capillary loss score had the highest diagnostic accuracy at discriminating patients with an SSc-spectrum disorder from patients with RP of different etiology and from controls, as defined by ROC curve analysis [AUC (95% CI)=0.925 (0.893-0.956)], followed by dilatation score [AUC (95% CI)=0.904 (0.807-0.938)] and giant score [AUC (95% CI)=0.856 (0.810-0.902)]. By contrast, micro-hemorrhages [AUC (95% CI)=0.722 (0.669-0.786)] and ramification scores [AUC (95% CI)=0.588 (0.521-0.654)] did not perform equally well. Notably, clinical correlations of capillaroscopic parameters in SSc were less likely to be consistent over time, when longitudinally assessed at two different time points by univariate and multivariate analysis. Binary logistic regression analysis indicated that baseline capillaroscopic pattern could predict occurrence of a combined adverse disease outcome (FVC deterioration>10% and/or DLOC deterioration>15% and/or mRSS deterioration>3.5 and/or first occurrence of digital ulceration and/or death), after a mean±SD follow up of 3.29±1.45 years in 94 SSc patients with available follow-up data (OR=3.43, p=0.031 for active versus early pattern, OR=8.778, p=0.007 for late versus early pattern).

Conclusion: Dilatation score performs best of all semi-quantitative NVC parameters in diagnosing SSc and although clinical correlations of capillaroscopic findings change over time, an active or late capillaroscopy pattern at baseline is associated with an adverse prognosis.


SENSITIVITY TO CHANGE AND RESPONSIVENESS TO TREATMENT OF RENAL RESISTIVE INDEX (IR) IN SYSTEMIC SCLEROSIS (SSC)
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Background: RRI may identify any problem with the blood flow in the renal artery and in the parenchyma. It was shown to be increased in SSc patients and being in relationship with both vascular and fibrotic SSc-related manifestations.

Objectives: to test sensitivity to change and responsiveness to treatment in RRI in SSc.

Methods: patients fulfilling the 2013 ACR/EULAR classification criteria for SSc were enrolled from two SSc-care units, if RRI was determined at least twice since being diagnosed. Data regarding SSc clinical manifestations, instrumental and laboratory evaluation for renal, cardiac and cardio-vascular involvement, as well as ongoing immunosuppressive and vasoactive/vasodilating treatment, were collected both at baseline and follow-up RRI measurements.

Results: 230 patients (aged 57 [48-67] years, 12.6% males) were enrolled in the study, with baseline RRI value of 0.68 (IQR 0.70). In a mean 3.4 years follow-up, RRI showed a median change (ΔRRI) of +0.02 (IQR 0.05). ΔRRI was significantly correlated with ΔsPAP (R=0.173, p=0.023) and it was significantly higher in patient with new onset of pulmonary arterial hypertension (0.08 ± 0.02 vs 0.03 ± 0.05; p=0.038). On Cox univariate regression analysis, time from disease onset, ΔsPAP and ΔRRI predicted new PAH, while ΔsPAP was the only independent predictors at multivariate regression analysis. Regarding treatment, Sildenafil exposure determined a significantly lower increase in ΔRRI, with a progressive long-term effect. Conversely, CCBs and iloprost treated patients showed a significantly higher increase of ΔRRI, which was determined by a higher DU burden. No difference was seen when immunosuppressive treatment was evaluated.

Conclusion: RRI is sensitive to change and reflects, in particular, the worsening of cardio-pulmonary vascular involvement (increase in ΔsPAP and new PAH onset). Moreover, it shows a possible protective effect of Sildenafil in reducing pulmonary vasculature manifestations.

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