traditional risk factors (smoke, hypertension, dyslipidemia) and treatment with aspirin and hydroxychloroquine.

**Conclusion** Our results confirmed that Italian lupus patients suffer a high incidence of CV disease compared with general population. However, this incidence was lower than that detected in North European and American lupus cohorts significantly lower capillary density (7.97 [7.19; 8.72] vs. 8.92 (8.19; 9.34), p<0.05). Dilatation point and giant capillary point was significantly higher in the RP-SLE subgroup (0.36 [0.13; 0.69] vs 0.13 [0.06; 0.28] p<0.05, 0.06 [0.00;0.28 vs. 0.00 [0.00; 0.00] p<0.001).

**Conclusion** SSc capillary pattern is present in SLE as well, most of these particular patients had Raynaud’s phenomenon. Patients having both SLE and RP have lower capillary density and worse capillary structure. SLE patients capillary density is higher than the density found in SSc controls.

**Results** was 1.97 (1.19; 3.13) in SSc subgroup. The median microangiopathia evaluation score (MES) in this group, the median capillary density was 6.66 (5.2; 7.94) in this study.

**Methods** 318 systemic autoimmune patients and 25 healthy controls were collected, 73 fulfilled SLE classification criteria. Patients with and without RP were compared. 89 patients fulfilled SSc classification criteria, the median capillary density was 6.66 (5.2; 7.94) in this group, the median microangiopathy evaluation score (MES) was 1.97 (1.19; 3.13) in SSc subgroup.

**Results** 23 patients had pure ‘idiopathic’ SLE, 36 fulfilled SLE plus another classification criteria, 11 SLE plus two other, 2 SLE plus three other and 1 SLE plus four other. Median capillary density was 8.23 (7.4; 8.94), the median MES was 1.00 (0.56; 1.47); the median giant capillary number was 0.00 (0.00; 0.75) in the entire SLE group. 6.9% of all SLE patients had SSc early pattern, 1.4% SSc active pattern, 20.6% had SSc late pattern and 71.2% had no SSc pattern. Among patients having SSc pattern all except two had RP. Comparison of capillaroscopy of SLE patients with and without RP showed that patients in the former group had worse capillary structure.

**Background** Capillaroscopy is a noninvasive method for evaluating nailfold abnormalities and differentiating between primary and secondary Raynaud syndrome (RP). It is widely investigated in systemic sclerosis (SSc) but not in systemic lupus erythematosus (SLE). SSc pattern is described with decreased capillary density, haemorrhage, neoangiogenesis and avascularity.

**Objective** Evaluate capillaroscopic pattern and clinical features in SLE patients; examine the influence of RP on capillaroscopic pattern and capillary density.

**Methods** 318 systemic autoimmune patients and 25 healthy controls were collected, 73 fulfilled SLE classification criteria. All patients underwent detailed nailfold capillaroscopic investigation. Density, intercapillary distance was recorded as well as the progression and diagnostic parameters described by Cutolo in semiquantitative manner. Presence of RP was investigated by a detailed questionnaire. Patients with and without RP were compared. 89 patients fulfilled SSc classification criteria, the median capillary density was 6.66 (5.2; 7.94) in this group, the median microangiopathy evaluation score (MES) was 1.97 (1.19; 3.13) in SSc subgroup.

**Results** 23 patients had pure ‘idiopathic’ SLE, 36 fulfilled SLE plus another classification criteria, 11 SLE plus two other, 2 SLE plus three other and 1 SLE plus four other. Median capillary density was 8.23 (7.4; 8.94), the median MES was 1.00 (0.56; 1.47); the median giant capillary number was 0.00 (0.00; 0.75) in the entire SLE group. 6.9% of all SLE patients had SSc early pattern, 1.4% SSc active pattern, 20.6% had SSc late pattern and 71.2% had no SSc pattern. Among patients having SSc pattern all except two had RP. Comparison of capillaroscopy of SLE patients with and without RP showed that patients in the former group had worse capillary structure.

**Conclusion** LN in Hispanic SLE patients represents an early and severe manifestation with higher incidence. It imposes poorer prognosis during first years of disease duration.

**Background** Lupus nephritis among Hispanic SLE patients have been identified with poor outcomes when it is compared to other populations; so, we aimed to identify lupus nephritis characteristics and its outcomes in an inception cohort of Hispanic SLE patients. Patients and methods two-hundred twenty-three patients with SLE of recent-onset were studied. At baseline, standardized medical history and laboratory tests were done; follow-up visits occurred quarterly, and information about renal disorder, disease activity, damage accrual and comorbidities was updated annually. Main outcome was the development of renal disorder since SLE diagnosis, incidence of LN and ESRD over time, and mortality associated with renal disease.

**Results** At entry into the cohort, age of SLE patients [mean (SD)] was 27.3 (9.1) years and 90% were female. One-hundred thirty-one (59%) patients developed lupus nephritis during 9.95 years of follow-up; incidence-rate 59/1000 py, most events (78%) were developed within the first year of diagnosis. Patients with lupus nephritis had lower baseline BMI, less frequency of arthritis, and higher hypertension. There were no differences on age at lupus diagnosis, gender and baseline comorbidities between lupus patients with and without renal involvement. Among patients with renal biopsy, 80% had ISN/RPS Class IV and V alone or in combination. Twenty-eight (21%) developed ESRD, five of them (18%) have been received renal transplantation. Severe renal disease was strongly associated with poor outcomes in this cohort.

**Conclusion** LN in Hispanic SLE patients represents an early and severe manifestation with higher incidence. It imposes poorer prognosis during first years of disease duration.