Conclusions: TUS differed between study arms, favouring RTX. This encourages further research into SGUS as an imaging biomarker in PSS clinical trials.

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AB0445 HYDROXYCHLOROQUINE'S IMPACT IN RENAL BIOPSY AND OUTCOMES OF LUPUS NEPHRITIS

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Background: Given its immunomodulatory effects, hydroxychloroquine use is recommended in systemic lupus erythematosus (SLE). It is associated with a lower rate of appearance and of relapse of lupus nephritis (LN). LN is classically classified using ISN/RPS classification, but others indexes, such as the ones described by Austin and Hill, allow for the quantification of SLE activity in the kidney tissue.

Objectives: To analyze the association between the use of hydroxychloroquine and the activity of LN in the kidney biopsy.

Methods: Retrospective single center study of consecutive SLE and biopsy proven LN patients, diagnosed from 2010 to 2016. We evaluated the following outcomes: clinical remission, renal function and proteinuria at end of follow-up (g/24h). Complete remission was defined as a reduction of proteinuria to <0.5g/24h, inactive urinary sediment and serum creatinine <115% of baseline; partial remission same parameters, except proteinuria <1g/24h if initial value <3h/24h, or reduction to <3g/24h if initial value >3g/24h. Kidney biopsies were evaluated by the INS/RPS LN classification and the morphological indexes described by Austin and Hill, obtained after histomorphological review of renal biopsies. The studied predictor was the use of hydroxychloroquine. Statistical analysis was performed with STATA software, using one-way ANOVA, Qui2 and Pearson/Sperman test were appropriate.

Results: During 6 years, there were 46 biopsy-proven LN cases, 84,8% (n=39) woman, median 35 years old (27–42,5) and 57,6% (n=19) caucasian. 39 patients were already known to have SLE, 7,44 (1,13–12,3) years previously. Of those 39 patients, 46% were under hydroxychloroquine and 77% under other immunosuppression.

The median follow-up was 31,9 (13,2–45,6) months. Based on biopsy findings, 35 patients were started on immunosuppression – induction in 50% of cases with MMF and in 50% with cyclophosphamide; maintenance in 81% with MMF, the rest with azathioprine. Complete remission was achieved in 58% of patients, 27% achieving partial remission. We observed 4 LN relapses. At the end of FUP, we saw a 96% (n=44) patient survival, with a median serum creatinine of 0,8 mg/dl (0,7–0,99), eGFR 99,8 ml/min (71,2–116,8) and proteinuria of 0,6 g/24h (0,2–1,6). From those 46 patients, 30 were under immunosuppressive therapy at time of LN presentation, and 60% (n=18) were also under hydroxychloroquine. Table 1 summarizes the clinical findings:

With the use of hydroxychloroquine, we observed a lower histomorphological activity, as represented by a lower Hill biopsy index, and tendency towards lower Activity index. We also saw a tendency towards lower proteinuria.

Conclusions: Our data reinforces the recommendations of using hydroxychloroquine for its adjuvant role in SLE patients, as we saw a lower histomorphological activity in kidney biopsy, and a trend towards lower proteinuria.

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AB0446 ADDITIVE INHIBITION OF INTERFERONS, B AND T CELL ACTIVATION AND TFH-RELATED CYTOKINE CXCL13 BY LEFLUNOMIDE AND HYDROXYCHLOROQUINE SUPPORTS RATIONALE FOR COMBINATION THERAPY IN PSS PATIENTS

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Abstract AB0445 – Table 1

Hydroxychloroquine use	Class II	Class III	Class IV	Class V	Hill Biopsy Index	Activity Index	Chronicity Index	End of FUP eGFR (ml/min)	End of FUP serum creatinine (mg/dl)	End of FUP proteinuria (g/24h)
Yes (n=18)	39%	28%	28%	22%	0,86 (0,2 – 1,8)	1 (0 – 4,25)	0,75 (0 – 3,25)	99,8 (81,9 – 112,1)	0,74 (0,7 – 0,8)	0,3 (0,1 – 0,6)
No (n=12)	8%	17%	58%	33%	1,52 (0,98 – 2,01)	7,25 (1,38 – 9)	2,25 (0,13 – 5,25)	98,6 (56,4 – 120,3)	0,78 (0,7 – 1,3)	0,75 (0,15 – 2,6)
p-value	0,09	NS	NS	NS	0,03	0,07	NS	NS	NS	0,09

Background: T and B cell-driven immunity is critically involved in immunopathology of pSS. Recently we demonstrated synergistic T and B-cell activation upon T cell triggering and TLR7/9-driven B cell activation in pSS patients, accompanied by synergistic induction of immunoglobulins and IFN- γ - and IL-17-producing T cells¹. In addition, TLR7/9-expressing activated pDCs associated with increased type I IFNs and IFN-inducible genes are increased pSS patients. Several studies have shown that the DMARDs leflunomide and hydroxychloroquine inhibit immune activation in pSS but only show moderate efficacy. However, LEF and HCQ target different pathways with overlapping, but also potentially additive mechanisms, where LEF primarily targets T and B cells and HCQ TLR7/9-driven B cell and pDC activation.

Objectives: To assess the additive effects of LEF and HCQ on CD4 T- and B-cell activation and production of interferons IFN- α and IFN- γ , Tfh-related cytokine CXCL13, as well as IgG and IgM *in vitro* employing SEB/TLR9-triggered PBMC.

Methods: PBMCs of healthy individuals (n=9) and of pSS patients (n=8) were cultured with antigen (SEB), TLR9 and their combination, in presence or absence of LEF, HCQ and their combination in clinical relevant concentrations. Proliferation of T and B cells and release of IFN- α , IFN- γ , CXCL13, IgG and IgM were measured.

Results: In line with robust T and B cell activation, IFN- γ , IFN α , CXCL13, IgG and IgM production was achieved by a combination of SEB and TLR9 (all at least p<0.001). LEF dose dependently inhibited B and T cell proliferation, Interferon, CXCL13 and immunoglobulin production. HCQ dose dependently inhibited B cell proliferation, IFN- α , CXCL13, and immunoglobulin production. T cell proliferation and IFN- γ production were inhibited by HCQ only at higher concentrations. At several suboptimal concentrations LEF and HCQ additively inhibited T cell proliferation both in healthy individuals and in pSS patients. (Figure 1). Significant additive effects were seen for all outcome measures except IFN- α . Since IFN α was already robustly inhibited by HCQ alone (eg. for pSS 90% at 3.3 μ M, p<0.001), only trends towards additive effects were observed.

Figure 1. HCQ and LEF additively inhibit T cell activation. T cell proliferation is dose dependently inhibited by LEF and by HCQ at 0.1 and 10 μ M (A). Significant additive inhibition of T cell proliferation is achieved by combination of suboptimal concentrations of LEF and HCQ both in HC and pSS patients (B). *and **** indicates statistical significance of p<0.05 and p<0.0001 resp. vs control, # indicates statistical significance of p<0.05 of the combination vs each of the single drugs.

Conclusions: LEF and HCQ robustly inhibited proliferation of T and B cells, cytokine production and immunoglobulin production with clear additive efficacy in both healthy individuals as in pSS patients. These data support the potential surplus value of combination therapy with LEF and HCQ for patients with pSS.

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AB0447 ANTIMALARIALS IMPROVE SURVIVAL OF SYSTEMIC LUPUS ERYTHEMATOSUS ON CHOLESTEROL: RESULTS OF A FIFTEEN-YEAR CHINESE MULTICENTER RETROSPECTIVE STUDY IN JIANGSU PROVINCE

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Background: Nowadays the importance of antimalarials, especially hydroxychloroquine (HCQ) and chloroquine (CQ), in treatment of systemic lupus erythematosus (SLE) has been demonstrated. However, few have examined the efficacy of HCQ and CQ on eastern Chinese SLE patients.

Methods: The analysis is based on 1372 patients who were enrolled in a retrospective study of 26 centers from January 1st, 1999 through December 31st, 2009, during which time is their first hospitalization. Baseline and follow-up clinical, laboratory and therapeutic data and survival status before April 30th, 2015 were recorded. Statistical analysis consist of Chi-square test, *t*-test, Kaplan-Meier curves and logrank test.

Results: Compared with 562 patients without HCQ or CQ treatment, the hazard ratio (HR) of deaths in 810 patients taking those was reduced (HR 0.52, 95% CI 0.38–0.70, *p*<0.001). 376 of these 1372 patients experienced their second hospitalization, during which treating group (165 of 376) showed lower blood level of total cholesterol (TG), compared to control group (4.47 (0.13) vs 5.03 (0.21), *p*=0.027), while no statistical difference of TG exists between the two groups' first hospitalization (*p*>0.05). Other metabolic data, such as systolic and diastolic blood pressure, fasting blood sugar, Triglyceride and uric acid were similar between the two groups in two times of hospitalization. On second inpatient visit, disease activity (SLEDAI, blood sedimentation rate, complement) and organ involvements (SLICC) of those who took antimalarials and no users have no significant differences.

Conclusions: Use of HCQ or CQ lower the risk of mortality and TG levels of eastern Chinese SLE patients.

Disclosure of Interest: None declared

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AB0448 USE OF DENOSUMAB IN PATIENTS WITH SYSTEMIC LUPUS ERYTHEMATOSUS: A REAL LIFE MULTI-CENTRIC EXPERIENCE

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Background: The occurrence of osteoporosis (OP) and fragility fractures (FF) is a frequent comorbidity in patients with systemic lupus erythematosus (SLE), mainly due to the concomitant presence of many risks factors for secondary osteoporosis (drugs, gender, concomitant diseases). The use of corticosteroids (CS) increases the risk of osteoporosis induced by corticosteroid (GIOP) and FF [1]. Denosumab is monoclonal antibody that binds to RANKL, inhibiting osteoclast formation and activation, used for both male and women OP to prevent FFx [2,3]. Previous RCT [2] reported a numerically higher serious adverse events of infections in the denosumab users. In particular severe skin infections were significantly more frequent [4]. The possible increase in infection risk might be a concern in subjects treated with CS. The use of denosumab in SLE with GIOP patients is recently reported in 17 cases within a RCT [5]. Anyway, no data regarding the tolerability and disease activity during therapy were reported

Objectives: Our aim was to analyze the prevalence, the tolerability, and relevant changes in disease activity or damage in a cohort of SLE patients treated with denosumab

Methods: among all SLE patients currently followed in two referral Center we selected the ones that received at least one dose of denosumab. Clinical, serological manifestations, concomitant diseases and therapies were collected from clinical charts. Damage index (SDI), activity index (SLEDAI-2K) were calculated when denosumab was introduced (first cycle) and at the last evaluation (last cycle). For statistical analysis Chi-squared test or Fisher exact test was used

Results: Among 793 patients in our cohorts, 21 (2.6%) were treated with denosumab. Demographic data reported a female prevalence (19 cases, 90%), mean age at onset of 38±12years, mean duration of disease of 28±10years and mean follow-up of 21±7.8years; most frequent manifestations were arthritis (90%), cutaneous (67%) and neurological (67%). All patients were still treated with CS (mean duration of 25±6.8 years) with a daily dose of 8.5±3.5mg/day. Other risk factors for OP were present in the majority of them: drugs (anticoagulants

Efficacy/adverse events	First cycle (N=21 pts)	Last cycle (N=21 pts)	P value
New fracture	Not applicable	0 (0%)	na
Mean dose of GC, mg/day (mean±DS)	6.9±3.4	5.8±3.1	ns
T score hip*	-2.6±1.4	-2±1	ns
T score lumbar spine *	-2.6±1.1	-2.3±0.8	ns
Hypovitaminosis D **	15 (71%)	13 (62%)	ns
SLEDAI-2k (mean±DS)	3.5±3.3	2.36±2.65	ns
SDI (mean±DS)	4 ±1.18	4.1±1.21	ns
Any adverse event	1	1	ns
Infective episodes	1 (not requiring antibiotic)	1 (not requiring antibiotic)	
Skin infection	0	0	ns
Flare SLE	0	1 (arthritis)	ns
Therapy withdrawal	na	0	ns

*available for 18 patients at first cycle and 13 at last evaluation, **=all patients received vitamin D supplementation; na=not applicable; ns=not significant

in 6pts, cyclosporin in 3 and antiepileptic in 2), chronic kidney disease (CKD) (4pts), hypovitaminosis D (15pts), anorexia, celiac disease and hemochromatosis (1pt, each). Indication for denosumab was OP with FFx in 17 cases: denosumab was used for a new FF during biphosphonates or after teriparatide. In the other 4 the indication was primary prevention with contraindications at the use of biphosphonates for concomitant severe CKD. Mean duration therapy was 4 cycles (range 1–8): data regarding disease activity, damage and adverse events are reported in table 1. No new FFx developed in any of the included patients at the last evaluation available.

Conclusions: In our cohorts denosumab is still used in few selected patients. However it could be considered as a valid option in GIOP patients because it was globally well tolerated and in our cohort it's efficacy in the prevention of new fragility fractures was confirmed

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AB0449 OFF LABEL OF BIOLOGICS IN CONNECTIVE TISSUE DISEASES. A SINGLE CENTER EXPERIENCE

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Background: Connective tissue diseases (CTDs) are a broad spectrum of autoimmune conditions including different entities such as systemic sclerosis (SSc), Sjogren syndrome (SSj), systemic lupus erythematosus (SLE) and autoimmune myositis. There are many patients that do not meet the classification or diagnostic criteria and they fall under the diagnosis of Undifferentiated connective tissue disease (UCTD). Treatment of refractory forms of these could be challenging and clinicians sometimes are forced to try off label drugs such as biologics on the basis of scarce literature support and the experience on different rheumatic disease.

Objectives: To evaluate duration of biologic drug administered off label for the treatment of CTD and investigate variables possibly associated to drug suspension.

Methods: We used ACE program to search among Mayo Clinic clinical records all the patients that had in their records the words "undifferentiated connective tissue disease" or "lupus" or "myositis" or "Sjogren" or "systemic sclerosis" or "scleroderma"(and acronyms) AND "infliximab" or "etanercept" or "golimumab" or "certolizumab" or "tocilizumab" or "abatacept" or "adalimumab" (and their brand names).

All records were checked for definite diagnosis and patients with uncertain ones where excluded. Also medications were checked, patients without any information about treatments where excluded. All the records of the selected patients were used to collect information about the off label biologic treatment but also clinical, serological and demographic variables.

Results: We collected data on 122 patients with connective tissue diseases, some of them had other concomitant autoimmune diseases with indication for biologic treatment (e.g. rheumatoid arthritis, etc). We analyze the group with some CTD alone (n=72) considering SLE (n=18 – 25%), inflammatory myositis (n=22 – 31%) and UCTD (n=32 – 44%). In this group the first biologic was etanercept in 55%, adalimumab in 18%, infliximab in 17%, golimumab in 1% and abatacept in 8%. The 22% tried also a second biologic and 4% a third. We consider for analysis the first treatment.

Mean treatment duration was 0.8 (±1.1)years. In our population 12,5% experienced a flare of CTD, 12,5% had infections, 18% had allergic reactions (of any type). The 44% experienced primary failure, 11% loss of efficacy, 31% had minor adverse events, 14% major ones (possible more than one reason for interruption).

Analysis showed no definite factors correlated with treatment duration or failure or adverse events, No difference due to type of CTD.

Conclusions: The study is retrospective and this limits the conclusions to be taken from it, however is one of largest population of off label treatments in CTDs