

Incomes	N	Mean	Std. Deviation	Median	Minimum	Maximum
<b>Medications</b>						
<b>Steroids</b>						
No	91	43955	10357	40565	17946	67091
Yes	12	39702	8927	39643	32285	63629
<b>Hydroxychloroquine</b>						
No	11	40530	7320	40159	31013	52924
Yes	92	43810	10525	40159	17946	67091
<b>Azathioprine</b>						
No	85	43303	10028	40159	17946	67091
Yes	18	44199	11541	41050	31977	66374
<b>Cyclophosphamide</b>						
No	102	43454	10304	40159	17946	67091
Yes	1	44010		44010	44010	44010
<b>Methotrexate</b>						
No	95	43282	10351	40159	17946	67091
Yes	8	45567	9358	41625	38887	66374
<b>Mycophenolate</b>						
No	86	43809	10774	40362	17946	67091
Yes	17	41689	7035	39808	32566	51055
<b>Belimumab</b>						
No	100	43369	10380	40159	17946	67091
Yes	3	46468	3779	44574	44010	50819
<b>Tacrolimus</b>						
No	100	43709	10284	40362	17946	67091
Yes	3	35124	4361	32646	32566	40159
<b>Any of the above*</b>						
No	4	40431	9947	38893	31013	52924
Yes	99	43582	10296	40159	17946	67091

\*p-value=0.592 (Mann-Whitney U Test)

#### 400 ANA-NEGATIVE SLE: RE-EVALUATION IN AN INTERNATIONAL INCEPTION COHORT

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**Background and aims** The prevalence of ANA-negative SLE is reportedly 5%–20%. Cytoplasmic or mitotic cell indirect immunofluorescence (IIF) patterns are usually reported as ANA-negative. This study examined the prevalence of ANA-negativity (no intracellular IIF pattern) and pure cytoplasmic and/or mitotic IIF patterns (CMP) in the Systemic Lupus International Collaborating Clinics (SLICC) inception cohort and examined demographic, clinical and autoantibody associations.

**Methods** Three groups were examined 1) ANA-positive (presence of nuclear IIF pattern), 2) ANA-negative (no IIF pattern), and 3) pure CMP. ANA were detected by IIF on HEp-2000 substrate, SLE-related autoantibodies by laser bead immunoassay, and anti-dsDNA and anti-dense fine speckles 70 (DFS70) by chemiluminescence immunoassay.

**Results** 1137 patients were included; 89.9% were female. 92.3% were ANA-positive, 6.2% were ANA-negative, and 1.5% had a CMP. In the multivariate analysis (Tables 1 and 2), patients from Canada (Odds Ratio (OR) 2.07 [95% CI: 1.28, 3.36]) or with anti-DFS70 (OR 4.45 [95% CI: 1.37, 14.39]) were more likely to be ANA-negative or have CMP. Patients of Asian descent (OR 0.34 [95% CI: 0.13, 0.86]) or with anti-dsDNA (OR 0.53 [95% CI: 0.30, 0.94]), anti-SSA/

Ro60 (OR 0.51 [95% CI: 0.30, 0.87]), or anti-UI-RNP (OR 0.35 [95% CI: 0.17, 0.70]) were less likely to be ANA-negative or CMP.

**Conclusions** In newly diagnosed SLE, the prevalence of ANA-negativity was at the lower end (6.2%) of the range previously published and an additional 1.5% had a CMP pattern. The prevalence of true ANA-negativity will likely decrease as future guidelines are expected to recommend that non-nuclear patterns, such as CMP, are also reported.

#### 401 HIGHEST FREQUENCY OF CLINIC VISITS AND HOSPITALIZATIONS IN SLE AMONG RHEUMATIC DISEASES: 8 YEAR CENSUS OF A TERTIARY RHEUMATOLOGY CENTRE

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**Background and aims** We describe the frequency of clinic visits and hospitalizations among rheumatic diseases seen at a tertiary Rheumatology centre in Manila, Philippines

**Methods** The University of Santo Tomas (UST) Hospital is a tertiary care centre, with specialised subspecialty training in Rheumatology. This study is derived from the patient census of UST Hospital Rheumatology Clinics from 2008 to 2015.

**Results** Mean age of the total 15 730 rheumatic disease patients (10 808, 69% females; 13 607, 86.5% adults; 2123, 13.5% paediatrics) was 47.51±21.55 (range <1–103). Most common rheumatic conditions were osteoarthritis (OA) (2828, 17.98%), gout/pseudogout (2378, 15.12%) and systemic lupus