

PO.7.155 STUDY OF THE COMORBIDITIES PRESENT IN THE PATIENTS INCLUDED IN THE REGISTRY OF SYSTEMIC LUPUS ERYTHEMATOSUS OF THE SPANISH SOCIETY OF RHEUMATOLOGY (RELESSER)

¹A Lois-Iglesias, ²R Iñigo, ³M Galindo Izquierdo, ⁴J Calvo-Alén, ¹V Balboa-Barreiro, ⁵C Mourriño, ⁶A Olivé, ⁷R Melero, ⁸A Fernández-Nebro, ⁸M Andrés, ²C Erasquin, ⁹E Tomero, ¹⁰C Fito, ¹¹E Uriarte, ¹M Freire, ¹²C Montilla, ¹³A Morasat, ¹⁴G Santos-Soler, ¹⁵A Boteanu, ¹⁶E León, ¹⁷J Narvaez, ¹⁸V Taboada, ¹L Silva, ¹⁹O Ibaranguoitia, ²⁰M Fernandez-Castro, ²¹A Hernandez-Beirian, ²²M Gantes, ²³B Hernández-Cruz, ²⁴J Pérez-Venegas, ²⁵A Pecondon, ²⁶N Lozano, ²⁷AP Cacheda, ²⁸G Bonilla, ²⁹V Torrente-Segarra, ³⁰I Castellvi, ³¹JJ Alegre, ³²J Calvet, ³³JL Marengo, ³⁴E Raya, ³⁵T Vázquez, ³⁶Q Víctor, ³⁷S Muñoz, ³⁸T Otón, ³⁹J Martínez Barrio, ⁵JM Pego-Reigosa*. ¹C.H.U.A. Coruña ~ A Coruña ~ Spain; ²H.U. Gran. Canaria Dr Negrín ~ Gran Canaria ~ Spain; ³H.U. 12 de Octubre ~ Madrid ~ Spain; ⁴H.U. Araba ~ Spain; ⁵H.U. Vigo ~ Spain; ⁶Germans Trias i Pujol ~ Barcelona ~ Spain; ⁷H.U. Málaga ~ Spain; ⁸C.H.U. Alicante ~ Alicante ~ Spain; ⁹H.U. Princesa ~ Madrid ~ Spain; ¹⁰H.U. Navarra ~ Spain; ¹¹H.U. Donosti ~ Spain; ¹²H.U. Salamanca ~ Spain; ¹³H.U. Príncipe de Asturias ~ Alcala de Henares ~ Spain; ¹⁴H.U. Marina Baixa ~ Alicante ~ Spain; ¹⁵H.U. Ramón y Cajal ~ Madrid ~ Spain; ¹⁶H.U. León ~ Spain; ¹⁷H.U. Bellvitge ~ Spain; ¹⁸H.U. Valdecilla ~ Santander ~ Spain; ¹⁹H.U. Basurto ~ Bilbao ~ Spain; ²⁰H.U. Puerta Hierro ~ Madrid ~ Spain; ²¹H. Insular Gran Canaria ~ Spain; ²²H.U. Canarias ~ Spain; ²³H.U. Virgen Macarena ~ Sevilla ~ Spain; ²⁴H.U. Jerez ~ Spain; ²⁵H.U. Miguel Servet ~ Zaragoza ~ Spain; ²⁶H.U. Virgen Arrixaca ~ Murcia ~ Spain; ²⁷H. Tenerife ~ Spain; ²⁸H.U. La Paz ~ Madrid ~ Spain; ²⁹CSAPG (Consorci Sanitari Alt Penedès Garraf) ~ Alt Penedes ~ Spain; ³⁰Alt Penedes ~ Spain; ³¹H.U. Peset ~ Valencia ~ Spain; ³²H.U. Parc Taulí ~ Sabadell ~ Spain; ³³H.U. Valme ~ Cádiz ~ Spain; ³⁴H.U. San Cecilio ~ Granada ~ Spain; ³⁵H.U. Ferrol ~ Spain; ³⁶H. Monforte ~ Spain; ³⁷H.U. Infanta Sofía ~ San Sebastian de los reyes ~ Spain; ³⁸H.U. Torrejón de Ardoz ~ Spain; ³⁹H.U. Grigorio Marañón ~ Madrid ~ Spain

10.1136/lupus-2022-elm2022.175

Background The survival of patients with systemic lupus erythematosus (SLE) has increased in recent years, but they have higher morbidity and mortality than the general population.

Purpose To study the prevalence of comorbidities in patients with SLE and its relationship with damage, gender and treatments received.

Methods Cross-sectional multicenter descriptive study of a cohort of adult patients with SLE.

Results We studied 3,656 patients, 90.3% women, mean age (\pm SD) at diagnosis of 35.2(\pm 14.7) years and duration of SLE of 142.6(\pm 100.8) months. We analyzed 27 comorbidities. 79.73% of the patients presented any, with the maximum accumulated being 14. The most frequent were smoking, dyslipidemia and arterial hypertension. 38.05% of patients accumulated damage. Males accumulated more comorbidities (85.48%

vs. 79.1%, $p=0.003$) and damage (47.03% vs. 37.11%, $p<0.001$). The first criterion for SLE appeared at a younger age in patients who did not have comorbidities: 27.73 (\pm 12.04) years vs. 34.47(\pm 14.76) years; $p<0.001$. We found that there is a positive correlation between the number of comorbidities and the number of systems with damage (Spearman's Rho = 0.478, $p<0.001$). There is a positive correlation between the number of comorbidities and damaged systems with the number of hospitalizations by disease activity (Rho=0.265 and 0.396 respectively, $p<0.001$ in both contrasts) as well as with the number of serious infections (Rho=0.299 and 0.307 respectively, $p<0.001$ in both contrasts). We found more patients without comorbidities in those who did not receive glucocorticoids (9.94% vs. 15.48%, $p<0.001$) and more patients with comorbidities in those who did not receive antimalarials (89.1% vs. 81.78%, $p<0.001$). There were significant differences in the presence of comorbidities in those treated with cyclophosphamide, mycophenolate, azathioprine or rituximab.

Conclusions A high percentage of patients with SLE have comorbidities. With few exceptions, they are more frequent in males. The onset of SLE was later in patients with more comorbidities. We found variations in comorbidities depending on the treatments received.

PO.7.156 DOES EXPERT OPINION MATCH THE DEFINITIONS OF LOW DISEASE ACTIVITY STATE? PROSPECTIVE ANALYSIS OF 500 PATIENTS FROM A SPANISH MULTICENTER COHORT

¹I Altabás González*, ²I Rúa-Figueroa, ²F Rubiño, ¹C Mourriño, ³R Menor Almagro, ⁴E Uriarte Isacelaya, ⁵E Tomer Muriel, ⁶TC Salman-Monte, ⁶I Carrión-Barberà, ⁷M Galindo, ⁷E Rodríguez-Almaraz, ⁸N Jiménez, ¹JM Pego-Reigosa. ¹Complejo Hospitalario Universitario de Vigo ~ Vigo ~ Spain; ²Hospital Universitario de Gran Canaria Doctor Negrín ~ Las Palmas de Gran Canaria ~ Spain; ³Hospital Universitario de Jerez de la Frontera ~ Cadiz ~ Spain; ⁴Hospital Universitario de Donostia ~ San Sebastián ~ Spain; ⁵Hospital Universitario de la Princesa ~ Madrid ~ Spain; ⁶Hospital del Mar ~ Barcelona ~ Spain; ⁷Hospital Universitario 12 de Octubre ~ Madrid ~ Spain; ⁸Grupo IRIDIS ~ Vigo ~ Spain

10.1136/lupus-2022-elm2022.176

OBJECTIVES To apply current definitions of Lupus Low Disease Activity State (LLDAS) to a large cohort and evaluate the

Abstract PO.7.156 Table 1 Reason of disagreement between patients that did not fulfill LLDAS definition and expert assessment as remission or low disease activity

Descriptors of disease activity	Not achievement of LLDAS Number (%) n=126 (100%)
1. SLEDAI-2 K \leq 4, with no activity in major organ systems (renal, CNS, cardiopulmonary, vasculitis, haemolytic anaemia, fever) and no gastrointestinal activity	59 (46.8%)
2. No new features of lupus disease activity compared to the previous assessment	74 (58.7%)
3. PGA (scale 0-3) \leq 1	4 (3.3%)
Immunosuppressive medications	
4. Current prednisolone (or equivalent) dose \leq 7.5 mg daily	16 (12.7%)
5. Well-tolerated standard maintenance doses of immunosuppressive drugs and approved biologic agents, excluding investigational drugs	

SLEDAI: systemic lupus erythematosus; CNS: central nervous system; PGA: physician global assessment