AB0550

CLINICAL-EPIDEMIOLOGICAL FEATURES OF PATIENTS WITH A LATE-ONSET LUPUS IN A TERTIARY CARE HOSPITAL

A. Crespo Golmar, C. Moriano, M. Garijo, C. Iñiguez, I. González, A. López, E. Díez, C. Álvarez, M. Martín, E. Vallejo, T. Pérez. *Rheumatology Department, University Health Care Complex of León, León, Spain*

Background: Different frequency of clinical and serological manifestations has been detected according to the age of onset of the patients with Systemic lupus erythematosus (SLE). According to the literature, senile SLE manifests between 6% and 18% of the patients with lupus.

Objectives:

1. To identify and analyse the clinical-serological and epidemiological features of senile SLE in our environment.
2. To determine the average survival time and mortality in these patients, identifying its main cause.

Methods: Observational retrospective study of 319 patients diagnosed with SLE (according to ACR 1992 and SLICC 2012 criteria) at the Hospital of León between 1997–2017 and with an age of onset ≥ 65 years, obtaining a total of 68 patients with senile SLE.

Results: The mean age at diagnosis was 75.4 ± 12.1 years, with a female/male ratio of 2.4. The most frequent manifestations were as joint (63.2%) and haematological manifestations in the form of leuco-lymphopenia (55.9%). The hemolytic anaemia only appeared in 2.9% of the cases and the thrombocytopenia in 25%. 36.8% of patients showed photosensitivity and 29% had other cutaneous manifestations, being the malar erythema the most prevalent type (60%), followed by the discoid lupus erythematosus (20%) and the subacute lupus (15%). Alopecia was only observed in 4.4%. Lupus nephritis was detected in the form of proteinuria in 4.4% of the patients, and only one patient had microscopic haematuria. Lung involvement was uncommon (8.8%), taking precedence the UIP (33.3%) over the rest of the pulmonary manifestations. Only 11.1% of the patients with senile SLE had serositis, being in the form of pleuritis in 75% of the cases, pericarditis in the 37.5% and ascites in the 12.5%. Regarding the neurological involvement, 5 patients showed polyneuropathy and 1 had chorea. Likewise, the frequency of Sjögren, Raynaud and secondary antiphospholipid syndrome was of 16.7%, respectively.

The most important serological findings were: 97.3% ANA; 44.1% DNA and 20.6% hypocomplementemia, with 54.4% of the patients having serological activity. Only 5.9% had anti-Sm. Antiphospholipid antibodies were positive in 41.2% of the cases, with 4.4% of them showing triple positivity.

The average survival time was of 13.7 years (SD: 10.9–16.5). Out of the total patients, 14 died (20.59%), mostly due to infectious etiology (35.7%) and 14.28% due to disease activity. Other less common causes were neoplasia or ischaemic heart disease (7.14% respectively).

Conclusions:

- The late-onset SLE prevails in our environment, one of every 5 patients diagnosed with SLE in our consulting room is older than 65 years.
- It is found most often in women and it is confirmed a lower male/female ratio than expected.
- Joint and haematology manifestations and cutaneous involvement in the form of malar erythema define the clinical profile of our patients with senile SLE, with the renal involvement or the presence of serositis being uncommon.
- Half of the patients had serological activity at the onset, having hypocomplementemia only in 1 out of 5 cases.
- Infections were the first cause of mortality in our sample with an average survival time of around 13 years.

Disclosure of Interest: None declared

DOI: 10.1136/annrheumdis-2018-eular.7538

AB0551

OBSTETRICAL MORBIDITY RELATED TO ANTI-SSA ANTIBODIES: DATA FROM A FRENCH MONOCENTRIC RETROSPECTIVE STUDY

L. Plaçais¹, F. Carrat², L. Carbillon³, P. Nicaise⁴, A. Benbara⁵, O. Fain¹, A. Mekinian¹. ¹Internal Medicine; ²Epidemiology and Public Health, Saint Antoine hospital; ³Obstetrics and Gynecology, Jean Verdier hospital; ⁴Auto-immunity and hypersensitivity, Bichat hospital; ⁵Obstetrics and Gynaecology, Jean Verdier hospital, Paris, France

Background: Obstetrical morbidity is an issue among autoimmune diseases patients. Anti-SSA positivity is well known for its effect on fetus and risk of

congenital heart block but no evidence has been yet found for its effect on obstetrical morbidity

Objectives: To investigate anti-SSA positivity effect among a large monocentric cohort of obstetrical morbidity patients and to evaluate the efficacy of different treatment regimens

Methods: All women who were seen from January 2010 to January 2015 in Jean-Verdier University Hospital for obstetrical morbidities were retrospectively included. All patients had been tested for anti-nuclear antibodies and anti-SSA/Ro and anti-SSB/La antibodies.

Results: We included 244 women (median age 34 years [interquartile range 21–53]) with 869 pregnancies overall. In 27 (11%) patients with anti-SSA antibodies, the median age at the time of pregnancy was 29.5 (17–40) years, with mean number of 3.66 pregnancies per woman. For these 27 patients, 83/99 pregnancies (84%) had an adverse obstetrical outcome: fetal loss (n=65; at a median of 20 (4–38) weeks' gestation), preeclampsia (n=15), intrauterine growth restriction (n=7), prematurity (n=7), and congenital heart block (n=2). In comparing women with obstetrical antiphospholipid syndrome and unexplained adverse complications, there was no major difference in frequency of recurrent miscarriages, preterm deliveries, preeclampsia or fetal loss. Among factors associated with obstetrical outcome and fetal loss, aspirin and hydroxychloroquine treatments were significantly associated with a favourable obstetrical outcome: odds ratio 0.05 [95% confidence interval 0.01; 0.37] (p=0.003) and 0.15 [0.02; 0.98] (p=0.04).

Conclusions: Women with unexplained recurrent obstetrical complications should be screened for anti-SSA antibodies. The benefit of aspirin and hydroxychloroquine treatment to improve the obstetrical outcome should be confirmed in prospective studies.

Disclosure of Interest: None declared

DOI: 10.1136/annrheumdis-2018-eular.5580

AB0552

THE CLINICAL AND LABORATORY FEATURES OF ANTIPHOSPHOLIPID ANTIBODY POSITIVE PATIENTS WITH OR WITHOUT SYSTEMIC LUPUS ERYTHEMATOSUS

A. Yazici¹, A. Omma², S. Yilmaz³, S. Yavuz⁴, K. Ayar⁵, E. Kasapoglu⁶, B. Kisacik⁷, A. Cefle¹, D. Erkan⁸. ¹Rheumatology, Kocaeli University School of Medicine, Kocaeli; ²Rheumatology, Numune Hospital; ³Rheumatology, Gulhane Training and Research Hospital, Ankara; ⁴Rheumatology, Bilim University School of Medicine, Istanbul; ⁵Rheumatology, University of Health Sciences Yüksek İhtisas Training and Research Hospital, Bursa; ⁶Rheumatology, Göztepe Training and Research Hospital, Istanbul; ⁷Rheumatology, Medicalpark Hospital, Gaziantep, Turkey; ⁸Barbara Volcker Center for Women and Rheumatic Diseases, Hospital for Special Surgery, Weill Cornell Medicine, New York, USA

Background: A multicenter antiphospholipid antibody (aPL) clinical database was created in 2016 with the participation of six rheumatology centres around the country. The purpose has been to better define aPL-related clinical manifestations and management strategies; and also to establish a resource for future clinical studies.

Objectives: In this first analysis, we compared the clinical and laboratory features of aPL-positive patients with and without systemic lupus erythematosus (SLE).

Methods: The demographic, clinical, laboratory, treatment characteristics of the aPL-positive patients with/without other systemic autoimmune diseases (SAID) are recorded at enrollment according to a predefined protocol. The inclusion criteria are positive aPL (lupus anticoagulant test [LA], anticardiolipin antibody [aCL], and/or anti β_2 -glycoprotein-I antibody [β_2 GPI]) based on the Updated Sapporo Antiphospholipid Syndrome (APS) Classification Criteria at least twice within one year prior to enrolment.^{1,2} For the purpose of this analysis, we only included aPL-positive patients without other autoimmune diseases (primary aPL/APS) and aPL-positive SLE patients (SLE aPL/APS).

Results: As of January 2018, 105 aPL-positive patients were recruited (mean age: 42.6 ± 10.1 [min-max: 19–70]; 83 [79%] female; and 67 [64%] with another SAID). Ten patients were excluded from the analysis due to their SAID history other than SLE. Of the remaining 95 patients, 38 (40%) had primary aPL/APS; 57 (60%) fulfilled the ACR SLE Classification Criteria; 42 (44%) had thrombotic APS (TAPS) (8 arterial, 24 venous, and 8 both); 21 (22%) had obstetric APS (OAPS); 22 (23%) had both TAPS and OAPS (7 arterial, 14 venous, and 1 both); and 10 (11%) had no TAPS/OAPS. Fifty percent of the patients had history of at least one non-criteria aPL-manifestation. Demographics, clinical and laboratory manifestations, and medications were similar between primary aPL/APS and SLE aPL/APS patients except increased frequency of autoimmune hemolytic anaemia, aCL IgG, and hydroxychloroquine use in SLE aPL/APS patients (table 1).