diary processed with FOOD CONS Foods software which allows to study in detail their eating habits. Nutritional state, muscle strength and basal metabolic rate were assessed. Alcoholism or drug abuse, diabetes mellitus, specific dietary models, treatment with drugs and/or food supplements with anti-inflammatory and/or antioxidant activities were considered exclusion criteria. Multivariate linear regression was performed with R project for Statistical Computing.

RESULTS:

<table>
<thead>
<tr>
<th>N=40</th>
<th>N=26</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, median (range) 53 (25-80) 33 (25-71)</td>
<td>BMI, median (range) 21 (19-23) 25.1 (19-33.7)</td>
</tr>
<tr>
<td>MedDietscore, median (range) 23 (26-43) 33 (23-40)</td>
<td>ESPDAI, median (range) 7 (0-16) 10 (0-16)</td>
</tr>
<tr>
<td>ESPSPI dryness, median (range) 6 (0-8) 6.3 (1.6-9)</td>
<td>ESPSPI dryness, median (range) 6 (0-10) 6.2 (0-10)</td>
</tr>
<tr>
<td>Focus score, median (range) 2.5 (0-9.6) 1.7 (0-8.6-24)</td>
<td>ASM kg, median (range) -16.8 (13.3-21.7)</td>
</tr>
<tr>
<td>IPAG meters, median (range) 1386 (99-11865)</td>
<td>6MWT meters, median (range) 595 (336-680)</td>
</tr>
</tbody>
</table>

BMI, body mass index; ASM: appendicular skeletal mass; IPAG: International Physical Activity Questionnaire; 6MWT, six minute walking test.

MIDiet was administered to 40 female SS outpatients. Even if not reaching significance, patients with a higher focus score in their MSG have a lower value of MIDiet score (p = 0.058, r = -1.00).

In 26 patients, daily food questionnaire shows that their diet consists of 43% of daily energy comes from proteins. Fat consumption is higher compared to the Italian population5. Six patients had a reduction in muscle mass; sarcopenia is not associated to ESSDAI (p = 0.610).

In 26 patients, daily food questionnaire shows that their diet consists of 43% of daily energy comes from proteins. Fat consumption is higher compared to the Italian population5. Six patients had a reduction in muscle mass; sarcopenia is not associated to ESSDAI (p = 0.610).

Focus score was performed with R project for Statistical Computing.

 modeled with respect to the antiphospholipid antibodies (aPL) profiles.

Background: In patients with the antiphospholipid syndrome (APS), concomitant systemic autoimmune rheumatic diseases (SARD) are common and often associated with more disease associated damage.

Less is known about the prevalence of non-rheumatic autoimmune diseases (NRAID) in patients with APS.

Objective: To evaluate the incidence and prevalence of concomitant autoimmune diseases (AID) in a cohort of APS patients. The risk of AID was also evaluated with respect to the antiphospholipid antibodies (aPL) profiles.

Methods: This retrospective cohort study comprises consecutive patients identified with APS through review of electronic medical records at Karolinska University Hospital, Sweden between 2014 and 2020. Exclusion criteria were misdiagnosis and age <18. Descriptive statistics was used for baseline data and multivariable Cox proportional hazard regression analysis to investigate the risk factors to develop new onset AID. Ethical approval was obtained from the Swed- ish Ethical Review Authority (2020-02333).

Results: Of 271 included patients, 66% were women and the median age at diagnosis of APS was 43 years (IQR 31–55). At inclusion, 130 (48%) patients presented with other AID; 101 (37%) of them had a concomitant SARD while 54 (19%) had a NRAID. Systemic lupus erythematosus (SLE) was the most frequent in 30% of patients, followed by autoimmune thyroid disease (ATD) in 10% of patients.

In addition, 35 (13%) APS-patients developed AID during the study period, corres- ponding to an incidence rate of 28.4 (95% CI: 19.3-40.3) per 1.000 person-years with mean time at risk of 4 (s2.2) years. Twenty-one (8%) patients developed a SARD and further 14 (5%) were diagnosed with a NRAID.

The cumulative incidence for AID was significantly higher in patients with high titers of IgG aPL. Patients that developed SARD had significantly higher median titers of a-j2GPI IgG isotype, p = 0.05. In the NRAID group, median a-j2GPI and aCL IgG isotypes were significantly increased, p = 0.02 and p = 0.04, respectively. The hazard ratio to develop diagnosis of AID was significantly increased in patients with higher titers of the IgG aPL aPL (HR 2.4 95% CI; 1.1-5.3). Obstet- ric APS manifestations were associated with a significantly increased hazard ratio of 2.8 (95% CI; 1.7-77) to develop SARD, and also trendwise for AID, as a compound variable.

During the study period, 52 patients had at least one new APS manifestation, as defined by the Sydney criteria (1). In comparison to patients without new mani- festations, these patients had significantly higher median titers of aPL of the IgG isotype, and concomitant AID at first visit (p<0.01, p=0.02, respectively).

Conclusion: APS patients are at high risk to develop other AID, and APS patients with concomitant AID had an increased risk to develop new clinical APS manifestations. These findings might be helpful when considering risk stratifica- tion and alternate treatment options in this patient group.

REFERENCES:


Disclosure of Interests: I have no acknowledgements to declare.

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AB0319 HEALTH-RELATED QUALITY OF LIFE ASSESSED BY LUPUSQOL IN 1102 LUPUS PATIENTS FROM RUSSIAN FEDERATION (RF), KIRGHIZSTAN AND KAZAKHSTAN (RENAISSANCE COHORT)

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Background: Health related quality of life (HRQoL) is an aspect that is generally underexplored in the routine clinical practice but it is attracting more and more attention. Lack of concordance between physician derived outcomes of disease activity and patient HRQoL outcomes is commonly found in SLE. This is due to the fact that the former is influenced by the immunologic pathogenic inflammatory mechanisms while the latter may be affected by a variety of psychological, social and co-morbid conditions.

Objectives: The aim of this study was to compare health-related quality of life assessed by LupusQol in 1102 Lupus patients (pts) from the Russian Federa- tion, Kirghizstan and Kazakhstan (RENAISSANCE cohort).

Methods: 1102 SLE pts who fulfilled SLICC 2012 criteria were enrolled into the study. The SLEDAI 2K index activity, SLICC damage index, and HRQoL using the LupusQol questionnaire (validated for the Russian-speaking population in 2018) were evaluated. Descriptive statistics, Spearman’s correlation coefficients were performed to analyze the data.

Results: 400/600/102 Lupus pts from RF, Kirghizstan and Kazakhstan respectively were studied. The pts were predominantly female (91%/93%/98%) and of indigenous nationality (Russian 83%/ Kirghiz 88%/ Kazakh 78%) with a means/SD age of 34.2±11.3, 35.0±12.2, 33.3±10.5 years (p=0.36; p=0.96 respectively). The mean disease duration (Me) in RF/Kazakhstan was 6 [3;12]/ 5[2;9] years (p<0.15), in Kirghizstan - 2[0,6; 7] years(p≤0,05). Kazakh pts (p=0,00) had higher disease activity (SLEDAI 2K) 17±6.8 than Kirghiz pts 15±1.8 and Russian pts 9,4±8.1. SLICC damage index 2,39±1,61 in Kazakh pts (p=0,00) was higher than in Russian 1,57±1,54 and Kirghiz 0,79±0,32 pts. 54% pts from RF and 58% from Kazakh had high education. Only 33% pts from Kirghiz also had high education. 45%/24%/33% pts from RF, Kirghizstan and Kazakhst- an respectively had steady-state condition, 45%/40%/42% pts had a perceived disabil- ity benefits. HRQoL assessed by LupusQol was low in all patients with SLE. The mean scores for each of the domains of LupusQol are shown in Table 1.

The statistically significant lowest HRQoL in all 8 domains is obtained among patients with concomitant AID had an increased risk to develop new clinical APS manifestations. These findings might be helpful when considering risk stratifica- tion and alternate treatment options in this patient group.

REFERENCES:


Disclosure of Interests: I have no acknowledgements to declare.

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