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Mental Health in Patients with Axial Spondyloarthritis: Increasing Our Understanding of the Disease. Results from the Spanish Atlas

Marco Garrido-Cumbrera¹, Victoria Navarro-Compán², David Galvez-Ruiz¹, Carlos Jesus Delgado Dominguez¹, Pilar Font-Ugalde³, Olta Brace¹, Pedro Zarco⁴, Jorge Chacon-Garcia¹ and Pedro Plazuelo-Ramos⁵, ¹Universidad de Sevilla, Seville, Spain, ²Rheumatology, Hospital Universitario La Paz, Madrid, Spain, ³Rheumatology service, IMIBIC/Reina Sofia Hospital/University of Cordoba, Cordoba, Spain, ⁴H Fundación Alcorcón, Alcorcón, Spain, ⁵CEADE, Madrid, Spain

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Session Title: Epidemiology and Public Health Poster II: Rheumatic Diseases Other than Rheumatoid Arthritis

Session Time: 9:00AM-11:00AM

Background/Purpose: This study's aim was to assess the association between sociodemographic characteristics, disease progression, and mental health comorbidity with risk of mental disorders (RMD).

Methods: In 2016 a sample of 680 axSpA patients was interviewed as part of the Spanish Atlas. To quantify the RMD, Goldberg's General Health Questionnaire (GHQ-12) scale was employed. Possible RMD predictors analysed were: sociodemographic characteristics (age, gender, being part of a couple, patient association membership, job status); disease characteristics (BASDAI, spinal stiffness ranging from 0-3, functional limitation in 18 daily activities ranging from 0-3); and mental health comorbidities (depression and anxiety). All clinical variables showed a Cronbach's alpha coefficient guaranteeing the reliability of the scales used. First, a descriptive analysis was employed to describe the sample and study variables. Second, we performed univariate correlation and homogeneity analyses between each predictor (independent variable) and RMD (GHQ-12). Third, selection of variables that showed statistical significance in the univariate analyses in order to conduct a multiple hierarchical and stepwise regression analysis.

Results: All variables except educational level and thoracic stiffness showed significant univariate correlation with RMD. BASDAI, functional limitation and age showed higher coefficient ($R = 0.543$, $R = 0.378$, $R = -0.174$, respectively). Multiple Hierarchical regression analysis showed as sociodemographic variables explained in great detail the RMD ($R^2 = 83.2\%$). By contrast, having established sociodemographic as a control variable, the inclusion of depression and anxiety to the model increase the R^2 value to just 0.6% ($p = 0.001$), while the inclusion of variables related to the disease characteristics add 5.5% ($p = 0.000$) to the GHQ-12 punctuation variability. The only variables presenting a significant coefficient different from 0 were BASDAI (0.52 , $p = 0.000$) and functional limitation (0.14 , $p = 0.004$). This suggests that once the sociodemographic and mental comorbidity variables are established, a change to BASDAI levels or functional limitation impacts the GHQ-12

score. In the stepwise regression analysis, four variables (BASDAI, functional limitation, association membership, cervical stiffness) showed a significant relation to GHQ-12 and explained the majority of RMD variability. BASDAI displayed the highest explanatory degree ($R^2 = 0.875$).

Table 1. Sample characteristics (n = 474, unless other specified).

Variables	Values (means \pm SD or percentage)
<i>Age, mean \pm SD</i>	45.43 \pm 10.78
<i>Sex, No. of men</i>	233 (49.16%)
<i>Having a couple, No. of participants (N=444)</i>	386 (86.94%)
<i>Education level, No. of university studies</i>	185 (39.30%)
<i>Job status, No. of unemployed</i>	68 (14.35%)
<i>Association Membership</i>	227 (47.89%)
<i>BASDAI, mean \pm SD (N=442)</i>	5.49 \pm 2.17
<i>Cervical stiffness, No. (N=447)</i>	201 (44.97%)
<i>Thoracic stiffness No. (N=435)</i>	186 (42.76%)
<i>Lumbar stiffness No. (N=458)</i>	288 (62.88%)
<i>Functional Limitation, mean \pm SD (N=473)</i>	27.54 \pm 12.78
<i>Depression, No. (%) (N=474)</i>	99 (20.89)
<i>Anxiety, No. (%) (N=474)</i>	134 (28.27)
<i>GHQ-12, mean \pm SD</i>	18.30 \pm 8.01

Conclusion: Patients at certain sociodemographic levels are more prone to present a higher BASDAI. Taking these conditions for granted, the degree of disease progression measured by BASDAI is a good indicator of RMD. Therefore, in those with higher disease activity, psychiatric evaluation and intervention should be considered within the medical treatment.

Disclosure: M. Garrido-Cumbrera, None; V. Navarro-Compán, None; D. Galvez-Ruiz, None; C. J. Delgado Dominguez, None; P. Font-Ugalde, None; O. Brace, None; P. Zarco, None; J. Chacon-Garcia, None; P. Plazuelo-Ramos, None.

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