

OP0172-PARE **WORKING TOGETHER FOR CHILDREN AND FAMILIES LIVING WITH RHEUMATIC AND MUSCULOSKELETAL DISEASES: THE EUROPEAN NETWORK FOR CHILDREN WITH ARTHRITIS (ENCA)**

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Background: The European Network for Children with Arthritis (ENCA) is a pan-European network led for and by parents and young people with rheumatic and musculoskeletal diseases (RMDs). ENCA's activities are delivered voluntarily by parents and young people passionate about improving the care and treatment of children and young people (CYP) with RMDs. This is done through education, innovation, networking and empowerment. ENCA is associated with the Paediatric Rheumatology European Society (PReS).

Objectives: ENCA aims to facilitate the exchange and dissemination of information, knowledge and best practice with regards to paediatric RMDs across Europe, working in partnership with national patient organisations for CYP and families living with RMDs. ENCA also aims to provide international awareness, engagement and research opportunities.

Methods: ENCA is managed by a board of elected members, who are all volunteers with national patient organisations. ENCA hosts an annual conference in conjunction with PReS, inviting attendance from national patient organisations. Aside from a newly refreshed website, ENCA hosts a closed Facebook and WhatsApp group for national patient organisation representatives, enabling real-time communication and the instant sharing of knowledge and best practice. ENCA hosts an annual 'Fun with JIA challenge', aimed at empowering CYP with RMDs to share their story; and has been involved in establishing a new international awareness campaign for 2019, titled 'World young Rheumatic Disease (WORD) day'. In addition, ENCA cooperates with a number of international organisations with shared goals in rheumatology, representing the paediatric voice.

Results: Since its inception in 2002, ENCA has enhanced the communication and shared learning between national patient organisations, by embedding the voice of CYP and families into research and advocacy activities. The 'Fun with JIA challenge' delivered since 2017 has provided CYP with RMDs across Europe with the opportunity to share their stories through film. In addition, ENCA's cooperation with PReS has recently facilitated the planning and launch of WORD day on March 18th, 2019. The need for a specific paediatric RMD awareness day was identified as a priority from national representatives, in light of the relatively limited attention that paediatric RMDs receive. WORD day helped to raise parental and professional awareness about paediatric RMDs.

Conclusion: As parents and young people living with RMDs, ENCA has enabled individuals and national patient organisations to develop their knowledge in relation to paediatric RMDs. ENCA provides several opportunities for networking and the sharing of best practices between national patient organisations and professionals, through informal and formal activities focused on paediatric RMDs. International networking helps individuals and communities to navigate through challenges with the support of others with a shared-goal.

REFERENCE:

- [1] Egert Y, et al. PARE0005|Brief Overview of ENCA (European Network for Children with Arthritis). *Ann Rheum Dis* 2016; 75:1321.

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OSTEOARTHRITIS: RESEARCH IN MOTION

OP0173 **EROSIVE HAND OSTEOARTHRITIS ASSOCIATES WITH LOW OCCURRENCE OF KNEE OSTEOARTHRITIS**

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Background: Erosive hand osteoarthritis (OA) is considered a more severe form of hand OA that includes pain and inflammation (1,2). However, more data are needed regarding the different hand OA phenotypes, especially erosive and non-erosive, to conclude if they represent two different subsets of hand OA or even different degrees of affectation of the same disease (1,3,4)

Objectives: To compare the clinical pattern of patients with erosive hand OA with those that do not show this phenotype

Methods: The study was conducted in the Prospective Cohort of Osteoarthritis A Coruna (PROCOAC). This cohort consists of 1107 subjects, of which 747 suffered from radiographic hand OA. We split the cohort into patients with and without erosive hand OA, and analyzed both clinical and demographic data within each group, together with the assessment of the AUstralian CANadian Osteoarthritis Hand (AUSCAN) index. The study consisted in a univariate analysis comparing different variables between both groups, followed by a stepwise logistic regression analysis taking into account the significant confounder variables analyzed in the Univariate approach

Results: The mean age of the cohort was 63.23±8.85 years and consisted of 627 females and 120 males. Of the 747 OA patients, 179 had erosive hand OA and 568 did not. The univariate analysis revealed that patients with erosive hand OA were younger (60.25±8.10 vs 64.16±8.88;p<0.001), smokers (p=0.005), with lower body mass index (p=0.005), increased inflammatory episodes (p<0.001), with family history or suffering from psoriasis (p=0.021) and less number of damaged joints (p<0.001). In addition, metabolic syndrome (MetS) was not associated with erosive hand OA and, despite the higher number of females suffering from hand OA, no significant differences between erosive and non-erosive phenotypes in terms of gender were detected.

The regression model confirmed the strong association of age (OR=0.9734; 95% CI=0.947-0.999; p=0.044), inflammation (OR=5.096; 95% CI=2.868-9.058; p<0.001), number of damaged joints (OR=0.490; 95% CI=0.307-0.782; p<0.003) and, interestingly, showed MetS as a risk factor too (OR=2.146; 95% CI=1.250-3.682; p=0.006). Regarding the number of damaged joints, a lower frequency of both knee and hip OA was detected in patients with erosive hand OA compared with non-erosive patients (24.4% vs 54.7% for knee OA prevalence and 11.6% vs 16.7% for hip OA prevalence); however, only knee OA occurred at a significant lower frequency (OR=0.267; 95% CI=0.182-0.393; p<0.001). In addition, the three subscales of the AUSCAN score showed significantly higher mean values in patients with the erosive phenotype: pain (57.67±29.30 vs 43.86±31.45; p<0.001), function (55.61±27.68 vs 40.53±29.48; p<0.001) and stiffness (55.40±32.26 vs 40.95±36.23;p=0.001)

Conclusion: The erosive hand OA phenotype is associated with more inflammation in younger patients, and worst AUSCAN score. Contrarily to the non-erosive phenotype, the presence of erosions associates with a lower occurrence of other forms of OA, specially knee OA. In our cohort, erosive hand OA seems to be more localized in the hand and it also seems to be associated with psoriasis. This different profile could serve the clinicians to provide personalized prevention strategies, and the researchers to investigate the pathogenesis of this specific phenotype

REFERENCES:

- [1] Haugen, et al. *Osteoarthritis Cartilage* 2016;24:647-654
 [2] Marshall, et al. *Ann Rheum Dis* 2015;74:136-141
 [3] Dolzani, et al. *Clin Exp Rheumatol* 2011;29:1006-1009
 [4] Punzi, et al. *Ann Rheum Dis* 2005;64:955-957

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