laboratories, to assess the suitability of the new candidate standard and calibrate it against W1066.

Results: The results of the collaborative study will be used to propose a unitage to the new candidate standard in RF and ACPA for consideration at the WHO Expert Committee for Biological Standardisation.

Conclusion: Standardisation of diagnostic laboratory tests involves the development of a physical sample for each analyte. The use of these international reference standards as primary calibrants promotes the harmonisation of test results between different assays. Because the assays use a similar scale, their results are easily understood, which ultimately increases the impact of published research and clinical findings on our knowledge of the disease.

REFERENCES

- [1] WHO Expert Committee on Biological Standardization, sixty-seventh report. Geneva: World Health Organization; 2017 (WHO Technical Report Series; no. 1004). 2017

Disclosure of Interests: None declared

DOI: 10.1136/annrheumdis-2019-eular.4068

AB1135

THE MANAGEMENT OF INTERSTITIAL LUNG DISEASES: THE IMPORTANCE OF THE RHEUMATOLOGIC EXPERTISE IN MULTIDISCIPLINARY MEETINGS

Enrico De Lorenzo¹, Gerlando Natalello¹, Lucrezia Verardi¹, Silvia Laura Bosello², Elisa Gremese^{1,2}. ¹Catholic University of the Sacred Heart, Institute of Rheumatology, Rome, Italy; ²Fondazione Policlinico Universitario A. Gemelli IRCCS, Department of Rheumatology, Rome, Italy

Background: Multidisciplinary Team (MDT) meetings are the current "gold standard" in interstitial lung disease (ILD) diagnosis. Requisite participants are respiratory physicians, radiologists and pathologists. A rheumatologist physician is not routinely involved in MDT even if up to 20% of ILD are related to connective tissue disease, rheumatoid arthritis or systemic vasculitis.

Objectives: The aim of this study is to evaluate the prevalence and predictors of systemic rheumatological diseases in a cohort of patients with ILD, evaluated by a rheumatology specialist on the advice of MDT in a university hospital.

Methods: Thirty-two patients with ILD, evaluated at dedicated MDT were referred to a rheumatologist in 2018, usually for autoantibodies positivity or clinical history suspected for a rheumatological disorder. Rheumatologic evaluation included physical examination, routine blood and urine tests, serum levels of C3 and C4, ANA, Rheumatoid Factor (RF), ANCA, anti-Sm, anti-RNP, anti-Ro/SSA, anti-La/SSB, anti-Sm, anti-Jo1, anti-dsDNA and anti-CCP antibodies. Family history of autoimmune diseases, presence of rheumatologic red flags (Raynaud's phenomenon, photosensitivity, inflammatory skin manifestations, sicca syndrome, recurrent fever, inflammatory arthralgias, paresthesias, oral or genital aphthosis, thrombosis and recurrent miscarriages), respiratory symptoms and pulmonary function test were also evaluated. When indicated, capillaroscopy, joint imaging and salivary gland biopsy were performed.

Results: Twenty-one patients (65.6%) were female. At the time of rheumatological evaluation, the patients had a mean age of 64.4±12.6 years with a mean period of 21.1±39.6 months since the first identification of ILD on a CT scan. Twenty-five (78.1%) and 18 (56.3%) patients respectively complained exertional dyspnea and cough, 6 patients (18.8%) were on oxygen therapy and 13 (40.6%) had a FVC<80%. The pattern on CT scan suggested a UIP, NSIP and OP respectively in 18 (56.3%), 6 (18.8%) and 2 (6.3%) patients. Ten patients (31.3%) were diagnosed with a defined rheumatological condition (4 with Sjogren Syndrome, 3 with scleroderma, 1 with Rheumatoid Arthritis, 1 with Mixed Connective Tissue Disease and 1 with Granulomatosis with Polyangiitis); 7 (21.9%) were diagnosed with Interstitial Pneumonia with autoimmune feature (IPAF). A rheumatological diagnosis was not statistically associated with any specific rheumatologic red flag, routine laboratory abnormalities or family history of systemic autoimmune disease. Three out of 13 capillaroscopies and 3 out of 9 salivary glands biopsies performed had diagnostic findings. The only immunologic abnormalities associated to a rheumatologic diagnosis were RF positivity (20.0%vs54.5%, p=0.024) and p-ANCA positivity (10.0%vs40.9%, p=0.042), both with low specificity (respectively 15.0% and 5.0%) and sensitivity (respectively 55.6% and 42.9.0%). After diagnosis, 11 patients (34.4%) started immunosuppressive therapy and 7 (21.9%) started antifibrotic therapy.

Conclusion: Single clinical or laboratory abnormalities are not strongly associated to a rheumatological diagnosis in patients with ILD, suggesting

that only a comprehensive rheumatological evaluation allows correct classification of the disease associated with ILD and is mandatory to make or exclude a diagnosis. In this study, we evaluated only preselected patients by ILD-MDT, but the results indirectly suggest that direct participation of rheumatologist to MDT is advisable to increase accuracy and reduce delay in diagnosis and treatment.

Disclosure of Interests: Enrico De Lorenzo: None declared, Gerlando Natalello: None declared, Lucrezia Verardi: None declared, Silvia Laura Bosello: None declared, Elisa Gremese Consultant for: AbbVie, BMS, Celgene, Janssen, Lilly, MSD, Novartis, Sanofi, UCB, Roche, and Pfizer, Speakers bureau: BMS, Speakers bureau: Roche, Speakers bureau: AbbVie, BMS, Celgene, Janssen, Lilly, MSD, Novartis, Sanofi, UCB, Roche, and Pfizer

DOI: 10.1136/annrheumdis-2019-eular.6364

AB1136

CD26: A POTENTIAL NOVEL HISTOLOGICAL MARKER OF IDIOPATHIC INFLAMMATORY MYOPATHIES

Rebecca De Lorenzo¹, Clara Sciorati¹, Antonella Monno¹, Francesco Bonomi², Silvia Cavalli², Antonio Francesco Di Naro³, Stefano Previtali⁴, Patrizia Rovere-Querini¹. ¹San Raffaele Scientific Institute, Laboratory of Innate Immunity and Tissue Remodelling, Milan, Italy; ²Vita-Salute San Raffaele University, Milan, Italy; ³Adienne Pharma and Biotech, Lugano, Switzerland; ⁴San Raffaele Scientific Institute, Experimental Neurology Institute (INSPE), Milan, Italy

Background: Idiopathic inflammatory myopathies (IIM) are a heterogeneous group of acquired skeletal muscle disorders including polymyositis (PM), dermatomyositis (DM), inclusion body myositis (IBM) and immune-mediated necrotizing myopathy (IMNM), characterized by immune-mediated muscle damage¹. Activated T cells are the predominant inflammatory infiltrates in muscle biopsies of PM and DM patients and the lack of T regulatory cells (T_{regs}) has been implicated in the persistence of muscle damage. CD26 is an intrinsic membrane glycoprotein and a serine exopeptidase involved in the activation of T lymphocytes and amplification of inflammatory cytokines production. The enzymatically active form of CD26 is selectively expressed by activated T cells and has been described as a negative selection marker for human T_{regs}²⁻⁵.

Objectives: The aims of this study were to evaluate the expression of CD26 in muscle biopsies of IIM patients and to correlate it with patients' clinical and histological features.

Methods: Immunofluorescence was used to evaluate CD26 expression and co-localization with CD3 and CD31, markers of T cells and endothelial cells respectively, in muscle biopsies of 6 DM, 6 PM, 3 IBM and 3 IMNM patients and of 6 healthy controls.

Results: We found that CD26 is preferentially expressed in muscle biopsies of IIM patients with respect to controls and that its level of expression is higher in DM patients. In muscle biopsies of IIM patients, CD26 is distributed not only in the extracellular matrix surrounding myofibers and infiltrating leukocytes, but also at the level of T cell membranes and endothelial cells. Specifically, CD26 co-localization with CD31 is more prominent in DM muscle biopsies. We could not find any association between vessel morphology in terms of size and shape and CD26 endothelial expression, suggesting that CD26 is expressed at the perivascular level independently of the degree of vessel dysfunction. With regard to clinical features, we found that CD26 is more expressed in patients presenting the typical DM rash. Moreover, CD26 expression was found not to be significantly associated with the degree of muscle weakness nor with the presence of interstitial lung disease, dysphagia, myalgias or mechanic's hands. As for histological data, higher levels of CD26 expression were found in biopsies with perivascular inflammatory infiltrates, especially T lymphocytes and macrophages.

Conclusion: Our data suggest that CD26 may represent a suitable marker for the diagnosis of IIM and a potential novel target for selective immune-therapies.

REFERENCES

- [1] Dalakas, M. C. (2015). "Inflammatory Muscle Diseases." *N Engl J Med* 373(4): 393-394.
- [2] Goebels N, Michaelis D, Engelhardt M, et al. Differential expression of perforin in muscle-infiltrating T cells in polymyositis and dermatomyositis. *J Clin Invest* 1996;97:2905-10.
- [3] Waumans Y, Baerts L, Kehoe K, Lambeir A M, De Meester I. The dipeptidyl peptidase family, prolyl oligopeptidase, and prolyl carboxypeptidase in the immune system and inflammatory disease, including atherosclerosis. *Front Immunol* 2015; 6:387.