

This is a repository copy of GRAPPA 2019 Project Report.

White Rose Research Online URL for this paper: https://eprints.whiterose.ac.uk/165787/

Version: Accepted Version

#### Article:

Goel, N, Coates, LC orcid.org/0000-0002-4756-663X, De Marco, G orcid.org/0000-0003-2406-161X et al. (11 more authors) (2020) GRAPPA 2019 Project Report. The Journal of Rheumatology Supplement, 96. pp. 53-57. ISSN 0380-0903

https://doi.org/10.3899/jrheum.200129

© 2020. This is an author produced version of an article published in The Journal of Rheumatology Supplement. Uploaded in accordance with the publisher's self-archiving policy.

#### Reuse

Items deposited in White Rose Research Online are protected by copyright, with all rights reserved unless indicated otherwise. They may be downloaded and/or printed for private study, or other acts as permitted by national copyright laws. The publisher or other rights holders may allow further reproduction and re-use of the full text version. This is indicated by the licence information on the White Rose Research Online record for the item.

#### Takedown

If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing eprints@whiterose.ac.uk including the URL of the record and the reason for the withdrawal request.



eprints@whiterose.ac.uk https://eprints.whiterose.ac.uk/

# For submission to The Journal of Rheumatology

# **GRAPPA 2019 Project Report**

Niti Goel, Laura C. Coates, Gabriele De Marco, Lihi Eder, Oliver FitzGerald, Philip Helliwell, Ying Ying Leung, Walter P. Maksymowych, Philip J. Mease, Mikkel Østergaard, Denis O'Sullivan, Denis Poddubnyy, Christopher T. Ritchlin, Dafna D. Gladman

**ABSTRACT**: At the 2019 annual meeting of the Group for Research and Assessment of Psoriasis and Psoriatic Arthritis (GRAPPA), members received updates on several ongoing efforts. Among them were updates on research, including the trainee symposium, pilot research grants, and Collaborative Research Network; GRAPPA's patient research partners; education, including the slide collection; treatment recommendations; and additional undertakings related to advancing the understanding of disease aspects, including the Outcome Measures in Rheumatology (OMERACT)-GRAPPA outcome measure, axial, ultrasound enthesitis projects, as well as the early psoriatic disease systematic literature review and magnetic resonance imaging.

Key Indexing Terms: Psoriasis, Psoriatic Arthritis, Education, Research, GRAPPA Source of Support: None

Author Information: N Goel, MD, [ORCID: 0000-0001-5869-5157], Patient Research Partner, Adjunct Assistant Professor, Duke University School of Medicine, Durham, North Carolina, USA, agwngw1@gmail.com; LC Coates, MB ChB, PhD, [ORCID: 0000-0002-4756-663X], Department of Orthopaedics, Rheumatology, and Musculoskeletal Sciences, University of Oxford, Oxford, United Kingdom, laura.coates@ndorms.ox.ac.uk; G De Marco, MD, [ORCID: 0000-0003-2406-161X], Post-CCT Clinical Research Fellow, UK National Institute for Health Research (NIHR) Leeds Biomedical Research Centre, Leeds Teaching Hospitals National Health Service (NHS) Trust, Leeds Institute of Rheumatic and Musculoskeletal Medicine, University of Leeds, Leeds, United Kingdom, g.demarco@leeds.ac.uk; L Eder, MD, PhD, [ORCID: 0000-0002-1473-1715], Assistant Professor of Medicine, University of Toronto and Women's College Hospital, Toronto, Ontario, Canada, Lihi.EDER@wchospital.ca; O FitzGerald, MD, FRCP(UK), FRCPI, [ORCID: 0000-0002-6607-6070], Newman Clinical Research Professor, Conway Institute for Biomolecular Research, University College Dublin, Dublin, Ireland, oliver.fitzgerald@ucd.ie; PS Helliwell, DM, PhD, FRCP, UK NIHR, [ORCID: 0000-0002-4155-9105], Institute of Rheumatic and Musculoskeletal Medicine, University of Leeds, Leeds, United Kingdom, and Bradford Hospitals UK NHS Foundation Trust, Bradford, United Kingdom, p.helliwell@leeds.ac.uk; YY Leung, MB ChB, MD, [ORCID: 0000-0001-8492-6342, Associate Professor, Duke-NUS Medical School, Singapore, Department of Rheumatology and Immunology, Singapore General Hospital, Singapore, katyccc@hotmail.com; WP Maksymowych, MB, ChB, FRCPC, [ORCID: 0000-0002-1291-1755], Professor of Medicine, University of Alberta, Edmonton, Canada, walter.maksymowych@ualberta.ca; PJ Mease, MD, [ORCID: 0000-0002-6620-0457], Rheumatology Research, Swedish Medical Center and University of Washington School of Medicine, Seattle, Washington, USA, pmease@philipmease.com; M Østergaard, MD, PhD, DMSc, [ORCID: 0000-0003-3690-467X], Professor of Rheumatology at University of Copenhagen and the

National Hospital (Rigshospitalet), Denmark, mo@dadInet.dk; D O'Sullivan, BE, [ORCID: 0000-0002-6607-6070], Patient Research Partner, Our Lady's Hospice & Care Services, Dublin, Ireland, denis\_osullivan@ymail.com; D Poddubnyy, MD, MSc (Epi), [ORCID: 0000-0002-4537-6015], Professor of Rheumatology, Charité-Universitätsmedizin Berlin, and German Rheumatism Research Centre, Berlin, Germany, denis.poddubnyy@charite.de; CT Ritchlin, MD, MPH, [ORCID: 0000-0002-2602-1219], Professor of Medicine, Division of Allergy, Immunology, and Rheumatology, University of Rochester Medical Center, Rochester, New York, USA, Christopher\_Ritchlin@URMC.Rochester.edu; DD Gladman, MD, FRCPC, [ORCID: 0000-0002-9074-0592], Professor of Medicine, University of Toronto, Senior Scientist, Krembil Research Institute, Director, Psoriatic Arthritis Program, University Health Network, Toronto Western Hospital, Toronto, Ontario, Canada, dafna.gladman@utoronto.ca.

**Corresponding Author**: Niti Goel, MD, 4809 Taproot Ln, Durham, NC 27705, USA, phone 1 919 407 0477; email agwngw1@gmail.com

Running Footline: GRAPPA 2019 Project Report

**Word Count**: 2886 words, including text (2795) and abstract (91), but excluding title page

Members of the Group for Research and Assessment of Psoriasis and Psoriatic Arthritis (GRAPPA) continue to pursue the core objectives of GRAPPA's mission, specifically, to provide education, identify research assessment tools, and pursue research in disease pathophysiology. At the 2019 annual GRAPPA meeting in Paris, France, members received updates on research, including the trainee symposium, pilot research grants, and Collaborative Research Network (CRN); GRAPPA's patient research partners (PRPs); education, including the slide collection; treatment recommendations; and additional undertakings related to advancing the understanding of disease aspects, including the Outcome Measures in Rheumatology (OMERACT)-GRAPPA outcome measure, axial, and ultrasound enthesitis projects, as well as the early psoriatic arthritis (PsA) systematic literature review and magnetic resonance imaging (MRI).

#### **Research Committee**

Dr. Oliver FitzGerald reported that the GRAPPA CRN held a very productive premeeting. The pre-meeting was held to review progress on the development of an electronic case report form (eCRF), standardized operating procedures (SOPs), and shared database to be used in CRN studies; to obtain feedback on the SOPs needed for sample collection; and to propose an investigator-initiated study that would allow for testing of both the eCRF and SOPs. The appointment of a coordinator for the CRN was announced with the remit of the coordinator being to develop the eCRF and SOPs. As a result of the meeting, the CRN committee planned to proceed with a more detailed treatment response study design to guide the development of the eCRF and SOPs. Continued progress on the PsA Biomarkers of Joint DAMage (BioDAM) project was reported, especially with relation to obtaining samples from various industry-sponsored trials in PsA. This included a signed contract with Lilly, with shipping of samples from the ixekizumab PsA program planned in fall 2019; a contract under discussion with Amgen to access samples from the SEAM study; and a contract under discussion with Pfizer related to the tofacitinib program to predict treatment response.

GRAPPA is now registered in the European Union (EU) in The Netherlands to allow GRAPPA to participate in EU-funded research programs. As part of this effort, GRAPPA continues to pursue a psoriatic disease-specific call for research in areas of major unmet need with the EU-based Innovative Medicines Initiative (IMI). While the call for proposals was issued in late 2018, it will likely not proceed until early 2020. Meanwhile, contact was made with the United States (US) National Institutes of Health (NIH) regarding the possibility of an Accelerated Medicines Partnership (AMP)-type program in psoriatic disease.

Dr. Christopher Ritchlin reported that there were 26 applications for the pilot research grants compared to 22 in 2018. Once again, 3 were awarded at \$25,000 each to Maria Angioni (Italy) for her study evaluating the pharmacogenetics of treatment response in PsA; Alla Ishchenko (Belgium) on metabolomic profiling in psoriatic disease; and Zhenrui Shi (USA) on the preclinical analysis of CCR6 and CCL20 in mouse and human joints. The Research Committee will send a new request for applications in early 2020 with additional information and details provided at that time.

Additionally, results to date from the 3 research projects awarded in 2018 were presented. Dr. Fardina Malik (USA) presented on the effect of medium-chain fatty acid (MCFA) supplementation on PsA and psoriasis. She noted that 39% of psoriatic disease patients report the use of complementary and alternative medicine. She explored how MCFA supplementation, specifically pure extra virgin coconut oil (PEVCO), regulates psoriatic disease. Theoretically, MCFA in immune regulation could alter gut microbiota, have an antimicrobial effect, modulate T cell function, increase interleukin (IL)-10 levels, and decrease levels of tumor necrosis factor alpha  $(TNF\alpha)$ . In the study, patients with active psoriasis with or without PsA began PEVCO supplementation for 6 weeks. They underwent assessment of their skin, joints, and microbiome, and they completed the Multi-dimensional Health Assessment Questionnaire. So far, 14 individuals have been recruited with a median age of 42.5 years (57% female). While no changes in clinical disease activity were seen, after 3 weeks of MCFA supplementation, a change in the levels of firmicutes and clostridia was seen in the gut microbiome as well as decreases in the level of IL-22.

Kim Wervers and Dr. Hannah den Braanker (The Netherlands) presented on the response to methotrexate (MTX) and a quest for personalized medicine in PsA. Their objectives were to determine how often a good response to MTX occurs in daily practice in early PsA patients and what differences exist between responders versus non-responders. They evaluated 219 PsA patients with 1 year or less of oligo/polyarthritis of whom 183 (84%) started on MTX at the highest tolerated dose within 6 months of diagnosis. Ninety patients (49%) continued MTX monotherapy at 1 year. Of these patients, 44 (49%) achieved minimal disease activity (MDA) by 6 months, and only 33 maintained MDA status at 1 year. MTX monotherapy was

06 January 2020

concluded to be insufficient therapy for early PsA. While folate and vitamin B12 levels were similar at baseline, differences between responders and non-responders included significantly higher baseline levels of IL-23, TNF $\alpha$ , interferon gamma, granulocyte-macrophage colony stimulating factor, and IL-10. Further, levels of IL-23 and IL-10 evolved differently on MTX monotherapy in responders versus non-responders. Further work is ongoing to characterize the patients.

Dr. Philip Helliwell presented Graham Chapman's project on behalf of the EffeCt of Psoriatic Arthritis on Plantar shEar Stress (ESCAPES) study group. This ongoing study aims to examine the hypothesis that abnormal shear stresses in the foot are a contributory factor in the development of dactylitis of the toes in PsA. As part of this project, a new custom-made shear stress device was manufactured and is now ready for laboratory in vivo measurements. The project is planned to be completed by summer 2020.

# MRI

Drs. Walter Maksymowych and Mikkel Østergaard presented an update on the ongoing work with whole-body MRI methodology for the objective assessment of inflammation in synovium, tendon, and bone, together with large joint-scoring methodologies for the assessment of inflammation in the hip and knee.

A preliminary scoring method has been developed for whole-body MRI that assesses synovitis and osteitis in 83 peripheral joints and 33 entheses (0-3 grading scheme per joint or enthesis) with good reliability attained in a preliminary exercise of 8 patients. The sensitivity to change and discriminatory ability between spondyloarthritis (SpA) patients treated with TNF-inhibitors and placebo have been demonstrated. This method is expected to be particularly relevant in patients with PsA.

The Hip MRI Inflammation Scoring System (HIMRISS), a granular method for scoring bone marrow lesions in the hip, has been developed and is based on the use of software that applies an overlay to segment the femoral head and acetabulum. This allows multiple regions to be scored directly on a web-based interface of consecutive MRI slices through the joint (scoring range 0-100), dispensing entirely with the use of scoring spreadsheets. Excellent reliability has been documented in patients with osteoarthritis (OA) of the hip for detection of lesions at a cross-sectional level, and very good reliability has been documented in detecting change 8 weeks after intra-articular glucocorticoid injection in MRIs from 90 patients assessed by 8 readers.

The system is now ready for testing in PsA (e.g., assessing the discrimination between treatment groups in clinical trials). Segmentation based on hip joint effusion/synovitis is also performed. A quantitative assessment method, which is hoped to be superior to semi-quantitative assessment, is being developed. Work on automation and comparisons with ultrasound for measuring effusion are also being performed. For the knee, a similar quantitative method, the Knee MRI Inflammation Scoring System (KIMRISS), has been compared with the MRI OA Knee Score (MOAKS) scoring system in patients with knee OA and has been shown to have excellent reliability and better reproducibility.

Further, the group has been working on a literature review on the utility of MRI in enthesitis. Lastly, an enthesitis assessment system, including exact definitions of the various pathologies involved, has been developed and validated,

Page **8** of **14** 

with good reproducibility and sensitivity to change. The next steps are to apply the MRI techniques in clinical trials and PsA patient cohorts as well as to further develop and validate the methods on a modular plane.

### PRPs

The 2019 annual GRAPPA meeting was the seventh at which PRPs have been present. There were 11 PRPs who attended (including 2 new members) from North America, South America, Europe, and Asia. The PRP pre-meeting included presentations on precision medicine and composite endpoints. PRPs were active participants in the meeting's individual and breakout sessions. Individual PRPs also presented during the MTX panel debate. The PRPs continue to be active in several ongoing GRAPPA activities including the CRN, GRAPPA-OMERACT working group and subgroups, Best Practices, and education.

From a governance perspective, the PRPs reported that the PRP Handbook was completed, approved by the GRAPPA Executive Committee, and adopted. Along with the PRP Policies and Procedures, these 2 documents not only define how the PRP group functions and is governed but also support increased group effectiveness. Reflecting the governance, the PRP network has now been able to fully implement the Chair, Chair-elect, and Immediate Past Chair model, with each position held for 2 years.

#### **Education Committee**

As a core objective of GRAPPA's mission, GRAPPA members around the world provide psoriatic disease education to many people, including other health care professionals, patients and their families, and stakeholders. On behalf of the Education Committee, Dr. Philip Mease discussed a variety of educational activities, including educational symposia, the slide library, and educational videos. He also discussed newer efforts that are aimed at virtual opportunities, such as videoconferencing and podcasts, to reach broader non-rheumatology/dermatology audiences.

In the last decade, many events that have occurred around the world have included accredited continuing medical education (CME) and non-CME symposia. The predominantly US-based CME initiatives have been the GRAPPA-SpA Research and Treatment Network (SPARTAN) PsA-SpA symposia, offered as 2hour, half-day, and full-day symposia, and the GRAPPA Rheum-Derm PsA-psoriasis symposia. Educational symposia are also conducted around the globe with industry support. Symposia in India have partnered up to 3 international GRAPPA faculty with up to 3 regional faculty and include an ultrasound component. Other events that have occurred or are planned for 2019-2020 include (1) a symposium in Taiwan (November 2019); (2) symposia in Mumbai (November 2019 and February 2020); (3) a symposium in Dubai (September 2019); (4) events at the Asia Pacific League of Associations for Rheumatology (APLAR) meeting (symposium and GRAPPA meeting [April 2019, Brisbane, Australia; and April 2020, Kyoto, Japan]); (5) events at the Pan American League of Associations for Rheumatology (PANLAR) meeting (symposium and GRAPPA meeting [April 2019, Quito, Ecuador; and April 2020, Miami, Florida, USA]); and (6) events at the African League of Associations for Rheumatology (AFLAR) meeting (September 2019, Mauritius).

Guidelines were also provided to qualify an educational symposium as GRAPPA-approved. GRAPPA member involvement is required in planning the symposium agenda. GRAPPA staff could assist with the administrative footwork (e.g., grant applications, identifying audiences, travel logistics, and supporting document creation). GRAPPA faculty tasks include content creation and updates, as well as presentation. Further, participating faculty will have the opportunity to receive mentoring to improve cultural sensitivity during presentations.

GRAPPA aims to provide education in less-resourced areas of the world and is exploring the provision of virtual educational opportunities with videoconferencing and podcasts. Other educational endeavours include the GRAPPA slide set, which is available for use by members. The popular GRAPPA smartphone application (app) launched in 2018 and is available in 14 languages. Work is ongoing to obtain grant funding to cover the addition of new languages. The website also continues to host educational content with a plan to provide the GRAPPA videos for members to use for educational and noncertification purposes in the next year.

### **OMERACT-GRAPPA Outcome Measure Project**

After updating the core domain set for PsA in 2016, the GRAPPA-OMERACT Outcome Measure Working Group has continued its work to develop the core outcome measurement set. The presenters (Drs. Ying Ying Leung, Ana-Maria Orbai, William Tillett, and Dafna Gladman) reminded attendees that at OMERACT 2018, the 66/68 swollen and tender joint count was endorsed to measure musculoskeletal (MSK) disease activity-peripheral arthritis, and the PsA Impact of Disease 12-item questionnaire (PsAID12) was provisionally endorsed to evaluate health-related quality of life. The working group subsequently prioritized the determination of core measures for 4 additional domains: MSK disease activity-enthesitis, physical function, fatigue, and structural damage. Workstreams for each domain are led by a GRAPPA-OMERACT steering committee member and include at least 2 PRPs. Identified instruments that assess each domain will be appraised using the OMERACT Filter 2.1, a set of standards by which to evaluate each instrument for truth, discrimination, and feasibility. Instruments will be proposed for the core outcome measurement set based on the available evidence and will be presented for endorsement by OMERACT. A new endorsement process from OMERACT is planned that will allow approvals through a web-based portal in real time rather than through the biennial face-to-face OMERACT congress.

#### **Axial Project**

Drs. Dafna Gladman and Denis Poddubnyy presented the ongoing Assessment of SpA international Society (ASAS)-GRAPPA axial project, which aims to develop a consensus definition of axial involvement in PsA. From December 2018 through January 2019, a web-based survey was conducted among ASAS and GRAPPA members that aimed to identify the parameters relevant to the definition of axial involvement in PsA using an established decision-making algorithm (Potentially All Pairwise RanKings of all possible Alternatives [PAPRIKA] method with practical implementation through the 1000minds web application). The survey was completed by 186 ASAS and GRAPPA members (106 ASAS members and 123 GRAPPA members, with 43 members belonging to both societies).

According to the survey results, the following 4 parameters that reflect objective signs of inflammatory involvement of the axial skeleton received the highest ranking: (1) presence of structural damage on a radiograph (X-ray) of the sacroiliac joints (radiographic sacroiliitis of at least grade II bilaterally or grade III unilaterally); (2) presence of structural damage on an X-ray of the spine; (3) presence of subchondral bone marrow edema/osteitis on an MRI of the sacroiliac

Page **12** of **14** 

joints compatible with SpA; and (4) presence of bone marrow edema/osteitis on MRI of the spine compatible with SpA. These parameters represented the most likely candidates for the future consensus definition. Currently, a prospective study is planned in PsA with comprehensive imaging of the axial skeleton to generate data to validate the expected definition.

### **Ultrasound Enthesitis Project**

Dr. Lihi Eder presented an update on the GRAPPA Diagnostic Ultrasound Enthesitis Tool (DUET) project. The objective of this GRAPPA initiative is to develop a sonographic tool to enable the early diagnosis of PsA, especially as sonography is more sensitive and specific than clinical examination in detecting enthesitis. From pilot studies, a candidate set of entheses was selected. Utilizing these entheses, a larger study is planned within the GRAPPA membership to ensure a diverse population is evaluated to test this potential diagnostic tool. The protocol has been reviewed by the steering committee, and training materials (videos, acquisition atlas) are being developed. A post-meeting workshop was held to understand potential barriers to study conduct in various countries. The possibility of performing local reading in addition to the planned central reading was also explored. Members from Europe, Israel, India, as well as North America were present. Challenges within each system were discussed, as well as potential solutions. Timelines to seek study funding were clarified, with the planned study start in mid-2020.

### Early Psoriatic Disease Systematic Literature Review

Dr. Gabriele De Marco reported on the set up of a systematic literature review project entitled "Non-topical pharmacological treatment of early, untreated (disease modifying antirheumatic drug [DMARD]-naïve, systemic therapy-naïve) psoriatic disease: a systematic review." Most trials have been limited with a focus on severe disease of the skin or joints, without looking at more comprehensive indices in outcomes assessment. The target condition was defined so that clinical studies investigating (1) MSK and/or cutaneous manifestations; or (2) untreated psoriatic disease in its early stage could be included. Case reports or studies assessing the effects of topical therapies were excluded. The range of outcome measures considered was broad in order to assess across the spectrum of disease manifestations and to maximize the sensitivity of the search strategy. The resources accessed to find the available evidence covered a time range from 1946 to May 2019 and spanned across electronic databases, trial registers, and conferences proceedings. The flow diagram presented during the talk highlighted the substantial number of records (144,299) that were identified through electronic database searches. The diagram also provided the status of the ongoing selection activities performed by the team and the timeline for publication.

# Conclusion

This paper summarizes GRAPPA's recent work on several projects. These projects are part of GRAPPA's ongoing mission to address educational and unmet research needs for psoriatic disease, to create opportunities for networking within the psoriatic community, and to optimize patient care through collaborative care networks and treatment recommendations.