902 Friday, 14 June 2019 Scientific Abstracts

FRI0425

LITERATURE REVIEW OF PATIENT PERSPECTIVES ON THE MANAGEMENT AND TREATMENT OF PSORIATIC ARTHRITIS

Annelies Boonen¹, Neil Betteridge², Andreas Pinter³, Julie Hill⁴, Savita Anand⁵, Colleen Mchorney⁶, Catherine Reed⁴. ¹Maastrict University, Maastrict, Netherlands; ²Neil Betteridge Associates, London, United Kingdom; ³University Hospital Frankfurt, Frankfurt, Germany; ⁴Eli Lilly, Windlesham, United Kingdom; ⁵Evidera, London, United Kingdom; ⁶Evidera, Bethesda, United States of America

Background: A patient-centred approach to the management and drug treatment of psoriatic arthritis (PsA) has been advocated by a multidisciplinary group of experts to improve skin and joint symptoms and health-related quality of life (HRQoL).¹

Objectives: To examine perspectives of patients with PsA on: (1) disease management and treatment goals, (2) disease management and treatment satisfaction, and (3) treatment adherence, including reasons for discontinuation. Areas of interest were those related to medication, symptom resolution, everyday living and overall HRQoL.

Methods: A targeted literature review was conducted to identify peer-reviewed literature on patient experience with PsA management and drugtreatment. English-language articles published between 1 January 2010–8 October 2018 reporting qualitative or quantitative evidence from cross-sectional or longitudinal observational studies were identified from searches conducted using MEDLINE (via PubMed) and Embase. Selection criteria included adult patients with PsA (self-reported or clinician-diagnosed); drug-treatment studies could consider only regulatory-approved treatments for PsA and other studies had to provide evidence of patient perspectives on disease management and treatment goals, experiences and/or satisfaction. Studies involving paediatric/adolescent populations were excluded, as were results for PsA were not distinguishable from other diseases.

Results: The literature search identified 266 titles, of which 48 duplicates were removed. The remaining 218 abstracts were screened: 58 full-text articles were assessed for eligibility and 16 articles were selected for fulltext review. Of these 16 articles, 9 were primarily related to patient perspective on disease management, 6 to patient satisfaction and 1 to treatment adherence; some articles covered more than one of these objectives. None of the articles studied whether explicit consideration of treatment goals from the patient perspective would influence management or outcome of care. Symptom resolution, reduced fatigue, improved sexual relations, improved HRQoL and ability to participate in daily activities were consistently identified by patients as important aspects for disease management and daily living with PsA. Articles on patient satisfaction focused largely on general satisfaction with medication rather than satisfaction specifically with holistic PsA management. Notwithstanding, symptom resolution was clearly linked to greater patient satisfaction with medication. Patient dissatisfaction with PsA treatment was influenced by their attitude towards treatment, concerns about PsA medication, the physician-patient relationship and lack of patient involvement in decisionmaking. Treatment adherence has not been widely explored but mainly relates to perceptions about, and experiences with, medications, including efficacy and adverse events.

Conclusion: This literature review identified a lack of research on patient perspectives of PsA management and treatment goals. It also highlighted the lack of patient involvement in determining management and/or setting personal goals, which may ultimately affect satisfaction. There remains a lack of clarity on PsA symptoms and other disease- or patient-related parameters that impact patient satisfaction/dissatisfaction and patient-centred reasons for treatment discontinuation. The findings of this review will be used to develop a PsA patient survey to further explore patient perspectives to improve care in PsA.

REFERENCES:

[1] Betteridge N, et al. J Eur Acad Dermatol Venereol 2016; 30: 576-85.

Disclosure of Interests: Annelies Boonen: None declared, Neil Betteridge Consultant for: Amgen, Eli Lilly, Grunenthal, GSK, Heart Valve Voice, Janssen, Roche, Sanofi Genzyme and Sanofi Regeneron, Speakers bureau: Amgen, Eli Lilly, Grunenthal, GSK, Heart Valve Voice, Janssen, Roche, Sanofi Genzyme and Sanofi Regeneron, Andreas Pinter Consultant for: AbbVie, Almirall Hermal, Amgen, Biogen Idec, Boehringer Ingelheim, Celgene, GSK, Eli Lilly and Company, Galderma, Hexal, Janssen, LEO Pharma, Medac, Merck Serono, Mitsubishi, MSD, Novartis, Pfizering-Plough and UCB Pharma, Speakers bureau: AbbVie, Almirall Hermal, Amgen, Biogen Idec, Boehringer Ingelheim, Celgene, GSK, Eli Lilly and Company, Galderma, Hexal, Janssen, LEO Pharma, Medac, Merck Serono, Mitsubishi, MSD, Novartis, Pfizer, Tigercat Pharma, Regeneron,

Roche, Sandoz Biopharmaceuticals, Schering-Plough and UCB Pharma, Julie Hill Shareholder of: Eli Lilly and Company, Employee of: Eli Lilly and Company, Savita Anand Consultant for: Contracted by Eli Lilly and Company to conduct the literature review., Colleen McHorney Consultant for: Contracted by Eli Lilly and Company to conduct the literature review., Catherine Reed Shareholder of: Eli Lilly and Company, Employee of: Eli Lilly and Company

DOI: 10.1136/annrheumdis-2019-eular.5105

FRI0426

IXEKIZUMAB MAKES REMISSION AND LOW DISEASE ACTIVITY POSSIBLE IN PATIENTS WITH PSORIATIC ARTHRITIS: TWO-YEAR RESULTS IN TNF INADEQUATE RESPONDERS OR BIOLOGIC-NAÏVE PATIENTS

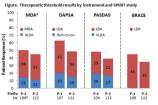
Laura C. Coates¹, Alexis Ogdie², Prashanth Sunkureddi³, Lisa Kerr⁴, Matthew Hufford⁴, Philip Helliwell⁵. ¹University of Oxford, Oxford, United Kingdom; ²Penn Medicine, Philadelphia, United States of America; ³Clear Lake Rheumatology, League City, United States of America; ⁴Eli Lilly and Company, Indianapolis, United States of America; ⁵Univ of Leeds, School of Medicine, Leeds, United Kingdom

Background: Psoriatic arthritis (PsA) is a heterogeneous inflammatory disease that can involve peripheral and axial joints, skin, and entheses. A number of validated composite indices have been developed, not only to measure overall disease activity for PsA, but also to provide thresholds for treat-to-target goals. These include MDA (Minimal Disease Activity), DAPSA (Disease Activity in PsA), and PASDAS (PsA Disease Activity) score). We have previously demonstrated that higher proportions of PsA patients treated with ixekizumab (IXE), a monoclonal antibody that selectivity targets interleukin-17A, achieved therapeutic thresholds defined by MDA, DAPSA, and PASDAS versus placebo (PBO) up to Week 24.

Objectives: To explore the extent to which IXE can help biologic-naïve and tumor necrosis factor inhibitor (TNFi) inadequate responder patients achieve treat-to-target goals, as defined by composite indices incorporating multiple disease domains, through 108 weeks of treatment.

Methods: Data were analyzed from all patients initially randomized to 80 mg IXE every 4 weeks after a 160-mg starting dose in 2 double-blind, PBO-controlled phase III trials investigating the efficacy and safety of IXE. For SPIRIT-P1 (NCT01695239), patients (N=107) were bDMARD naïve. For SPIRIT-P2 (NCT02349295), patients (N=122) had an inadequate response or were intolerant to TNFis. The following composite measures and definitions were used: MDA and Very Low Disease Activity (VLDA) (see Figure); DAPSA Low Disease Activity (VLDA) (≤14 and >4) and remission (≤4); PASDAS LDA/VLDA (≤3.2/≤1.9); and GRACE (GRAppa Composite score) LDA (≤2.3). Modified nonresponder imputation (mNRI; missing data treated as nonresponse for patients discontinued due to lack of efficacy or adverse events; multiple imputation for all other missing data) was used for all analyses.

Results: Therapeutic threshold results at Week 108 are summarized in the Figure. Whether measured using MDA, DAPSA, PASDAS, or GRACE, the proportions of IXE-treated patients achieving designated therapeutic thresholds were sustained through 2 years of treatment. Efficacy was similar between SPIRIT-P1 and SPIRIT-P2.



Data labels show proportion of patients achieving the indicated treatment threshold P-1 and P-2, SPRIT-2 and P-2, No. patients included in the nankyi (modified nonresponder imputation applied). MODA was achieved if 5 to 6, and VLDA was achieved if 7 of 7, of the following criteria were met-tender join count s1; shooting joint count s1; shortisis Area and Severity index total score s1 or body surface area 53%; patient's assessment of pain visual madies cale (VAI) 53; patient's global assessment of disease activity VAS 250; Health Assessment Classifications 10x46 s. 50.3 and tender enthease) point's Lassessed by the Lede Sintherist index 10x46 s. 50.3 and tender enthease plorist is Lassessed by the Lede Sintherist index 10x46 s. 50.3 and tender enthease plorist is Lassessed by the Lede Sintherist index 10x46 s. 50.3 and tender enthease plorist is Lassessed by the Lede Sintherist index 10x46 s. 50.3 and tender enthease plorist is Lassessed by the Lede Sintherist index 10x46 s. 50.3 and tender enthease plorist is set to the sintherist index 10x46 s. 50.3 and tender enthease plorist is set to the sintherist index 10x46 s. 50.3 and tender enthease plorist is set to the sintherist index 10x46 s. 50.3 and tender enthease plorist index

Conclusion: High proportions of IXE-treated patients, whether biologic naïve or TNFi inadequate responders, achieved treat-to-target goals, as defined by composite indices incorporating multiple disease domains, through 2 years of treatment.

REFERENCES:

[1] Coates L, et al. Ann Rheum Dis. 2018;77(Suppl):A375.