ORIGINAL RESEARCH

An 8-week multimodal treatment program improves symptoms of knee osteoarthritis: a real-world multicenter experience

Larry E Miller[1,2]
Jon E Block[2]

[1]Miller Scientific Consulting, Inc, Arden, NC, USA; [2]The Jon Block Group, San Francisco, CA, USA

Purpose: To report outcomes from a 5-year real-world clinical experience with a multimodal treatment program in patients with symptomatic knee osteoarthritis (OA).

Methods: Patients with symptomatic, radiographically confirmed knee OA resistant to traditional conservative treatments underwent a supervised 8-week multimodal treatment program consisting of low-impact aerobic exercise, muscle flexibility exercises, joint mobilization, physical therapy modalities, muscle strengthening and functional training, patient education, and a series of 3 or 5 weekly hyaluronic acid injections. Patients were evaluated at admission, 4 weeks, and 8 weeks. Patient-reported outcomes included knee pain severity using an 11-point (0–10) numerical scale and the Western Ontario and McMaster Universities Osteoarthritis Index.

Results: A total of 3,569 patients completed an 8-week treatment course between January 2008 and April 2013 at 66 dedicated treatment centers in the United States. Knee pain severity assessed on a numeric scale decreased 59% on average, from 5.4±2.9 to 2.2±2.2 ($P<0.001$). Western Ontario and McMaster Universities Osteoarthritis Index subscores decreased by 44% to 51% (all $P<0.001$) during the 8-week program. The percentage of patients achieving the threshold for Western Ontario and McMaster Universities Osteoarthritis Index minimally perceptible clinical improvement was 79% for the Pain subscale, 75% for Function, and 76% for Stiffness. Favorable patient outcomes were reported in all subgroups, regardless of age, sex, body mass index, disease severity, or number of treatment cycles.

Discussion: A real-world 8-week multimodal treatment program results in clinically meaningful improvements in knee OA symptoms, with excellent generalizability across a broad range of patient characteristics.

Keywords: hyaluronic acid, knee, multimodal, nonsurgical, osteoarthritis, physical therapy

Introduction

Osteoarthritis (OA) of the knee is the leading cause of disability in adults[1–3] and is characterized by progressive joint pain and dysfunction secondary to articular cartilage and subchondral bone damage, synovitis, osteophyte formation, and joint space loss.[4] The prevalence of OA is expected to increase by 40% by 2025,[5] with a projected economic burden of $128 billion per year in the United States.[6–8]

Nonsurgical treatments for knee OA commonly include weight loss, lateral wedge insoles, analgesics, bracing, physical therapy, and hyaluronic acid (HA) injections. However, the independent clinical benefit offered by these therapies is questionable,[9] and the use of a single modality to treat knee OA does not reflect typical clinical practice. Because the etiology of knee OA is multifactoral,[10] it is plausible that multimodal treatment regimens may be necessary to effectively assuage the symptoms

Correspondence: Jon E Block
The Jon Block Group, 2210 Jackson St, Suite 401, San Francisco, CA 94115, USA
Tel +1 415 775 7947
Fax +1 415 928 0765
Email jonblock@jonblockgroup.com

of this disease. Although multimodal nonsurgical therapy is widely advocated and unanimously supported in knee OA treatment position statements,[11] research in this area is mainly restricted to clinical trials with limited generalizability to the general population. Furthermore, the components of such multimodal programs are highly variable, and comparisons among interventions are difficult.

Intraarticular injection of HA enhances the viscoelastic properties of synovial fluid and inhibits proinflammatory pathways.[12] A recent meta-analysis of randomized controlled trials reported that HA products approved in the United States were safe and significantly improved knee pain and function in knee OA patients for up to 6 months.[13] Because a major determinant of knee OA progression is excessive and/or aberrant knee joint loading patterns,[14] physical rehabilitation programs aimed at improving muscle strength, flexibility, and proprioception may serve to lower knee joint loading forces and decrease OA symptom severity. The purpose of this article is to report outcomes from a 5-year real-world clinical experience with a multimodal treatment program in patients with symptomatic knee OA.

Methods
Patients
Patients with symptomatic, radiographically confirmed knee OA resistant to traditional conservative therapies were enrolled from 66 dedicated treatment centers (affiliated with OsteoArthritis Centers of America[SM]) in 25 states across the United States. The percentage of patients enrolled according to city size was 6% for cities with a population smaller than 10,000, 57% for a population of 10,000 to 99,999, 29% for a population of 100,000 to 999,999, and 7% for a population of 1,000,000 or more.

Baseline assessments included medical history and a complete clinical and orthopedic examination. Standing weight-bearing X-rays confirmed the OA diagnosis, and disease severity was classified using the Kellgren-Lawrence (K-L) grading scale[15] (0, normal; I, possible osteophyte, no joint space narrowing; II, definite osteophyte, possible joint space narrowing; III, multiple osteophytes, definite joint space narrowing, sclerosis, and possible deformity of bone ends; and IV, large osteophytes, marked joint space narrowing, severe sclerosis, and definite deformity of bone ends). Patients with a K-L grade of 0 were not considered for this treatment program. A diagnostic arthrogram was performed to rule out contraindications (eg, meniscal tear, ligamentous instability) and to ensure potential for tri-compartmental HA flow by confirming the absence of

potentially obstructive osteophytes, particularly in patients with grade IV OA.

Multimodal treatment program
Patients underwent an 8-week proprietary multimodal treatment program consisting of low-impact aerobic exercise, muscle flexibility exercises, joint mobilization, physical therapy modalities, muscle strengthening and functional training, patient education, and a series of weekly HA injections. The combinatorial program components were performed two to three times per week and divided into four distinct 2-week phases. The initial goals of the program were to reduce knee pain, encourage consistent aerobic exercise, improve muscle flexibility and joint range of motion, increase circulation in the knee joint, and reduce knee joint loading forces with a knee brace and/or wedged insoles. A typical phase 1 session began with low-impact aerobic exercise of approximately 10 minutes' duration. Muscle flexibility exercises emphasized the hamstrings, quadriceps, gastroc/soleus complex, hip flexors, and hip rotators. Joint mobilization techniques were used to improve range of motion at the patellofemoral and tibiofemoral joints, as well as the lumbar spine and hip. Therapeutic modalities such as vasopneumatic compression, electrical stimulation, cold laser, and/or ice were administered as needed. Muscle strengthening exercises were prescribed and supervised by physical therapists and included closed- and open-chain activities with a focus on knee flexors/extensors, hip abductors and rotators, ankle inverters/everters, and trunk rotators. As the program advanced, aerobic exercise duration increased from 10 to 20 or more minutes. Muscle flexibility exercises and joint mobilization techniques continued. Muscle strengthening exercise intensity progressively increased from slow-speed movements at 60% of a one-repetition maximum to faster speed movements at 75%–80% of a one-repetition maximum, as tolerated. Balance, proprioception, and closed-chain functional movements bearing partial body weight were gradually introduced. At the end of the 8-week combinatorial program, patients were encouraged to engage in daily low-impact aerobic activity and functional exercises outside of the program.

Patients underwent 1 week of physical therapy to establish baseline knee pain and function scores before undergoing a series of weekly HA injections for 3 or 5 weeks, depending on the type of HA product. HA injections were administered under fluoroscopic guidance to improve injection accuracy.[16] Specifically, before each HA injection, 1 cc contrast medium was injected in the joint space and tricompartmental flow was confirmed. The needle was always inserted into

the joint capsule on the side of maximal pain. The most commonly used HA products in this series were Hyalgan (Fidia Farmaceutici, Abano Terme, Italy; 57%) and Supartz (Bioventus LLC, Durham, NC, USA; 43%), with Euflexxa (Ferring Pharmaceuticals Inc, Parsippany, NJ, USA) and Synvisc (Sanofi, Bridgewater, NJ, USA) rarely used (each <1%).

In accordance with most United States payer guidelines for HA injections in knee OA, patients were eligible to undergo multiple 8-week treatment programs if the initial treatment course provided significant pain relief, the patient experienced recurring pain after therapy was discontinued, and a minimum of 6 months had passed between treatment courses.

Outcomes

Patients were evaluated at admission, 4 weeks, and 8 weeks. Patient-reported outcomes included knee pain severity using an 11-point (0 to 10) numerical scale and the Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) version 3.1.[17] The WOMAC is a validated tool that measures pain and dysfunction associated with OA of the lower extremities by assessing 5 pain-related items, 17 functional items, and 2 stiffness items. Each item is based on recall during the previous 48 hours and is scored on a 0–4 scale on which 0 represents none and 4 represents extreme. All WOMAC scores were normalized to a 0–100 scale, with a higher score representing a worse outcome.

Data analysis

Continuous data were reported as mean ± standard deviation, and categorical data were reported as frequencies and percentages. In patients with bilateral disease, the knee with the highest pain severity on the numeric scale was selected for analysis purposes. Longitudinal changes in clinical outcomes were assessed with repeated measures analysis of variance. We defined the minimum perceptible clinical improvement (MPCI) as an improvement from a baseline of 9.7 points or higher for WOMAC Pain, 9.3 points or higher for WOMAC Function, and 10.0 points or higher for WOMAC Stiffness scores.[18] Stepwise forward logistic regression was used to identify factors that predicted patients who achieved the MPCI for each WOMAC subscale. Univariate predictors with a P-value lower than 0.10 were included in the multivariate analysis. A P-value lower than 0.05 was considered statistically significant. Data were analyzed using Predictive Analytics Software (version 18; IBM Corporation, Armonk, NY, USA).

Results

A total of 3,569 patients completed an 8-week treatment course between January 2008 and April 2013. Baseline patient characteristics are detailed in Table 1. Patients were predominantly (55%) women and elderly (mean age, 70 years), with a mean body mass index of 31 kg/m^2, and suffered from moderate knee OA pain and dysfunction with a severity comparable to patients undergoing total knee arthroplasty.[19,20]

Knee pain severity assessed on a numeric scale decreased 59% on average during the 8-week treatment period, from 5.4±2.9 at pretreatment to 2.2±2.2 after the treatment course (P<0.001; Figure 1). The percentage of patients achieving improvements of 10%, 30%, and 50% compared with baseline values were 87%, 79%, and 70%, respectively.

WOMAC subscores decreased by 44%–51% (all P<0.001; Figure 2). The mean (±95% confidence interval [CI]) absolute improvement from baseline was 25±1 for Pain, 24±1 for Function, and 24±1 for Stiffness. The percentage of patients demonstrating any improvement in WOMAC scores was 85% for Pain, 87% for Function, and 76% for Stiffness. The percentage of patients achieving the WOMAC MPCI was 79% for Pain, 75% for Function, and 76% for Stiffness.

Similarly favorable patient outcomes were identified regardless of sex, age, body mass index, or K-L grade, with the percentage of patients achieving the MPCI ranging from

Table 1 Baseline patient characteristics

Variable	Value (N=3,569)
Female sex (n=3,554), n (%)	1,948 (55)
Age, mean ± standard deviation, years	70±11
Weight, mean ± standard deviation, kg	89±20
Body mass index, mean ± standard deviation, kg/m^2	31±7
Disease involvement, n (%)	
Bilateral	2,593 (73)
Unilateral	976 (27)
Kellgren-Lawrence grade (n=2,928), n (%)	
I	112 (4)
II	453 (15)
III	1,279 (44)
IV	1,084 (37)
Knee Pain Severity, 0–10 scale, mean ± standard deviation	5.4±2.9
WOMAC Pain, 0–100 scale, mean ± standard deviation	49±20
WOMAC Function, 0–100 scale, mean ± standard deviation	50±20
WOMAC Stiffness, 0–100 scale, mean ± standard deviation	55±24

Abbreviation: WOMAC, Western Ontario and McMaster Universities Osteoarthritis Index.
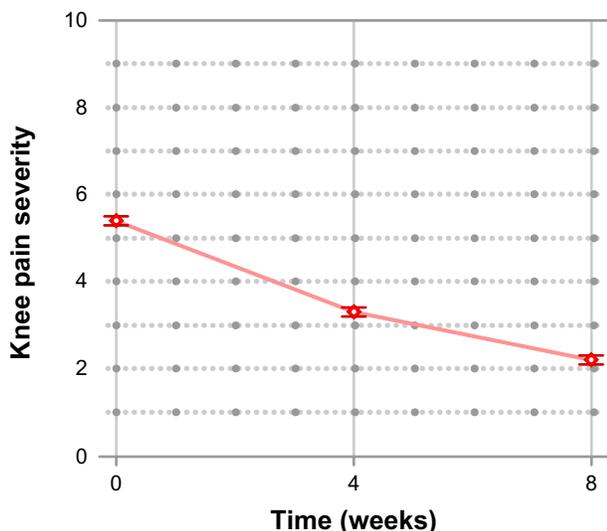
Figure 1 Improvement in knee pain severity using a numeric scale after a multimodal knee osteoarthritis program.

Table 2 Percentage of patients achieving the WOMAC minimal perceptible clinical improvement

Variable	Pain	Function	Stiffness
Sex			
Male	76	73	72
Female	80	77	78
Age, years			
<65	81	78	78
65–74	78	75	77
≥75	74	70	70
Body mass index, kg/m²			
≥30 kg/m²	79	76	76
<30 kg/m²	79	75	76
Kellgren-Lawrence grade			
I	79	76	76
II	80	77	76
III	79	76	76
IV	76	73	73

Abbreviation: WOMAC, Western Ontario and McMaster Universities Osteoarthritis Index.

70% to 81%, depending on subgroup and WOMAC subscore (Table 2). Longitudinal changes in WOMAC Pain scores by K-L grade are shown in Figure 3. Importantly, mean improvement in WOMAC Pain was 53%–55% in patients with K-L grade I–III and 46% in patients with K-L grade IV. Similar improvements across K-L grades were observed for WOMAC Function and Stiffness scores (data not shown).

Data were available for 158 patients who underwent a subsequent 8-week treatment cycle. Treatment outcomes in these patients remained favorable, although marginally inferior to those observed in the initial cycle, including a mean improvement of 51% in knee pain severity and MPCI achievement rates of 71% for Pain, 59% for Function, and 68% for Stiffness.
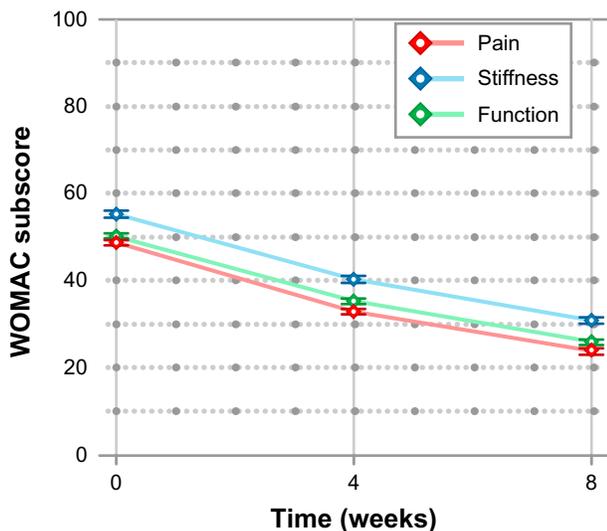
In multivariable logistic regression, younger age (per 1 year of age; odds ratio [OR], 0.99; 95% CI, 0.98–1.00) and female sex (OR, 1.32; 95% CI, 1.08–1.61) predicted Pain MPCI achievement, younger age (per 1 year of age; OR, 0.99; 95% CI, 0.98–1.00) predicted Function MPCI achievement, and younger age (per 1 year of age; OR, 0.99; 95% CI, 0.98–1.00) and female sex (OR, 1.48; 95% CI, 1.23-1.80) predicted Stiffness MPCI achievement. Body mass index and K-L grade had no independent influence on clinical outcomes.

No injuries related to the rehabilitation program necessitating program withdrawal or serious adverse events related to the HA injections were reported in any patient during this 5-year clinical experience.
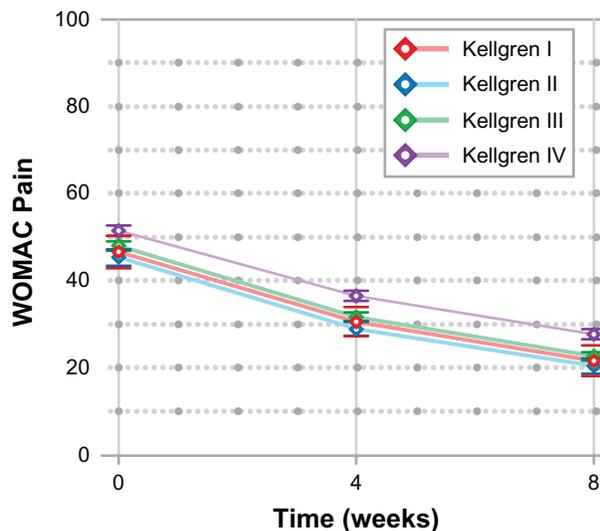
Figure 2 Improvement in Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) subscores after a multimodal knee osteoarthritis program.

Figure 3 Improvements in Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) pain scores by Kellgren-Lawrence classification.

Discussion

Knee OA is a prevalent musculoskeletal condition in older adults. Despite numerous available treatment options, none demonstrate clinically meaningful symptom amelioration and acceptable patient participation rates.[9,21] A 5-year real-world experience with more than 3,500 patients demonstrates the effectiveness of a proprietary multimodal knee OA program that incorporates low-impact aerobic exercise, muscle flexibility exercises, joint mobilization, physical therapy modalities, muscle strengthening and functional training, patient education, and a series of HA injections. These findings are important, as there are few satisfactory treatment options available to the OA patient, particularly in those with advanced disease.[21] Isolated conservative treatments for knee OA are not clinically effective for pain or function,[22] whereas surgical intervention such as arthroplasty is clinically effective, although only 1 in 5 patients are willing to undergo this invasive procedure.[21]

Interestingly, about 75% of the patients in the current series presented with K-L grade III or IV, and as a whole, patients experienced symptom severity levels at program entry similar to patients who undergo unicompartmental or total knee arthroplasty.[19,20] We observed short-term improvements in WOMAC subscores of 44%–51%, which are comparable to the magnitude of improvement in patients who undergo unicompartmental or total knee arthroplasty (range, 43%–73%).[19,23] These data suggest that even patients with advanced knee OA have a favorable prognosis with a multimodal treatment program without resorting to surgical interventions. It is plausible that if efficacy were maintained over the course of several treatment courses, arthroplasty might be delayed in patients with end-stage disease. Some patients in this program have undergone three or more treatment cycles, each separated by at least 6 months, but the sample size was too small to conduct a meaningful statistical analysis on this subset. Additional research is warranted to identify the subset of therapies used in this research that is responsible for eliciting the majority of the favorable treatment effect.

The strengths of this research are that a large number of patients were included in the analysis and that, as patients were treated as part of real-world clinical practice as opposed to a clinical trial, the external validity of these findings is noteworthy. Limitations of this research included lack of outcome reporting beyond the 8-week treatment period and the potential for placebo effect bias, as there were no untreated patients to serve as a comparator group.

Overall, an 8-week multimodal treatment program results in clinically meaningful improvements in knee OA symptoms, with excellent generalizability across a broad range of patient characteristics, including those with advanced disease.

Disclosure

References

1. Felson DT, Naimark A, Anderson J, Kazis L, Castelli W, Meenan RF. The prevalence of knee osteoarthritis in the elderly. The Framingham Osteoarthritis Study. *Arthritis Rheum*. 1987;30(8):914–918.
2. van Saase JL, van Romunde LK, Cats A, Vandenbroucke JP, Valkenburg HA. Epidemiology of osteoarthritis: Zoetermeer survey. Comparison of radiological osteoarthritis in a Dutch population with that in 10 other populations. *Ann Rheum Dis*. 1989;48(4):271–280.
3. Peat G, McCarney R, Croft P. Knee pain and osteoarthritis in older adults: a review of community burden and current use of primary health care. *Ann Rheum Dis*. 2001;60(2):91–97.
4. Dieppe PA, Lohmander LS. Pathogenesis and management of pain in osteoarthritis. *Lancet*. 2005;365(9463):965–973.
5. Woolf AD, Pfleger B. Burden of major musculoskeletal conditions. *Bull World Health Organ*. 2008;1(9):646–656.
6. Buckwalter JA, Saltzman C, Brown T. The impact of osteoarthritis: implications for research. *Clin Orthop Relat Res*. 2004;(Suppl 427): S6–S15.
7. Jackson DW, Simon TM, Aberman HM. Symptomatic articular cartilage degeneration: the impact in the new millennium. *Clin Orthop Relat Res*. 2001;(Suppl 391):S14–S25.
8. Samson DJ, Grant MD, Ratko TA, Bonnell CJ, Ziegler KM, Aronson N. Treatment of primary and secondary osteoarthritis of the knee. *Evid Rep Technol Assess (Full Rep)*. 2007;(157):1–157.
9. Crawford DC, Miller LE, Block JE. Conservative management of symptomatic knee osteoarthritis: a flawed strategy? *Orthop Rev (Pavia)*. 2013;5(1):e2.
10. Johnson VL, Giuffre BM, Hunter DJ. Osteoarthritis: what does imaging tell us about its etiology? *Semin Musculoskelet Radiol*. 2012; 16(5):410–418.
11. Zhang W, Moskowitz RW, Nuki G, et al. OARSI recommendations for the management of hip and knee osteoarthritis, part I: critical appraisal of existing treatment guidelines and systematic review of current research evidence. *Osteoarthritis Cartilage*. 2007;15(9): 981–1000.
12. Dahl LB, Dahl IM, Engström-Laurent A, Granath K. Concentration and molecular weight of sodium hyaluronate in synovial fluid from patients with rheumatoid arthritis and other arthropathies. *Ann Rheum Dis*. 1985;44(12):817–822.
13. Miller LE, Block JE. US-Approved Intra-Articular Hyaluronic Acid Injections are Safe and Effective in Patients with Knee Osteoarthritis: Systematic Review and Meta-Analysis of Randomized, Saline-Controlled Trials. *Clin Med Insights Arthritis Musculoskelet Disord*. 2013;6:57–63.
14. Waller C, Hayes D, Block JE, London NJ. Unload it: the key to the treatment of knee osteoarthritis. *Knee Surg Sports Traumatol Arthrosc*. 2011;19(11):1823–1829.
15. Kellgren JH, Lawrence JS. Radiological assessment of osteo-arthrosis. *Ann Rheum Dis*. 1957;16(4):494–502.
16. Berkoff DJ, Miller LE, Block JE. Clinical utility of ultrasound guidance for intra-articular knee injections: a review. *Clin Interv Aging*. 2012;7:89–95.

17. Bellamy N, Buchanan WW, Goldsmith CH, Campbell J, Stitt LW. Validation study of WOMAC: a health status instrument for measuring clinically important patient relevant outcomes to antirheumatic drug therapy in patients with osteoarthritis of the hip or knee. *J Rheumatol*. 1988;15(12):1833–1840.

18. Ehrich EW, Davies GM, Watson DJ, Bolognese JA, Seidenberg BC, Bellamy N. Minimal perceptible clinical improvement with the Western Ontario and McMaster Universities osteoarthritis index questionnaire and global assessments in patients with osteoarthritis. *J Rheumatol*. 2000;27(11):2635–2641.

19. Bachmeier CJ, March LM, Cross MJ, et al; Arthritis Cost and Outcome Project Group. A comparison of outcomes in osteoarthritis patients undergoing total hip and knee replacement surgery. *Osteoarthritis Cartilage*. 2001;9(2):137–146.

20. Becker R, Döring C, Denecke A, Brosz M. Expectation, satisfaction and clinical outcome of patients after total knee arthroplasty. *Knee Surg Sports Traumatol Arthrosc*. 2011;19(9):1433–1441.

21. London NJ, Miller LE, Block JE. Clinical and economic consequences of the treatment gap in knee osteoarthritis management. *Med Hypotheses*. 2011;76(6):887–892.

22. Crawford DC, Miller LE, Block JE. Conservative management of symptomatic knee osteoarthritis: a flawed strategy? *Orthop Rev (Pavia)*. 2013;5(1):e2.

23. Lyons MC, MacDonald SJ, Somerville LE, Naudie DD, McCalden RW. Unicompartmental versus total knee arthroplasty database analysis: is there a winner? *Clin Orthop Relat Res*. 2012;470(1):84–90.