Efficacy and tolerability of celecoxib and naproxen versus placebo in Hispanic patients with knee osteoarthritis

Margaret Noyes Essex Regina Behar Michael A O'Connell Pritha Bhadra Brown

Pfizer Inc, New York, NY, USA

Background: Celecoxib is an effective treatment for osteoarthritis (OA). However, information on its efficacy and safety profile in different racial/ethnic groups is limited. Noticeable differences among racial groups are found in other disease states, but a thorough investigation of OA is lacking. The objective of this study was to determine if celecoxib 200 mg once daily is as effective as naproxen 500 mg twice daily in the treatment of OA of the knee in Hispanic

Methods: Hispanic patients aged ≥45 years with knee OA were randomized to receive celecoxib 200 mg once daily, naproxen 500 mg twice daily, or placebo for 6 weeks. The primary efficacy variable was the change in Patient's Assessment of Arthritis Pain at 6 weeks compared with baseline. Secondary variables were change in Patient's and Physician's Global Assessments of Arthritis from baseline to week 6/early termination, change in Western Ontario and McMaster Universities OA Index (WOMAC) from baseline to week 6/early termination, change in American Pain Society pain score, Pain Satisfaction Scale, Patient Health Questionnaire (PHQ-9), and measurements of upper gastrointestinal tolerability.

Results: In total, 239 patients completed the trial (96 celecoxib, 96 naproxen, 47 placebo). Celecoxib was as effective as naproxen in reducing OA pain (least squares mean change from baseline [standard error] -39.7 [2.7] for celecoxib and -36.9 [2.6] for naproxen). Patient's and Physician's Global Assessments of Arthritis, WOMAC scores, upper gastrointestinal tolerability, Pain Satisfaction Scale, and PHQ-9 showed no statistically significant differences between the celecoxib and naproxen groups. The incidence of adverse events and treatment-related adverse events were similar among the treatment groups.

Conclusion: Celecoxib 200 mg once daily was as effective as naproxen 500 mg twice daily in the treatment of signs and symptoms of knee OA in Hispanic patients. Celecoxib was shown to be safe and well tolerated in this patient population.

Keywords: nonsteroidal anti-inflammatory drugs, cyclo-oxygenase-2, race, ethnicity

Introduction

It is predicted that nearly one in three US residents will be Hispanic by 2060. The exact prevalence of osteoarthritis (OA) in the Hispanic population is unknown, but research suggests that between 12% and 22% of Hispanics have arthritis, of which OA is the most common form.^{2,3} The prevalence is lowest (12%) in Cubans/Cuban Americans and highest (22%) in Puerto Ricans.³ Research also suggests that severe pain, work impairment, and poor outcomes are higher in this population compared with Caucasians.⁴ Older Hispanics with OA of the knee and obesity have an unduly elevated loss of quality-adjusted life-years.⁵ In addition, a negative impact on the

Correspondence: Margaret Noyes Essex Medical Affairs, Pfizer Inc, 235 East 42nd Street, New York, NY 10017, USA Tel +I 2I2 733 8018 Email margaret.essex@pfizer.com

how to request permission may be found at: http://www.dovepress.com/permissions.php

perception of quality of life has been reported in association with pain severity, age, and poor socioeconomic status in Venezuelan patients with OA.⁶ This suggests that Hispanic patients with OA could benefit from early and effective medical intervention.

Limited information is available on how Hispanic patients with OA compare with other ethnic groups in their OA treatment approaches and responsiveness. The literature indicates that, compared with Caucasians, Hispanic patients are less likely to undergo total knee replacement and are more likely to use oral herbs and magnets/copper jewelry therapy. Another study has shown that these patients are less likely to receive treatment with a cyclo-oxygenase-2 selective non-steroidal anti-inflammatory drug (NSAID; odds ratio 0.47, P<0.01) and more likely to discontinue treatment early. 13

Celecoxib is a selective NSAID indicated for the treatment of signs and symptoms associated with OA.¹⁴ Its efficacy has been established, and it has a favorable gastrointestinal tolerability profile relative to nonselective NSAIDs, ie, fewer patients report gastrointestinal adverse events, such as dyspepsia, with celecoxib.¹⁵ The purpose of this study was to confirm the noninferiority of celecoxib to naproxen, a nonselective NSAID, with regard to analgesic effects and gastrointestinal tolerability in patients with OA who were of Hispanic descent.

Materials and methods

Objectives

The primary objective of this study was to determine whether celecoxib 200 mg once daily was as effective as naproxen 500 mg twice daily for the treatment of symptoms associated with OA of the knee in a Hispanic population. The secondary objective was to confirm the tolerability of celecoxib 200 mg once daily versus placebo in these patients. The use of complementary and alternative medicines in this population was also evaluated at baseline.

Study population

Patients aged ≥45 years and of self-reported Hispanic descent with OA of the knee (diagnosed according to American College of Rheumatology criteria¹⁶) who were determined to be in a flare state and had a functional capacity classification of I to III met the study eligibility criteria. Other inclusion criteria that applied have been described in a previously published report that assessed the response to NSAIDS in an African American population with OA.¹⁷ Briefly, patients actively being treated with an NSAID or other analgesic therapy discontinued treatment at least 48 hours prior to the

baseline assessments. Eligible patients indicated a Patient's Assessment of Arthritis Pain visual analog scale (VAS) score between 40 mm and 90 mm (range 0–100 mm) and had a minimum rating of 3 on the Physician's and Patient's Global Assessment of Arthritis at baseline. Exclusion criteria were the same as those described in the previous report.¹⁷

Study design

This was a 6-week, randomized, double-blind, placebocontrolled, active-comparator, parallel-group trial carried out in 31 US centers in compliance with the principles of Good Clinical Practice and the Declaration of Helsinki. Each study site received protocol approval from an institutional review board, and all patients gave written informed consent.

Four clinic visits were required, ie, at screening, baseline, week 2, and week 6. During the screening visit, patients underwent a physical examination and laboratory tests. Both the patient and physician provided an assessment of arthritis. Patients were randomized in a 2:2:1 ratio to one of three treatments, ie, celecoxib 200 mg once daily, naproxen 500 mg twice daily, or placebo, according to a predetermined computer randomization schedule. Patients were assigned their randomization number based on the order in which they enrolled in the study. Both the investigator and patient were blinded to the study medications and to placebo, and all assessments were made by individuals who had been blinded. Each study medication had a matching placebo that was of similar appearance (capsule size, color, smell, and taste).

Efficacy evaluation

The primary efficacy outcome was defined as the change from baseline to week 6 in the Patient's Assessment of Arthritis Pain, which was measured on a VAS of 0 mm (no pain) to 100 mm (worst pain). All pain assessments were based on the one knee selected by the patient to be the "index joint". Secondary outcomes included change in Patient's and Physician's Global Assessments of Arthritis and Western Ontario and McMaster Universities OA Index (WOMAC) from baseline to week 6, change in American Pain Society (APS) pain scores from baseline to day 7 (week 1), change in Pain Satisfaction Scale and Patient Health Questionnaire (PHQ-9) scores from screening to week 6, and measurement of upper gastrointestinal tolerability.

The population evaluable for efficacy was used for the primary efficacy analysis and the modified intent-to-treat population was used for secondary efficacy analyses. The modified intent-to-treat population included all patients who were randomized, received at least one dose of study

medication, and had at least one post-baseline follow-up efficacy measure. The efficacy evaluable population included modified intent-to-treat patients who had no major protocol violations, were assessed at both baseline and week 6 for the primary efficacy variable, had adequate treatment, and belonged to the protocol-specified ethnic group.

Safety evaluation

General clinical safety was assessed by monitoring treatmentemergent adverse events and serious adverse events and by physical examination. Upper gastrointestinal tolerability was assessed as described in a previously published study.¹⁷ Safety analyses were carried out in the safety population, which included all randomized patients who received at least one dose of study medication.

Statistical analysis

Sample size calculation was based on the maximum clinically acceptable difference for declaring noninferiority, which was compared with the lower bound of the two-sided 95% confidence interval (CI) for the difference between the two treatment groups. ¹⁷ A total of 120 patients per active treatment group were randomized to adjust for the differences between the intent-to-treat and efficacy evaluable populations to allow for any nonevaluable patients (eg, those who are lost to follow-up). Sixty patients were randomized to placebo in order to have 80% power to detect a difference of 15 mm between the active treatment group and placebo in the VAS score.

Change in VAS score from baseline to week 6 was analyzed using a general linear model with treatment and center effects in the model and baseline score as a covariate. Pairwise comparisons were conducted. Celecoxib was declared to be as effective as naproxen if the lower bound of the two-sided 95% CI of the treatment difference (naproxen—celecoxib) lay above—10 mm. ¹⁸ As a test of internal control, differences in the mean change in VAS score were also analyzed for celecoxib versus placebo and for naproxen versus placebo.

The 24-item WOMAC scale and subscales were analyzed using a general linear model with treatment and center effects in the model, and baseline WOMAC score as a covariate. The WOMAC total domain score (range 0–96) was the sum of the pain, stiffness, and physical function domain scores. Responses to the Patient's and Physician's Global Assessments of Arthritis were analyzed and the patients' conditions were classified as "improved", "no change", or "worsened" using the Cochran–Mantel–Haenszel test,

stratified by center. APS questions were analyzed using the Cochran–Mantel–Haenszel test stratified by center (question 1, "Have you experienced any pain in the past 24 hours?"; yes or no). Change from baseline for the remaining questions was analyzed using a general linear model with treatment, center, and baseline APS value (questions 2–5) as a covariate. Change in PHQ-9 score was analyzed using a general linear model with treatment, center, and screening PHQ-9 score as a covariate. Patient pain satisfaction was analyzed at screening and week 6 using the Cochran–Mantel–Haenszel test, stratified by center.

Safety and upper gastrointestinal events were analyzed in the safety population, defined as randomized patients receiving at least one dose of study medication. The incidence of upper gastrointestinal events was analyzed using two-tailed Fisher's exact tests.

Results

Patient disposition and baseline demographics

A total of 318 patients were randomized to treatment (127, 129, and 62 patients in the celecoxib, naproxen, and placebo groups, respectively), and 315 patients received treatment (Figure 1). Baseline characteristics of the patients were similar among the treatment groups, and are summarized in Table 1. Patients were mostly female (60%–72%), and with an age range of 40–88 years. Mean duration of OA ranged from 5.3 to 6.6 years. The majority of patients were assessed as "poor" or "very poor" at baseline on both the Patient's and Physician's Global Assessments of Arthritis, and were in the functional capacity classification of II and III. Mean WOMAC total domain scores ranged from 55.7 to 58.6.

There were no significant differences noted between groups with regard to responses to the Pain Satisfaction Scale at the screening visit except for question 8 ("have better relationships with others"), for which a statistically significant difference was seen in favor of the celecoxib group over the placebo group (P=0.015).

Responses to the Complementary and Alternative Medicines Questionnaire indicated that prescription medicines, self-determined over-the-counter medicines, and physician-recommended over-the-counter medicines were used by 69%, 45%, and 44% of the screened population, respectively (data not shown). "Store bought lotions, oils, and creams" (56%) were the most frequently used alternatives to conventional medical OA treatments, while other herbal, homemade, or household lotions or oils were used by fewer than 15% of individuals screened. Dietary modifications were also a

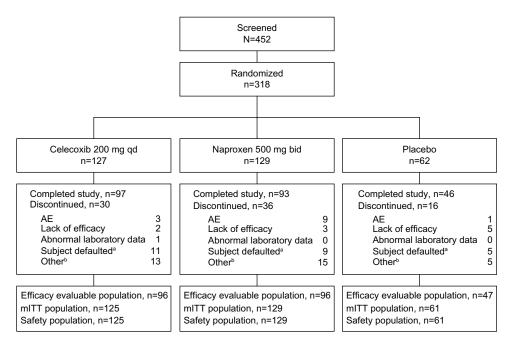


Figure I Patient disposition.

Notes: alncludes "lost to follow-up" and "patient no longer willing to participate"; bincludes "protocol violation".

Abbreviations: AE, adverse events; mITT, modified intent-to-treat; qd, once daily; bid, twice daily

common alternative to medical OA treatments; ≥50% of patients avoided alcohol and saturated fats or fried foods, 44% ate a high-fiber diet or whole grain foods, and 35%–38% avoided white flour, sugar, and/or red meats, and/or increased the amount of cabbage, broccoli, kale, and Brussels sprouts in their diets. There were also reports indicating the use of special vitamins, vitamin combinations, or minerals (27%) and glucosamine and/or chondroitin sulfate (23%). The use of other dietary supplements occurred in <12% of the screened individuals.

Further commonly-used alternative treatments included nutritional therapy (25%), massage (30%), and prayer (34%). Less than 18% of the population used herbal medicine, reflexology, acupuncture, chiropractic, and spiritual healing. Miscellaneous treatments such as venom, magnets, and biofeedback were used by <10% of the population.

Efficacy outcomes

Improvement in all three groups was seen on the primary efficacy outcome, ie, Patient's Assessment of Arthritis Pain (VAS, Table 2). Least squares mean changes from baseline to week 6 were –39.7 mm, –36.9 mm, and –28.6 mm in the celecoxib, naproxen, and placebo groups, respectively. The lower bound of the two-sided 95% CI of the treatment difference (naproxen – celecoxib) was above –10 mm (–3.8 mm). Hence, celecoxib was as effective as naproxen at reducing OA pain. Also, the *P*-values for celecoxib – placebo (*P*=0.0077)

suggest celecoxib is significantly more effective than placebo at relieving pain, as recorded on the Patient's Assessment of Arthritis Pain VAS.

The results for the secondary efficacy end points were similar between active treatments, supporting the noninferiority of celecoxib to naproxen. The outcomes of Patient's and Physician's Global Assessments of Arthritis are presented in Figures 2 and 3. Physicians described the arthritis condition of 60% of patients in the celecoxib group and 52% of patients in the naproxen group as "improved" by the week 6/early termination visit, compared with 46% of patients in the placebo group. Between-treatment differences were statistically significant in favor of celecoxib over placebo (P=0.0369). The OA condition of 57% of the patients in each active treatment group had "improved" by the week 6/early termination visit compared with 43% of patients in the placebo group. Between-treatment differences were not statistically significant (data not shown).

The mean change from baseline in the total and individual domain scores of the WOMAC indicated improvement from baseline in each treatment group (Table 3). Although differences between celecoxib and naproxen were not statistically significant, between-treatment differences for both celecoxib and naproxen compared with placebo were statistically significant for all but the stiffness domain.

For APS pain measurements, the number of patients who experienced pain within 24 hours prior to completion of the

Table I Baseline demographic and clinical characteristics

	Celecoxib	Naproxen	Placebo	P-value
	200 mg qd	500 mg bid	(n=62)	
	(n=127)	(n=129)		
Age, years,	59.6 (10.6)	60.5 (10.1)	61.7 (11.2)	0.396
mean (SD)				
(range)	(44-88)	(40–82)	(45–85)	
Sex, n (% female) ^a	92 (72)	82 (64)	37 (60)	0.166
Duration of OA,	5.3 (5.1)	6.4 (6.9)	6.6 (7.1)	0.193
years, mean (SD)				
Patient's Global Ass	sessment, n (%)	þ		
Very good	0	0	0	0.127
Good	l (<l)< td=""><td>I (<i)< td=""><td>0</td><td></td></i)<></td></l)<>	I (<i)< td=""><td>0</td><td></td></i)<>	0	
Fair	18 (14)	10 (8)	2 (3)	
Poor	93 (73)	98 (76)	52 (84)	
Very poor	15 (12)	20 (16)	8 (13)	
Physician's Global A	Assessment, n (%) ^b		
Very good	0	0	0	0.278
Good	I (<i)< td=""><td>I (<i)< td=""><td>0</td><td></td></i)<></td></i)<>	I (<i)< td=""><td>0</td><td></td></i)<>	0	
Fair	17 (13)	12 (9)	3 (5)	
Poor	103 (81)	109 (85)	56 (90)	
Very poor	6 (5)	7 (5)	3 (5)	
Functional capacity	classification, n	ı (%) ^b		
1	2 (2)	7 (5)	3 (5)	0.213
II	90 (71)	95 (74)	41 (68)	
III	34 (27)	27 (21)	16 (27)	
IV	0 `	0 ` ´	0 `	
VAS score, mm,	67.3 (13.0)	67.0 (12.6)	67.5 (11.5)	0.975
mean (SD)	()	, ,	(-/	
WOMAC total	57.4 (14.4)	55.7 (15.8)	58.6 (13.1)	0.335
score, mean (SD) ^c	` '	, ,	` /	

Notes: *Two patients were younger than the protocol-specified 45 years of age; a 40-year-old naproxen patient was discontinued from the study due to this protocol violation and a 44-year-old celecoxib patient completed the study; *percentages are calculated based on the number of randomized patients; *WOMAC total domain score is the sum of pain, stiffness, and physical function domain scores.

Abbreviations: qd, once daily; bid, twice daily; OA, osteoarthritis; SD, standard deviation; VAS, visual analog scale; WOMAC, Western Ontario and McMaster Universities OA Index.

questionnaire decreased from 100% at baseline to 91%, 84%, and 95% on day 7 for the celecoxib, naproxen, and placebo groups, respectively. In general, APS pain scores improved from baseline to day 7 (Table 3). There were statistically significant differences in favor of active treatments compared with placebo in total pain interference from baseline to day 7 (P<0.001).

Overall, a greater percentage of patients in the celecoxib and naproxen groups responded positively to the questions on the Pain Satisfaction Scale at the week 6/early termination visit compared with placebo. More of the patients using active treatment than those using placebo were happy with the duration and speed of pain relief and agreed or somewhat agreed that their study pain medication positively affected their physical health, outlook, ability to perform daily and leisure activities, independence, relationships, mood,

Table 2 Patient's assessment of arthritis pain (VAS) at week 6 (efficacy evaluable population)

	Celecoxib 200 mg qd (n=96)	Naproxen 500 mg bid (n=96)	Placebo (n=47)
Baseline, mean (SE)	65.5 (1.3)	66.8 (1.2)	69.7 (1.5)
Week 6, mean (SE)	24.3 (2.4)	25.2 (2.4)	35.4 (3.9)
Change from	-39.7 (2.7)	-36.9 (2.6)	-28.6 (3.6)
baseline, LSM (SE)			
	Naproxen –	Naproxen –	Celecoxib –
	Celecoxib	Placebo	Placebo
LSM (SE)	-2.8 (3.3)	-8.4 (4.1)	-11.1 (4.1)
95% CI	-3.8 to 9.3	-16.3 to -0.4	−19.3 to −3.0
P-value	0.4028	0.0407	0.0077

Abbreviations: CI, confidence interval; LSM, least squares mean; SE, standard error; VAS, visual analog scale; qd, once daily; bid, twice daily.

concentration, and ease of movement. Differences between celecoxib and naproxen were not statistically significant. Results on the PHQ-9 were similar between the celecoxib and naproxen groups (data not shown).

Safety outcomes

All 315 patients who received treatment were evaluated for adverse events. A total of 110 patients (28%–37% per treatment group) reported 157 adverse events. The incidence of adverse events was similar among the treatment groups, with most being mild to moderate in severity. Of the 157 adverse events, 70 were considered to be treatment-related (25, 36, and nine in the celecoxib, naproxen, and placebo groups, respectively). No deaths were reported. Treatment-related adverse events occurring in \geq 2% of patients are summarized in Table 4.

The majority of patients who reported adverse events complained of gastrointestinal system and psychiatric disorders. The most commonly occurring adverse events were depression, headache, abdominal pain, constipation, and dyspepsia. Of these, only depression occurred in >5% of the subject population: 11% of subjects overall reported depression (10%, 9%, and 20% in the celecoxib, naproxen, and placebo groups, respectively). There were few reports of upper gastrointestinal intolerability in this study. A total of eight patients (three in the celecoxib group, four in the naproxen group, and one in the placebo group) experienced an upper gastrointestinal event, defined as moderate or severe nausea, abdominal pain, and/or dyspepsia (Table 3). Between-treatment differences were not statistically significant (P=1.0000).

Thirteen patients in total discontinued the study as a result of adverse events, while ten were withdrawn because

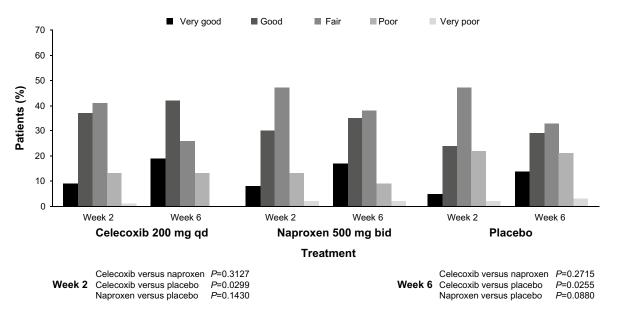


Figure 2 Patient's Global Assessment of Arthritis: modified intent-to-treat population. **Abbreviations:** qd, once daily; bid, twice daily.

of adverse events that were deemed to be treatment-related (two, seven, and one patients in the celecoxib, naproxen, and placebo groups, respectively). One patient in the naproxen group experienced a severe adverse event (gastrointestinal hemorrhage) that was considered related to the study treatment, resulting in discontinuation of the study medication.

Discussion

Despite the growing population of minority ethnic groups in the USA and a greater interest in their clinical experience of pain, non-white groups remain substantially underrepresented in clinical trials for a multitude of reasons, including mistrust of the health care system, sociocultural barriers, and a shortage of investigators of diverse ethnic backgrounds.^{19–23} This lack of representation poses a challenge with regard to the generalizability and external validity of clinical trial results and leaves a need for safety and efficacy data in minority groups.

Cultural and ethnic differences exist in the perception of pain and how it is treated from the perspectives of both patients and health care providers. In addition, the response to medication may differ in various ethnic populations because of genetic and metabolic factors.^{24,25} For these reasons, individual prescribers and payers are increasingly requesting efficacy and safety data for medications that have been studied in a greater variety of ethnic populations, so as

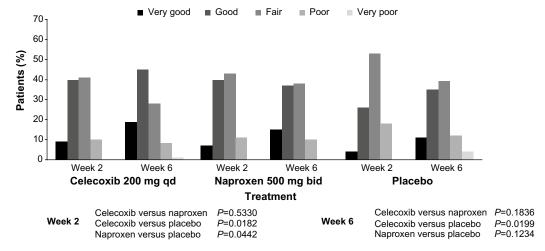


Figure 3 Physician's Global Assessment of Arthritis: overall ratings (modified intent-to-treat population). Abbreviations: qd, once daily; bid, twice daily.

Table 3 WOMAC, upper gastrointestinal tolerability, and APS pain scores

	Celecoxib 200 mg qd n=125	Naproxen 500 mg bid n=129	Placebo n=6 l	
Change from baseline in week 6/early termination WOMAC:				
mITT population				
Total				
LSM (SE)	-23.I (2.0)	-23.0 (I.9)	-16.0 (2.6)	
Pain				
LSM (SE)	-5.2 (0.4)	-5.I (0.4)	-4.0 (0.6)	
Stiffness				
LSM (SE)	-1.9 (0.2)	-1.9 (0.2)	-I.6 (0.2)	
Physical function				
LSM (SE)	-16.3 (1.4)	-16.0 (1.4)	-11.1 (1.9)	
	Celecoxib	Celecoxib	Naproxen	
	versus	versus	versus	
	naproxen	placebo	placebo	
P-values				
Total	0.9402	0.0232	0.0252	
Pain	0.9246	0.0089	0.0100	
Stiffness	0.9690	0.2173	0.2000	
Physical function	0.8857	0.0245	0.0307	
UGI tolerability during study (safety population)				

UGI event, n (%)	3 (2.4)	4 (3.1)	I (I.6)
	Celecoxib	Celecoxib	Naproxen
	versus	versus	versus
	naproxen	placebo	placebo
P-values	1.0	1.0	1.0

Mean change from baseline to day 7 in APS pain score – total pain interference (mITT population)

Baseline, n	124	128	61
Mean (SEM)	41.3 (1.30)	42.5 (1.21)	42.5 (1.71)
Day 7, n	107	96	55
Mean (SEM)	-16.0 (1.61)	-17.0 (1.60)	-8.3 (1.96)
	Celecoxib	Celecoxib	Naproxen
	Celecoxib versus	Celecoxib versus	Naproxen versus
			'

Abbreviations: APS, American Pain Society; LSM, least squares mean; mITT, modified intent-to-treat; SE, standard error; SEM, standard error of the mean; UGI, upper gastrointestinal; WOMAC, Western Ontario and McMaster Universities Osteoarthritis Index; bid, twice daily; qd, once daily.

to reflect the diversity of their beneficiary groups. The present study was conducted to further our understanding of the efficacy, safety, and tolerability of celecoxib in individuals of Hispanic descent.

In this population, mean changes in the Patient's Assessment of Arthritis Pain VAS improved in all three treatment groups. Given that the lower boundary of the two-sided 95% CI of the treatment difference (naproxen – celecoxib) was above –10 mm (–3.8 mm), celecoxib was as effective as naproxen at reducing OA pain. Compared with placebo, celecoxib was significantly more effective at relieving pain

Table 4 Treatment-related adverse events occurring in \geq 2% of patients (in decreasing order of occurrence)

	Celecoxib 200 mg qd (n=125)	Naproxen 500 mg bid (n=129)	Placebo (n=61)
AE by preferred term, n (%)			
Dyspepsia	4 (3)	5 (4)	I (2)
Depression	4 (3)	4 (3)	3 (5)
Abdominal pain	2 (2)	4 (3)	0

Abbreviations: AE, adverse event; bid, twice daily; qd, once daily.

(*P*=0.0077). Secondary efficacy findings were indicative of the noninferiority of celecoxib to naproxen, because similar results were seen on the Patient's and Physician's Global Assessment of Arthritis, WOMAC scores, Pain Satisfaction Scale, PHQ-9, and upper gastrointestinal tolerability. These findings are consistent with other studies showing comparable efficacy of celecoxib with naproxen and other NSAIDs.^{26–28} However, in these studies, the racial composition of the study subjects was primarily white or undefined.

Several recent publications have highlighted disparities in treatment approaches and outcomes in Hispanic patients with cardiovascular disease, 29,30 asthma, and depression. Hispanics are less likely to receive or use medications for asthma,31 cardiovascular disease,32 human immunodeficiency virus infection/acquired immunodeficiency syndrome, 33 mental illness,³⁴ or pain,³⁵ as well as prescription medications in general. 36,37 These disparities in pharmaceutical treatment are substantial and often persist, even after adjustment for differences in income, age, insurance coverage, and coexisting medical conditions.³⁸ There is a paucity of data evaluating differences in response to medications between Hispanic and non-Latino populations. 38 Emerging research demonstrates that genetic variations affect Hispanic Americans and may require dosage adjustments to achieve an optimal therapeutic effect.^{39,40} The published literature highlights that Hispanics are cautious about American medicines, in part because of concerns about addiction, and often initiate downward dosage adjustments to avoid even minor side effects. 41 Given the increasing percentage of Hispanic Americans in the US population, studying the efficacy and safety of various medications in Hispanic populations will become increasingly important to health care practitioners and payers as they make treatment and formulary decisions for their populations.³⁹ To meet the data needs of payers and health care practitioners, more studies such as this one, which prospectively evaluated a specific medication in a Hispanic population, or the implementation of measures to increase the participation of Hispanic patients in broader clinical trials, will be required.³⁹

Safety and tolerability are important considerations when prescribing analgesic therapies. As noted above, they may be even more important in the Hispanic community. The composite measures of upper gastrointestinal tolerability presented in this paper, in addition to individually recorded adverse events, may provide a more clinically relevant assessment of treatment for the practicing physician. With only three upper gastrointestinal tolerability events (defined as moderate or severe nausea, abdominal pain, and/or dyspepsia) reported in the celecoxib treatment group, coupled with the low incidence of patient discontinuation due to treatment-related adverse events, the tolerability profile of celecoxib was demonstrated to be favorable.

Conclusion

This prospective, well controlled study of Hispanic patients provides insight into the efficacy and tolerability of celecoxib in the effective management of OA symptoms in a minority population. Celecoxib 200 mg once daily was noninferior to naproxen 500 mg twice daily for treating the signs and symptoms associated with OA of the knee in this patient group. In addition, both celecoxib and naproxen were shown to be safe and well tolerated. Research into the role of race or ethnicity in the response to treatment is still needed.

Disclosure

This study was sponsored by Pfizer Inc. The authors are fulltime employees of Pfizer Inc. Editorial support was provided by K Bradford of PAREXEL and was funded by Pfizer Inc.

References

- United States Census Bureau. US census bureau projections show a slower growing, older, more diverse nation a half century from now. US Department of Commerce. Available from: https://www.census. gov/newsroom/releases/archives/population/cb12-243.html. Accessed October 22, 2013.
- Lawrence RC, Felson DT, Helmick CG, et al. Estimates of the prevalence of arthritis and other rheumatic conditions in the United States. Part II. Arthritis Rheum. 2008;58(1):26–35.
- Centers for Disease Control and Prevention. Prevalence of doctordiagnosed arthritis-attributable effects among Hispanic adults, by Hispanic subgroup – United States, 2002, 2003, 2006, and 2009. MMWR Morb Mortal Wkly Rep. 2011;60(6):167–171.
- Centers for Disease Control and Prevention. Racial/ethnic differences in the prevalence and impact of doctor-diagnosed arthritis – United States, 2002. MMWR Morb Mortal Wkly Rep. 2005;54(5):119–123.
- Losina E, Walensky RP, Reichmann WM, et al. Impact of obesity and knee osteoarthritis on morbidity and mortality in older Americans. *Ann Intern Med*. 2011;154(4):217–226.
- Chacon JG, Gonzalez NE, Veliz A, et al. Effect of knee osteoarthritis on the perception of quality of life in Venezuelan patients. *Arthritis Rheum*. 2004;51(3):377–382.
- 2008 National Population Projections; 1970, 1980, 1990 and 2000 Decennial Censuses. US Census Bureau. Available from: http://www.census.gov/prod/2011pubs/12statab/pop.pdf. Accessed March 7, 2013.

- Dunlop DD, Song J, Manheim LM, Chang RW. Racial disparities in joint replacement use among older adults. Med Care. 2003;41(2):288–298.
- Herman CJ, Dente JM, Allen P, Hunt WC. Ethnic differences in the use of complementary and alternative therapies among adults with osteoarthritis. *Prev Chronic Dis*. 2006;3(3):A80.
- Olson JC, Foland J. Tracking racial and ethnic disparities of knee replacement rates in Connecticut. Conn Med. 2005;69(4):211–215.
- Skinner J, Weinstein JN, Sporer SM, Wennberg JE. Racial, ethnic, and geographic disparities in rates of knee arthroplasty among Medicare patients. N Engl J Med. 2003;349(14):1350–1359.
- Suarez-Almazor ME, Souchek J, Kelly PA, et al. Ethnic variation in knee replacement: patient preferences or uninformed disparity? *Arch Intern Med.* 2005;165(10):1117–1124.
- 13. Dominick KL, Bosworth HB, Jeffreys AS, Grambow SC, Oddone EZ, Horner RD. Racial/ethnic variations in non-steroidal anti-inflammatory drug (NSAID) use among patients with osteoarthritis. *Pharmacoepidemiol Drug Saf.* 2004;13(10):683–694.
- Celebrex [US prescribing information]. New York, NY, USA: Pfizer Inc.; 2013.
- Niculescu L, Li C, Huang J, Mallen S. Pooled analysis of GI tolerability of 21 randomized controlled trials of celecoxib and nonselective NSAIDs. Curr Med Res Opin. 2009;25(3):729–740.
- Altman R, Asch E, Bloch D, et al. Development of criteria for the classification and reporting of osteoarthritis. Classification of osteoarthritis of the knee. Diagnostic and Therapeutic Criteria Committee of the American Rheumatism Association. *Arthritis Rheum*. 1986;29(8): 1039–1049.
- Essex MN, O'Connell M, Bhadra BP. Response to nonsteroidal antiinflammatory drugs in African Americans with osteoarthritis of the knee. *J Int Med Res*. 2012;40(6):2251–2266.
- 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.
- Wendler D, Kington R, Madans J, et al. Are racial and ethnic minorities less willing to participate in health research? *PLoS Med.* 2006;3(2):e19.
- Shavers VL, Lynch CF, Burmeister LF. Factors that influence African-Americans' willingness to participate in medical research studies. *Cancer*. 2001;91(Suppl 1):233–236.
- Rochon PA, Mashari A, Cohen A, et al. The inclusion of minority groups in clinical trials: problems of under representation and under reporting of data. *Account Res.* 2004;11(3–4):215–223.
- Hussain-Gambles M, Atkin K, Leese B. Why ethnic minority groups are under-represented in clinical trials: a review of the literature. *Health Soc Care Community*. 2004;12(5):382–388.
- Bartlett C, Doyal L, Ebrahim S, et al. The causes and effects of sociodemographic exclusions from clinical trials. *Health Technol Assess*. 2005;9(38):iii-iv, ix-x, 1-152.
- Cronstein BN. Pharmacogenetics in the rheumatic diseases. Bull NYU Hosp Jt Dis. 2006;64(1–2):16–19.
- Asanuma Y, Xie HG, Stein CM. Pharmacogenetics and rheumatology: molecular mechanisms contributing to variability in drug response. *Arthritis Rheum*. 2005;52(5):1349–1359.
- Bensen WG, Fiechtner JJ, McMillen JI, et al. Treatment of osteoarthritis with celecoxib, a cyclooxygenase-2 inhibitor: a randomized controlled trial. Mayo Clin Proc. 1999;74(11):1095–1105.
- Kivitz AJ, Moskowitz RW, Woods E, et al. Comparative efficacy and safety of celecoxib and naproxen in the treatment of osteoarthritis of the hip. *J Int Med Res*. 2001;29(6):467–479.
- 28. McKenna F, Borenstein D, Wendt H, Wallemark C, Lefkowith JB, Geis GS. Celecoxib versus diclofenac in the management of osteoarthritis of the knee. *Scand J Rheumatol*. 2001;30(1):11–18.
- Cooper-DeHoff RM, Zhou Q, Gaxiola E, et al. Influence of Hispanic ethnicity on blood pressure control and cardiovascular outcomes in women with CAD and hypertension: findings from INVEST. *J Womens Health (Larchmt)*. 2007;16(5):632–640.

- Cooper-DeHoff RM, Aranda JM Jr, Gaxiola E, et al. Blood pressure control and cardiovascular outcomes in high-risk Hispanic patients findings from the International Verapamil SR/Trandolapril Study (INVEST).
 Am Heart J. 2006;151(5):1072–1079.
- Lieu TA, Lozano P, Finkelstein JA, et al. Racial/ethnic variation in asthma status and management practices among children in managed medicaid. *Pediatrics*. 2002;109(5):857–865.
- 32. Herholz H, Goff DC, Ramsey DJ, et al. Women and Mexican Americans receive fewer cardiovascular drugs following myocardial infarction than men and non-Hispanic whites: the Corpus Christi Heart Project, 1988–1990. *J Clin Epidemiol*. 1996;49(3):279–287.
- Moore RD, Stanton D, Gopalan R, Chaisson RE. Racial differences in the use of drug therapy for HIV disease in an urban community. N Engl J Med. 1994;330(11):763–768.
- Harris KM, Edlund MJ, Larson S. Racial and ethnic differences in the mental health problems and use of mental health care. *Med Care*. 2005;43(8):775–784.
- Pletcher MJ, Kertesz SG, Kohn MA, Gonzales R. Trends in opioid prescribing by race/ethnicity for patients seeking care in US emergency departments. *JAMA*. 2008;299(1):70–78.

- Hahn BA. Children's health: racial and ethnic differences in the use of prescription medications. *Pediatrics*. 1995;95(5):727–732.
- Xu KT, Rojas-Fernandez CH. Ancillary community pharmacy services provided to older people in a largely rural and ethnically diverse region: a survey of consumers in West Texas. *J Rural Health*. 2003;19(1): 79–86
- 38. Reyes C, Van de Putte L, Falcón AP, Levy RA. Genes, culture, and medicines: bridging gaps in treatment for Hispanic Americans. National Alliance for Hispanic Health and the National Pharmaceutical Council. Available from: http://www.hispanichealth.org/assets/resource_library/hispanic_report04.pdf. Accessed October 28, 2013.
- Burroughs VJ, Maxey RW, Levy RA. Racial and ethnic differences in response to medicines: towards individualized pharmaceutical treatment. J Natl Med Assoc. 2002;94(Suppl 10):1–26.
- Huang SM, Temple R. Is this the drug or dose for you? Impact and consideration of ethnic factors in global drug development, regulatory review, and clinical practice. Clin Pharmacol Ther. 2008;84(3):287–294.
- Grissinger M. Cultural diversity and medication safety. P&T Community. Available from: http://www.ptcommunity.com/ptJournal/ fulltext/32/9/PTJ3209471.pdf. Accessed April 22, 2014.

International Journal of General Medicine

Publish your work in this journal

The International Journal of General Medicine is an international, peer-reviewed open-access journal that focuses on general and internal medicine, pathogenesis, epidemiology, diagnosis, monitoring and treatment protocols. The journal is characterized by the rapid reporting of reviews, original research and clinical studies across all disease areas.

A key focus is the elucidation of disease processes and management protocols resulting in improved outcomes for the patient. The manuscript management system is completely online and includes a very quick and fair peer-review system. Visit http://www.dovepress.com/testimonials.php to read real quotes from published authors.

 $\textbf{Submit your manuscript here:} \ \text{http://www.dovepress.com/international-journal-of-general-medicine-general-medicine-general-medi$

