

What Your Patients Need to Know About NSAIDs—A Case Study

When Prescribing NSAIDs...



USE THE
LOWEST
EFFECTIVE
DOSE



FOR THE
SHORTEST
PERIOD OF
TIME

Joan M.

Joan M. is a 62-year-old woman with plantar fasciitis who was prescribed a nonsteroidal anti-inflammatory drug (NSAID) and advised to use orthotic shoe inserts. At a follow-up visit, the patient reports that she has been experiencing fairly regular headaches, most likely associated with seasonal allergies, and is taking an over-the-counter (OTC) NSAID almost daily concurrent with her prescription NSAID.

Not an actual patient

Clinical Issues

- Concurrent use of >1 NSAID
- Inadvertent polypharmacy
- Use of higher dose of NSAID than prescribed

Case Commentary

Concomitant and typically inadvertent use of more than 1 NSAID at a time is a common practice among patients in the United States. In a survey of a random sample of the US public, 38% of respondents admitted to concurrent use of OTC and prescription NSAIDs.¹ This polypharmacy may occur because patients do not know what the term NSAID means or which products contain NSAIDs. They are also often unaware of the potential adverse events (AEs) associated with these agents, which include potentially severe gastrointestinal (GI), cardiovascular (CV), and renal toxicity.^{1,2}



Alliance for Rational
Use of NSAIDs

A Public Health Coalition

What Joan M. Should Know

Use of more than 1 NSAID concurrently is concerning because it increases the risk for NSAID-associated toxicities and may result in a preventable medication overdose. Additionally, the concomitant use of multiple NSAID products is outside of accepted standard of care, which explicitly directs NSAID use at the lowest effective dose for the shortest time required to achieve the desired therapeutic effect.³⁻⁷

- Compared with use of 1 NSAID, concurrent use of:
 - **>1 NSAID** approximately doubled the risk for acute renal failure (ARF)⁸
 - **2 NSAIDs** doubled odds of a GI bleed, and **≥3 NSAIDs** increased odds 12-fold⁹

In addition, taking more than 1 NSAID at a time increases the NSAID dose, which increases the risk for NSAID-associated GI, CV, and renal toxicity.

- Compared with low-dose NSAID use, high-dose NSAID use:
 - Increased the risk of upper GI complications by **70% to 230%**,^{10*†} and the risk of upper GI bleeding by approximately **3- to 6-fold**^{9*†}
 - Increased the risk of ARF by **50% to 240%**^{8,11*}
 - Increased the risk of myocardial infarction (MI) by **28%**¹²
- Risk of MI was increased 20% with NSAID use longer than 1 month, and 35% with use longer than 3 years¹²

Practical Considerations

The risks of NSAID use can be minimized if they are taken according to evidence-based guidelines—1 NSAID at a time at the lowest effective dose for the shortest period of time medically required to achieve the desired therapeutic effect.³⁻⁷

However, the availability and use of OTC NSAID products complicates patient compliance with recommendations and impacts their appropriate use. Therefore, patient education is critical when prescribing NSAIDs—patients should be counseled to use only 1 type of NSAID at a time and to take NSAIDs exactly as indicated or prescribed to help to avoid NSAID-associated AEs. It is also important to educate patients about which medications, both prescription and OTC, are in the NSAID class, and for health care providers to ask about OTC NSAID use as a standard part of every medication review to minimize risk and maximize therapeutic benefit.

The Alliance for Rational Use of NSAIDs—a public health coalition—aims to bridge the gap between guidance and clinical practice, educating health care professionals and the public at-large to ensure safe and appropriate use of NSAIDs.

Visit www.NSAIDAlliance.com to take an interactive survey about NSAID risks and learn more about the Alliance for Rational Use of NSAIDs.

*Relative risk or odds ratio varied depending on the NSAID used.

†NSAIDs used in these studies were not accompanied by gastroprotection.

References: 1. Wilcox CM, Cryer B, Triadafilopoulos G. Patterns of use and public perception of over-the-counter pain relievers: focus on nonsteroidal antiinflammatory drugs. *J Rheumatol*. 2005;32:2218-2224. 2. Lanias A, Garcia-Tell G, Armada B, Oteo-Alvaro A. Prescription patterns and appropriateness of NSAID therapy according to gastrointestinal risk and cardiovascular history in patients with diagnoses of osteoarthritis. *BMC Medicine*. 2011;9:38. 3. US Food and Drug Administration Web site. Public Health Advisory: FDA announces important changes and additional warnings for COX-2 selective and non-selective non-steroidal anti-inflammatory drugs (NSAIDs). <http://www.fda.gov/Drugs/DrugSafety/PostmarketDrugSafetyInformationforPatientsandProviders/ucm150314.htm>. Accessed September 18, 2012. 4. European Medicines Agency. *Public CHMP Assessment Report for Medicinal Products Containing Non-selective Non Steroidal Anti-inflammatory Drugs (NSAIDs)*. London, England: European Medicine Agency; November 7, 2006. 5. Anderson JL, Adams CD, Antman EM, et al. ACC/AHA 2007 Guidelines for the Management of Patients With Unstable Angina/Non-ST-Elevation Myocardial Infarction: a report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines (Writing Committee to Revise the 2002 Guidelines for the Management of Patients With Unstable Angina/Non-ST-Elevation Myocardial Infarction) developed in collaboration with the American College of Emergency Physicians, the Society for Cardiovascular Angiography and Interventions, and the Society of Thoracic Surgeons endorsed by the American Association of Cardiovascular and Pulmonary Rehabilitation and the Society for Academic Emergency Medicine. *J Am Coll Cardiol*. 2007;50:e1-e157. 6. Rostom A, Moayyedi P, Hunt R, for the Canadian Association of Gastroenterology Consensus Group. Canadian consensus guidelines on long-term nonsteroidal anti-inflammatory drug therapy and the need for gastroprotection: benefits versus risks. *Aliment Pharmacol Ther*. 2009;29:481-496. 7. Zhang W, Moskowitz RW, Nuki G, et al. OARSI evidence-based, expert consensus guidelines. *Osteoarthritis Cartilage*. 2008;16:137-162. 8. Schneider V, Lévesque LE, Zhang B, Hutchinson T, Brophy JM. Association of selective and conventional nonsteroidal anti-inflammatory drugs with acute renal failure: a population-based, nested case-control analysis. *Am J Epidemiol*. 2006;164:881-889. 9. Lewis SC, Langman MJS, Laporte J-R, Matthews JNS, Rawlins MD, Wilhelm B-E. Dose-response relationships between individual nonaspirin nonsteroidal anti-inflammatory drugs (NANSAs) and serious upper gastrointestinal bleeding: a meta-analysis based on individual patient data. *Br J Clin Pharmacol*. 2002;54:320-326. 10. Riera-Guardia N, Castellsague J, Calingaert B, et al. The SOS Project: nonsteroidal anti-inflammatory drugs and upper gastrointestinal complications. Meta-analysis of epidemiological studies. Presented at: 26th International Conference on Pharmacoepidemiology & Therapeutic Risk Management; August 2010; Brighton, England. 11. Griffin MR, Yared A, Ray WA. Nonsteroidal anti-inflammatory drugs and acute renal failure in elderly persons. *Am J Epidemiol*. 2000;151:488-496. 12. Garcia Rodriguez LA, Tacconelli S, Patrignani P. Role of dose potency in the prediction of risk of myocardial infarction associated with nonsteroidal anti-inflammatory drugs in the general population. *J Am Coll Cardiol*. 2008;52:1626-1636.

