

This second article in a three-part series on osteoarthritis discusses pharmacological and non-pharmacological management. The third part will be published online

OSTEOARTHRITIS PART 2 OF 3 | PART 3 WILL BE PUBLISHED ON NURSINGTIMES.NET ON 28 FEBRUARY

Osteoarthritis 2: pain management and treatment strategies

In this article...

- Introduction to the management of osteoarthritis (OA)
- Pharmacological and non-pharmacological management of OA

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Abstract Swift A (2012) Osteoarthritis 2: pain management and treatment strategies. *Nursing Times*; 108: 8, 25-27. Osteoarthritis (OA) is a painful, progressive joint disorder. This article discusses pharmacological management of OA, such as non-steroidal anti-inflammatory drugs and opioids, and non-pharmacological management, including weight reduction, acupuncture and joint replacement surgery. The third part, to be published online, will cover the physical, psychological and social impact of OA.

Osteoarthritis (OA) is a painful, progressive joint disorder that can cause stiffness, fatigue, depression and anxiety. Many people with the condition have difficulty in sleeping and carrying out everyday activities, and it can lead to loss of work and social isolation, reducing quality of life.

Most people who visit their GP with OA symptoms will have experienced pain and associated disability for three months or more; one in eight will be using a walking aid and 75% will have tried analgesics (Birrell et al, 2000). This shows many people with symptomatic OA try to self-manage the condition before seeking help.

People with the condition require an individualised treatment package.

Guidelines on the management of OA

use a combination of systematic reviews and Delphi techniques to form a consensus about how the condition should be treated (Zhang et al, 2010; NICE, 2008). In a systematic review, a group of experts appraise evidence and retain the high-quality papers to form the guidance. The Delphi technique involves a facilitator consulting a group of experts to reduce a lengthy list of potential treatments to a mutually agreeable smaller list of the “best” available treatments.

Pharmacological management

Medications used to manage OA pain include paracetamol, non-steroidal anti-inflammatory drugs (NSAIDs) and opioids.

Paracetamol

Paracetamol is usually the first-line pharmacological treatment for managing OA pain. Although it can be as effective as NSAIDs in some people, evidence suggests paracetamol has only a modest effect in reducing pain, and no effect on stiffness or function (Towheed et al, 2006). NICE recommends trying paracetamol first because, although NSAIDs provide better pain relief, paracetamol has fewer side-effects (NICE, 2008). Paracetamol is a relatively safe drug when taken at doses of less than 4,000mg per day, but it can cause gastric irritation if taken in high doses over a long period (Rahme et al, 2002).

According to NICE (2008), paracetamol and codeine combinations have no

5 key points

1 Osteoarthritis (OA) is a painful, progressive joint disorder

2 Paracetamol is usually the first line drug treatment for managing OA pain

3 Non-steroidal anti-inflammatory drugs are associated with gastric problems, dyspepsia and thrombus formation

4 Obesity is a risk factor for OA

5 More than 114,000 replacements or revisions for hip or knee joints were carried out in the UK last year



Heberden's nodes are a sign of OA

demonstrable benefit over paracetamol alone. Comparisons of paracetamol-dextropropoxyphene with NSAIDs tend to favour NSAIDs (NICE, 2008).

Non-steroidal anti-inflammatory drugs

NICE (2008) recommends using NSAIDs to manage OA pain, but they carry risks. Around 10-20% of people who use oral NSAIDs experience dyspepsia, and risk gastric complications (Wolfe et al, 1999).

The development of Cox-2 selective anti-inflammatory drugs reduced the incidence of gastric side-effects, but highlighted another problem of NSAIDs – the risk of thrombus formation, particularly with long-term use, in older people and in those with other prothrombotic risk factors. This increases the risk of myocardial infarction and stroke (Cannon et al, 2006). NSAIDs are

also associated with a slight increase in the risk of renal failure (Adam, 2011).

The risk of adverse events associated with NSAID use decreases significantly when topical gels are used instead of oral preparations. This has led to topical NSAIDs being suggested as a first-line treatment for managing OA (Zhang et al, 2010).

Opioids

Opioids can be extremely effective in the management of pain. However, their benefits can be outweighed by side-effects such as nausea, vomiting, dizziness and constipation, which affect a lot of people who use these drugs (Zhang et al, 2010). A recent review that explored the use of extended-release opioids in OA found they would help to improve sleep (Turk and Cohen, 2010).

Long-term use of opioids is associated with tolerance, where sensitivity to the analgesic effects of opioids decreases over time, and hyperalgesia, where sensitivity to painful stimuli increases. Tolerance leads to a need to increase the dose of the drug, which could increase adverse effects without relieving pain. This can make clinicians reluctant to use opioids to control long-term, non-malignant pain. However, some studies suggest concerns about this are overstated, and that very few people have this problem (Schneider and Kirsh, 2010).

Patients and practitioners can be fearful about the risks of addiction with opioid use. Portenoy (1990) developed criteria to identify addiction in people with chronic pain, taking into consideration the “normal” behaviours of a patient in pain that could be misconstrued as addiction (Box 1); addiction is diagnosed if drug-taking has persisted for at least a month and at least three of nine of the criteria are present. The criteria should be used with caution; a study by Hojsted et al (2010) found that, according to Portenoy’s criteria, almost one in five of a group of people with chronic pain were addicted to their opioid medication.

Tramadol

Tramadol is a synthetic analgesic that, in addition to having a morphine-like mode of action, is also a serotonin and noradrenalin reuptake inhibitor (Mongin, 2007).

This offers a theoretical advantage over other opioid preparations because it can increase the concentration of serotonin and noradrenalin in the central nervous system.

Its side-effects include nausea, vomiting, drowsiness, constipation and headache. In trials exploring the use of tramadol in OA, these side-effects were mild or moderate, most often occurring during the initial titration phase rather than the

maintenance phase (Langley et al, 2010). However, these reactions may reduce concordance and might also lead to secondary harms such as an increase in falls.

Tricyclic antidepressants

Tricyclic antidepressants (TCAs) work by blocking the uptake of serotonin and noradrenaline from synapses in the brain and spinal cord. These neurotransmitters are involved in inhibiting pain signals, and blocking their removal from synapses improves the body’s inhibition of pain signals. Theoretically, TCAs should be helpful in OA pain, but only one study has explored this. The study was inconclusive in terms of pain relief, but support is emerging for the benefits of TCAs in OA pain (NICE, 2008).

Corticosteroids or hyaluronic acid injections

Steroid or hyaluronic acid injections into the knee joints can bring partial pain relief for up to three months, and are often sought by people with OA (NICE 2008).

However, there is a lack of placebo-controlled trials, and some trials suggest efficacy is limited (Avouac et al, 2007). There is also a risk of joint infection, and steroids can cause blood glucose instability in people with diabetes (Habib and Safia, 2009).

Capsaicin

Capsaicin, which is a topical cream, causes localised burning pain but, as it takes a while for the body to synthesise the capsaicin, the period of burning is followed by pain relief. Using the cream twice daily often means the burning sensation does not return after the first application. A half-strength version is available for those who



Swollen knee in a man with osteoarthritis

find it hard to tolerate the full strength preparation (Schnitzer et al, 1995).

Nutraceuticals and herbal products

Glucosamine is one of the more popular nutraceutical products in the UK, but there is little evidence that it is effective and it is not recommended by NICE (2008).

There is some evidence that rose hip powder, devil’s claw and egg-shell membrane have some efficacy in pain relief, but research is in the early stages (Zhang et al, 2010; Ruff et al, 2009). There is also evidence some people benefit from eliminating certain foods from their diet, and from following eating plans such as a Mediterranean diet (Rayman and Pattison, 2008).

Non-pharmacological management

Non-pharmacological management strategies include: exercise; transcutaneous electrical nerve stimulation (TENS); acupuncture; heat or cold; weight reduction; joint supports; walking aids; surgery.

Exercise

People with OA need to maintain fitness, stamina and suppleness to facilitate confidence, independence and quality of life. Exercise can also help to alleviate pain, and fitness has a positive influence on post-operative recovery for those who have had joint replacements (Pisters et al, 2010).

Evidence suggests aerobic exercise leads to a better outcome than strengthening exercises, which are more effective than water-based exercises (Zhang et al, 2010). Physiotherapists can give advice about exercise, although most people just need advice to keep fit and active.

Improvements in physical function and pain can lead to improvements in depression (Lim et al, 2005).

TENS and acupuncture

Acupuncture and TENS have been shown to be helpful in relieving pain (Itoh et al, 2008).

BOX 1. PORTENOY’S CRITERIA FOR ADDICTION IN PEOPLE WITH CHRONIC PAIN

- Intense desire for the drug
- Overwhelming concern for its availability
- Compulsive drug use
- Unsanctioned escalation of dose
- Continued use despite significant side-effects
- Use of drugs to treat non-pain symptoms
- Unapproved use of drugs when no pain is present
- Efforts to manipulate the medical system or staff to obtain drugs
- Acquisition of drugs from elsewhere

Evidence suggests, while acupuncture can be expensive, it can help to alleviate pain and stiffness, and improve function (Zhang et al, 2010). TENS – the application of an electrical stimulus across sensory nerve pathways – stimulates the production of endorphins. This has been shown to have a positive effect on pain, and is a technique people with can do themselves (NICE, 2008).

Heat or cold

The application of superficial heat or cold to joints can provide short-term pain relief (Denegar et al, 2010).

Heat can be applied in numerous ways, including immersion in warm water, heat packs heated at home in the microwave, heat pads and wax. Cold is usually applied with an ice massage or by applying cold packs to the affected area.

Although there is little evidence to support the use of heat, anecdotal evidence supports it. Using a safe system is important; hot water bottles have been known to damage the skin as the temperature can be too high for direct application to skin. This is particularly important for older people.

Weight reduction

Obesity is a major risk factor in the development of OA, and is associated with a greater need for joint replacement surgery (Turley et al, 2006).

Weight reduction is thought to be a beneficial pain management strategy, but there is no strong evidence to support this. Advice on weight is based on expert opinion rather than randomised controlled trials or systematic reviews (Zhang et al, 2010).

Weight reduction is one of three core interventions in NICE (2008) guidance on OA, along with information and exercise. This could be due to the overall health benefits of a normal BMI, and because obesity is a contraindication for surgery. Obesity is associated with higher anaesthetic risk, greater risk of mechanical failure of the joint replacement and a greater risk of infection after surgery (Samson et al, 2010).

Braces, sleeves and walking aids

People with OA often use walking aids but many people do not obtain them by prescription (van der Esch et al, 2003). This creates a potential problem of incorrect technique and sizing, which can increase risk of falls and contribute to misalignment leading to the development of musculoskeletal pain in other places.

Use of walking aids or knee braces can help to reduce pain (NICE, 2008). Knee braces are more effective than neoprene sleeves (Brouwer et al, 2005).

Joint replacement

Replacement of the hip or knee joint can be an effective management strategy. In 2010, more than 114,000 hip and knee replacements or revisions were carried out in the UK (www.njrcentre.org.uk).

Surgery can be helpful in terms of pain reduction and restoration of function, but improvement is not guaranteed. Most people benefit from it, but around a quarter report ongoing moderate to severe pain five to eight years after surgery. Around one fifth of patients say they have not recovered sufficient function after surgery (Jones et al, 2000). Reasons include issues to do with the joints and prosthesis, and issues relating to the preoperative pain mechanism (Wylde et al, 2011; 2007).

Summary

The management of OA involves both medical and surgical options. Nurses can help patients decide which therapies will help relieve their pain, and improve function and quality of life. **NT**

The full version of this article can be found on nursingtimes.net

References

Adam WR (2011) Non-steroidal anti-inflammatory drugs and the risks of acute renal failure: number needed to harm. *Nephrology*; 16: 2, 154-155.

Avouac J et al (2007) Efficacy and safety of opioids for osteoarthritis: a meta-analysis of randomized controlled trials. *Osteoarthritis and Cartilage*; 15: 8, 957-965.

Birrell F et al (2000) Health impact of pain in the hip region with and without radiographic evidence of osteoarthritis: a study of new attenders to primary care. The PCR Hip Study Group. *Annals of the Rheumatic Diseases*; 59: 11, 857-863.

Brouwer RW et al (2005) Braces and orthoses for treating osteoarthritis of the knee. *Cochrane Database of Systematic Reviews*; 1: CD004020.

Cannon CP et al (2006) Cardiovascular outcomes with etoricoxib and diclofenac in patients with osteoarthritis and rheumatoid arthritis in the Multinational Etoricoxib and Diclofenac Arthritis Long-term (MEDAL) programme: a randomised comparison. *The Lancet*; 368: 9549, 1771-1781.

Denegar CR et al (2010) Preferences for heat, cold, or contrast in patients with knee osteoarthritis affect treatment response. *Clinical Interventions in Aging*; 5: 199-206.

Habib G, Safia A (2009) The effect of intra-articular injection of betamethasone acetate/betamethasone sodium phosphate on blood glucose levels in controlled diabetic patients with symptomatic osteoarthritis of the knee. *Clinical Rheumatology*; 28: 1, 85-87.

Hojsted J et al (2010) Classification and identification of opioid addiction in chronic pain patients. *European Journal of Pain*; 14: 10, 1014-1020.

Itoh K et al (2008) A pilot study on using acupuncture and transcutaneous electrical nerve stimulation (TENS) to treat knee osteoarthritis (OA). *Chinese Medicine*; 3: 2.

Jones CA et al (2000) Health related quality of life outcomes after total hip and knee arthroplasties in a community based population. *The Journal of Rheumatology*; 27: 7, 1745-1752.

Langley PC et al (2010) Adverse event profile of tramadol in recent clinical studies of chronic osteoarthritis pain. *Current Medical Research & Opinion*; 26: 1, 239-251.

Lim HJ et al (2005) Effects of home-based daily exercise therapy on joint mobility, daily activity, pain, and depression in patients with ankylosing spondylitis. *Rheumatology International*; 25: 3, 225-229.

Mongin G (2007) Tramadol extended-release formulations in the management of pain due to osteoarthritis. *Expert Review of Neurotherapeutics*; 7: 12, 1775-1784.

National Institute for Health and Clinical Excellence (2008) *Osteoarthritis: National Clinical Guideline for Care and Management in Adults*. London: NICE. www.nice.org.uk/cg59

Pisters MF et al (2010) Long-term effectiveness of exercise therapy in patients with osteoarthritis of the hip or knee: a randomized controlled trial comparing two different physical therapy interventions. *Osteoarthritis and Cartilage*; 18: 8, 1019-1026.

Portenoy RK (1990) Chronic opioid therapy in nonmalignant pain. *Journal of Pain and Symptom Management*; 5: S46-S62.

Rahme E et al (2002) Determinants and sequelae associated with utilization of acetaminophen versus traditional nonsteroidal antiinflammatory drugs in an elderly population. *Arthritis & Rheumatism*; 46: 11, 3046-3054.

Rayman MP, Pattison DJ (2008) Dietary manipulation in musculoskeletal conditions. *Best Practice & Research Clinical Rheumatology*; 22: 3, 535-561.

Ruff KJ et al (2009) Eggshell membrane in the treatment of pain and stiffness from osteoarthritis of the knee: a randomized, multicenter, double-blind, placebo-controlled clinical study. *Clinical Rheumatology*; 28: 8, 907-914.

Samson AJ et al (2010) Total knee replacement in the morbidly obese: a literature review. *ANZ Journal of Surgery*; 80: 9, 595-599.

Schneider JP, Kirsh KL (2010) Defining clinical issues around tolerance, hyperalgesia, and addiction: a quantitative and qualitative outcome study of long-term opioid dosing in a chronic pain practice. *Journal of Opioid Management*; 6: 6, 385-395.

Schnitzer TJ et al (1995) High strength capsaicin cream for osteoarthritis pain: rapid onset of action and improved efficacy with twice daily dosing. *Journal of Clinical Rheumatology*; 1: 5, 268-273.

Towheed TE et al (2006) Acetaminophen for osteoarthritis. *Cochrane Database of Systematic Reviews*; 1: CD004257

Turk DC, Cohen MJ (2010) Sleep as a marker in the effective management of chronic osteoarthritis pain with opioid analgesics. *Seminars in Arthritis and Rheumatism*; 39: 6, 477-490.

Turley M et al (2006) Non-fatal disease burden associated with excess body mass index and waist circumference in New Zealand adults. *Australian and New Zealand Journal of Public Health*; 30: 3, 231-237.

van der Esch M et al (2003) Factors contributing to possession and use of walking aids among persons with rheumatoid arthritis and osteoarthritis. *Arthritis & Rheumatism*; 49: 6, 838-842.

Wolfe MM et al (1999) Gastrointestinal toxicity of nonsteroidal antiinflammatory drugs. *New England Journal of Medicine*; 340: 24, 1888-1899.

Wylde V et al (2011) Persistent pain after joint replacement: Prevalence, sensory qualities, and postoperative determinants. *Pain*; 152: 3, 566-572, available from: PM:21239114.

Wylde V et al (2007) Total knee replacement: Is it really an effective procedure for all? *Knee*. available from: PM:17596949.

Zhang W et al (2010) OARSI recommendations for the management of hip and knee osteoarthritis: part III: changes in evidence following systematic cumulative update of research published through January 2009. *Osteoarthritis and Cartilage*; 18: 4, 476-499.