In 2014, an estimated 14,000 fatalities were due to opioid overdoses in cases in which the drug was prescribed for therapeutic purposes. The overall incidence was more than 28,000 deaths for all opioid-related deaths (prescription and illicit).

By comparison, the total opioid overdose death rate is higher than the annual US death rates from AIDS/HIV (6955) or oral cancer (>8000). Opioid prescribing by dentists is estimated to be 11% of the overall annual number of opioid prescriptions in the United States, indicating that approximately 1500 deaths annually may be attributable to unused opioids originally prescribed by dentists for therapeutic purposes. This number may be much higher as prescribed opioids are often reported as the “gateway” for heroin addiction. These projections suggest that dentists’ prescriptions for opioid analgesics may ultimately be associated with opioid overdose deaths. The overall burden is likely higher for management of acute dental pain as emergency department physicians also overwhelmingly prescribe opioid analgesics for nontraumatic dental conditions.

Below, we review the evidence that supports this conclusion and consider the role of patients’ expectations for receiving an opioid for acute dental pain that conflicts with the ethical imperative to “do no harm” for analgesic prescribing as part of dental care. We also provide alternative evidence-based therapeutic strategies for treating acute pain in dental practice. The data presented indicate opioids should be prescribed only as a last choice for acute dental pain. A failure for the dental profession to change from routine opioid prescribing for acute dental pain to more rational alternatives in the face of overwhelming evidence may continue to significantly contribute to the public health crisis in the opioid overdose epidemic.

**Analgesic Drug Prescribing**

The inadequate management of acute and chronic pain in the United States resulted in the implementation of Veterans Health Administration’s 1999 initiative “Pain as the 5th Vital Sign” for evaluating patients in medical environments and the much more recent 2016 National Pain Strategy to improve pain management and research in the United States. Recognition of the pain-relieving effects of antidepressant and anticonvulsant drugs for patients with chronic pain did result in the development of more effective drugs for neuropathic pain. The much-heralded advantages of...
Opioid analgesics are effective for relieving pain by acting at opiate receptors in the central nervous system and can be safely administered for acute and chronic pain under supervision of a health care practitioner. Repeated administration of an opioid drug eventually produces cellular changes that result in the need for increasing doses of the drug to produce the same effect (tolerance). Continued administration also results in molecular changes that require continuous receptor occupancy to avoid the unpleasantness of withdrawal (physical dependence) and an intense craving for the euphoria associated with opioid drugs (psychological dependence). These effects can be mitigated by administration of drugs in a therapeutic context. When withdrawal and dependence occur as part of drug abuse, the intense craving for opioid drugs promotes attempts to acquire drugs illicitly, which is referred to as addiction. Opioid drug abuse is partially heritable and can be precipitated from as little as a single dose to the progressive continuum of tolerance, dependence, and eventual addiction and overdose.

Variability in the composition and potency of illicit opioids increases the probability of inadvertent overdose,\textsuperscript{9} which, if not witnessed, can progress to respiratory depression, unconsciousness, and death. Rehabilitation from opioid dependence has had limited success while substitution of illicit opioids with methadone-therapy maintenance minimizes cravings and drug-seeking behavior but does not directly address the underlying physical and psychological dependence. Distribution of naloxone kits to emergency-medical technicians and families of drug-dependent individuals has improved resuscitation success but also provides a rationale for continued abuse on the assumption that respiratory depression can be readily reversed.\textsuperscript{10} Any opioid prescription for acute pain can progress to lifelong drug dependence or acute mortality due to drug overdose, if precautions are not taken.

In theory, opioid prescribing for acute pain contributes to greater availability of opioid drugs available for misuse, abuse, or diversion, thereby indirectly adding to the serious morbidity and mortality associated with prescription opioids. An estimated 12% of patients reporting back pain, dental pain, or headache in emergency departments are “doctor shopping” for an opioid prescription.\textsuperscript{11} Similarly, a survey of patients seeking care at an emergency dental clinic indicated that 37% of patients reported nonmedical use of prescription pain medications in the previous 30 days.\textsuperscript{12} Among factors contributing to the initiation of substance use, nonmedical use of prescription pain medications was the second most-prevalent illicit first-time drugs for persons (<12 years old) in the United States.\textsuperscript{13} The connection with dentistry is concerning. The prevalence of dental practices and pharmacies in counties within Indiana was identified as the leading predictor of enrollment in drug-abuse treatment programs in the state.\textsuperscript{14} This observation suggests that the availability of prescription opioids in a community (eg, unused medications in a household cabinet) is associated with higher rates of opioid abuse and that management of acute orofacial pain may eventually produce cellular changes that result in the need for increasing doses of the drug to produce the same effect (tolerance). Continued administration also results in molecular changes that require continuous receptor occupancy to avoid the unpleasantness of withdrawal (physical dependence) and an intense craving for the euphoria associated with opioid drugs (psychological dependence). These effects can be mitigated by administration of drugs in a therapeutic context. When withdrawal and dependence occur as part of OTC medications is often required. Analgesics for acute dental pain may be contributing to dependence rates.

Acute Dental Pain Management
Pain following a typical dental procedure such as a simple extraction is usually minor and transient and can be effectively managed with over-the-counter (OTC) drugs. When postprocedural pain follows a surgical extraction, implant placement, or endodontic procedure or when preexisting infection or inflammation has sensitized nociceptive processes, an analgesic with greater efficacy than OTC medications is often required. Analgesics for acute dental pain are usually orally administered, should produce minimal adverse events in ambulatory patients, and have minimal effects on cogni-
tion and alertness so that patients can return to the activities of daily living. These medications should be prescribed for a short period. Control of pain following a procedure that produces tissue injury and subsequent inflammation is best accomplished by targeting the etiology of the pain, ideally by attenuating the acute inflammatory process. Symptom management by interfering with the perception of pain with opioid drugs that act in the central nervous system does not target the cause of acute dental pain and often impairs alertness, cognition, and psychomotor function. In addition, ambulatory patients are more prone to adverse effects associated with motion, such as dizziness, nausea, and vomiting.\textsuperscript{15}

Before NSAIDs were introduced in the mid 1970s to the US market, orally administered analgesics were a compromise between the ceiling of analgesic efficacy associated with aspirin and acetaminophen and the undesirable effects of opioids when given to ambulatory patients. A standard dose of a peripherally acting agent (aspirin or acetaminophen) was combined with an orally effective opioid (codeine or oxycodone) at a dose selected to provide additive analgesia, often combined with adjunctive agents thought to potentiate analgesia (caffeine) or to counter the adverse effects of the opioid (promethazine). A critical appraisal of analgesic combinations by the Food and Drug Administration resulted in elimination of several adjuncts and reformulation of some combinations to meet regulatory requirements for safety and additive analgesia.\textsuperscript{16}

The treatment of acute pain was subsequently revolutionized by the introduction of NSAIDs and the development of the oral-surgery model\textsuperscript{17} for evaluating analgesic efficacy and side-effect liability for dental outpatients, rather than relying on extrapolation from cancer or general-surgery studies.\textsuperscript{18} Drugs and doses now used for acute postprocedural pain have been characterized through thousands of well-controlled clinical trials, providing a large and robust evidentiary basis for therapeutic recommendations.

### NSAID Analgesics

NSAIDs are widely used for acute dental pain as they are generally more efficacious than aspirin, acetaminophen, or codeine (Figure 1) due to the inflammatory cause of most dental pain and NSAIDs’ prominent anti-inflammatory effects. NSAID therapy is also preferable for ambulatory patients who generally experience a high incidence of side effects when given an opioid. NSAIDs modestly suppress the development of edema after surgical procedures,\textsuperscript{19} providing additional benefit without the potential liabilities of administering steroids. When used as directed, OTC dosing regimens for ibuprofen, ketoprofen, or naproxen sodium are safe and effective across a wide variety of dental-pain conditions.\textsuperscript{20} These conditions and the over 40 years of clinical experience with ibuprofen make NSAIDs the drug class of choice for dental pain for patients who do not have any contraindications to its use (Table 1).

Ibuprofen at a dose of 400 mg is superior to 650 mg of aspirin, 600 mg to 1000 mg of acetaminophen, combinations of aspirin and acetaminophen plus 60 mg of codeine, and 30 mg of dihydrocodeine when evaluated in the oral-surgery model.\textsuperscript{21,22} Administration of doses greater than 400 mg is not likely to result in greater peak analgesia, but the increased blood levels may modestly prolong the duration of effect.\textsuperscript{23} Other NSAIDs such as meclofenamate, naproxen sodium, and diclofenac result in analgesia that is comparable to ibuprofen, depending on the doses of each agent used. The use of NSAIDs also results in clinically significant analgesia for periodontal surgery,\textsuperscript{24} orthodontic tooth movement,\textsuperscript{25} and pain following endodontic procedures.\textsuperscript{26}

Ketorolac tromethamine is an NSAID approved for parenteral and nasal administration for short-term management of moderate- to severe pain, and several analgesic models have shown it is comparable in efficacy to 100 mg of meperidine and 10 mg to 12 mg of morphine.\textsuperscript{27,28} Parenteral administration of ketorolac produces less drowsiness, nausea, and vomiting than morphine;\textsuperscript{29} however, this route of administration limits its use for ambulatory patients to the initial dose prior to discharge. The ability of injectable ketorolac to overcome the slower onset of orally administered drugs, combined with analgesic efficacy comparable to that of parenteral opioids but with reduced side effects suggests that its use for a procedure likely to result in moderate-to-severe pain obviates the need to prescribe an opioid combination. The intravenous administration of ibuprofen or acetaminophen perioperatively may also minimize the need for the routine use of postoperative opioid pain relievers.

When an NSAID is combined with therapeutic strategies for preventing acute pain and minimizing its intensity, as described elsewhere,\textsuperscript{31} the rationale for prescribing an opioid combination

### TABLE 1

<table>
<thead>
<tr>
<th>Preexisting Medical Conditions</th>
<th>Possible Adverse Effects of NSAID</th>
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<tbody>
<tr>
<td>Gastric ulcer</td>
<td>Internal bleeding</td>
</tr>
<tr>
<td>Asthma</td>
<td>Asthmatic attack</td>
</tr>
<tr>
<td>Diabetes</td>
<td>High doses may cause hyperglycemia or hypoglycemia</td>
</tr>
<tr>
<td>Gout</td>
<td>Alternation in plasma urate levels</td>
</tr>
<tr>
<td>Influenza, varicella</td>
<td>Reye syndrome in children</td>
</tr>
</tbody>
</table>
| Hypocoagulation               | • Excessive bleeding following surgery  
|                               | • Increased risk of cardiovascular events (MI or stroke) |

Abbreviations: MI = myocardial infarction, NSAID = nonsteroidal anti-inflammatory drug.
to be taken as needed for acute pain is even less warranted. Table 2 summarizes a stepwise, multimodal approach to acute pain management that targets the inflammatory etiology of pain with NSAID administration to minimize the effects of inflammatory mediators released due to tissue injury and gene expression. Preventive administration of an NSAID and long-acting local anesthetic also blocks physiologic events that augment the effects of the inflammatory process. Adjunctive administration of acetaminophen provides additive analgesia without the deleterious effects of abuse-prone opioids such as oxycodone and hydrocodone. Prescribing tramadol is less likely to contribute to abuse, especially if a limited 3-day supply is prescribed without any refills. Administration of an opioid combination containing oxycodone or hydrocodone does not target inflammation, does not implement preventive strategies, and facilitates opioid misuse, abuse, and diversion that can lead directly to opioid overdoses.

**Opioid Combination Analgesics**

Many opioid analgesics have poor oral potency due to low oral bioavailability; opioids such as morphine, meperidine, and oxymorphone have such low oral potency that they are of little use in routine oral analgesic therapy. Notable exceptions are hydrocodone and oxycodone that are absorbed into the portal system and have much smaller first-pass metabolism by the liver. Codeine has better oral efficacy because there is significantly less loss due to first-pass metabolism and much is demethylated into morphine by the liver.

Another general problem with the opioid analgesics is their relatively high incidence of nausea and central nervous system (CNS) depression, which become more intense as the dosage is increased. Mild CNS depression manifested as sedation may sometimes be useful, but ambulatory dental patients generally want to be able to function normally after they leave the dental office. As a consequence, single-entity opioid analgesics are not the drugs of choice for the management of acute pain in ambulatory patients.

Codeine is commonly used in combination analgesics. Its effective oral-dose range is 30 mg to 90 mg, with 30 mg providing only minimal additive analgesia, 60 mg providing detectable analgesia with considerably more nausea and sedation, and 90 mg approaching the dose at which intolerable side effects appear. Codeine is available in combination with aspirin or acetaminophen. For most patients, 600 mg to 650 mg of either drug combined with 60 mg of codeine should provide some additive pain relief but with delayed and variable effects due to time needed for metabolism to morphine and genetic variability in this metabolic pathway. A combination of 400 mg of ibuprofen plus 60 mg codeine in comparison with 400 mg of ibuprofen alone results in modest, if any, additive analgesia for 1 to 2 hours. As a consequence of the weak evidence of therapeutic benefit in controlled clinical trials, a combination of these two drugs has never been approved for marketing in the United States.

Analgesic combinations containing oxycodone are generally perceived as being more effective than codeine-containing combinations based on the 10-fold to 12-fold greater potency attributed to oral oxycodone in comparison to oral codeine. However, the recommended dose of oxycodone in combination with aspirin or acetaminophen is 5 mg every 6 hours and should result in the same analgesia as 50 mg to 60 mg of codeine. Combinations of oxycodone with aspirin or acetaminophen result in greater analgesia that either drugs have alone, but are no more effective than an NSAID alone, while resulting in significantly higher incidence of adverse events in ambulatory patients. The results of several clinical trials and a systematic review indicate that 5 mg of oxycodone produces additive analgesia when combined with 400 mg of ibuprofen. Combining a 10-mg dose of oxycodone with 400 mg of ibuprofen produces additive analgesia but with a dose-related increase in drowsiness, nausea, and vomiting.

Hydrocodone/acetaminophen combinations are the most widely prescribed analgesics in the United States but no published data have indicated that they result in analgesia that is comparable to a prescription dose of an NSAID. A formulation combining ibuprofen 200 mg with hydrocodone 7.5 mg is also marketed, but the 200-mg ibuprofen dose is suboptimal and no evidence suggests that the analgesia provided by this combination is greater than

| TABLE 2
<table>
<thead>
<tr>
<th>Checklist for Prescribing Opioids for Acute Dental Pain</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>PATIENT EVALUATION</strong></td>
</tr>
<tr>
<td>- History of previous opioid use and experience</td>
</tr>
<tr>
<td>- Determination of personal and family history of drug abuse</td>
</tr>
<tr>
<td>- Avoidance if currently in a recovery program</td>
</tr>
<tr>
<td>- Drug history to assess potential drug interactions</td>
</tr>
<tr>
<td><strong>MULTIMODAL PAIN-MANAGEMENT STRATEGIES</strong></td>
</tr>
<tr>
<td>- Preventive NSAIDs (ie, naproxen sodium 550 mg, ibuprofen 600 mg)</td>
</tr>
<tr>
<td>- Long-acting local anesthesis/analgesia (0.5% bupivacaine with epi)</td>
</tr>
<tr>
<td>- Corticosteroids (dexamethasone 8 mg i.m. or i.v.)</td>
</tr>
<tr>
<td><strong>PATIENT EDUCATION AND COUNSELING</strong></td>
</tr>
<tr>
<td>- Severity of pain may not require the opioid prescription to be filled</td>
</tr>
<tr>
<td>- Possibilities of adverse effects (eg, nausea, vomiting, constipation)</td>
</tr>
<tr>
<td>- Significant potential for misuse, abuse, and overdose</td>
</tr>
<tr>
<td><strong>PATIENT COUNSELING</strong></td>
</tr>
<tr>
<td>- Follow the labeled instructions</td>
</tr>
<tr>
<td>- Avoid alcohol and sedative drugs while taking an opioid pain reliever</td>
</tr>
<tr>
<td>- Never share with a friend or family member</td>
</tr>
<tr>
<td>- Secure the prescription to avoid diversion</td>
</tr>
<tr>
<td>- Dispose of unused tablets properly:</td>
</tr>
<tr>
<td>- Combine with coffee grinds or kitty litter (this makes it less likely to be stolen). Place the mixture in a sealed bag. Do not flush.</td>
</tr>
<tr>
<td>- Return tablets through Drug Take-Back Programs or check with local governments about take-back initiatives. For disposal information, visit the US Office of Diversion Control’s website at deadiversion.usdoj.gov.</td>
</tr>
</tbody>
</table>
400 mg to 600 mg of ibuprofen alone. In patients with severe pain, combining one tablet of marketed fixed-dose combination with 200 mg to 400 mg of ibuprofen for a combined dose 400 mg to 600 mg of ibuprofen with 7.5 mg of hydrocodone should produce greater analgesia (and side effects) than an NSAID alone.

Patients’ Expectations and Opioids

Although local anesthesia effectively controls procedural pain, postoperative pain is driven by expression of inflammatory mediators for 2 to 3 days following a procedure and varies highly in intensity across patients due to a wide variety of influences. Our still-limited understanding of how these factors work together is inadequate for predicting, with certainty, the intensity of the patient’s postoperative pain. The usual approach is to select an appropriate analgesic regimen based on estimated pain intensity due to the extent of tissue injury, the patient’s medical history, the etiology of the nociceptive input (eg, inflammation), the patient’s self-report of past pain experiences, and the available choice of analgesic drugs. Patients’ expectations for a comfortable postoperative recovery with minimal impact on their activities of daily living are generally biased toward receiving the most efficacious drug. However, patients do not take into account the risk for common adverse events, possible idiosyncratic reactions when taking a drug for the first time or when administered in combination with another drug, the personal impact of opioid misuse and abuse, or the societal impact of drug dependence and drug-seeking behavior.

The possible discrepancy between wide inter-individual variability in pain and patients’ expectations for minimal postoperative discomfort presents a therapeutic dilemma. For some patients, the analgesic efficacy is adequate and the incidence of adverse events minimal. However, for those with high-intensity pain, analgesic efficacy is inadequate, and for others the side-effect liability is unacceptable. The potential for exposure to opioids prescribed for therapeutic purposes leading to drug misuse and possible drug dependence is also variable and unpredictable. Given the likely discrepancy between patients’ expectations and the need to balance analgesic efficacy with side-effect liability and potential for adding to the mortality associated with opioid overdose, alternative therapeutic strategies are needed. These include individualizing the analgesic drug regimen, using interventions that attempt to prevent pain rather than “manage” it after onset, and educating patients and their significant others about risks for opioid use, especially when administered to adolescents who are more vulnerable to drug abuse.

Table 3 provides suggestions to minimize opioid misuse or diversion that can be readily implemented before considering whether to prescribe an opioid combination analgesic.

Alternatives for Acute Dental Pain

Rather than routinely administering a fixed-dose opioid combination analgesic for all patients who are expected to have moderate-to-severe pain, a flexible analgesic strategy is recommended to optimize NSAID analgesia. This may also help to minimize opioid-induced adverse events and reduce the risk for individual patients being exposed to the risk for opioid abuse, leading to possible dependence and opioid overdose death (Table 3 and Table 4). Levels of pain that might be expected after a simple extraction or endodontic treatment to a tooth that was asymptomatic prior to treatment may be effectively managed with an OTC (mild pain)
medication or a prescription dose of an NSAID such as ibuprofen, ketoprofen, or naproxen. The analgesic can be administered prior to or immediately following the procedure to delay the onset of postoperative pain, minimize its intensity following the offset of local anesthesia, and attenuate the development of sensitization leading to hyperalgesia over the 48 to 72 hours following a procedure that produces tissue injury. For a surgical procedure such as removal of impacted third molars, implant placement, and endodontic treatment of a tooth that has already been sensitized due to subacute inflammation, preventive analgesia should be initiated with use of a long-acting local anesthesia and NSAID administration prior to or immediately following the procedure. This is to suppress the formation of prostaglandin E₂ that promotes postoperative pain, swelling, and the development of sensitization leading to hyperalgesia. The NSAID should be continued for 48 to 72 hours based on the dosing interval for the NSAID prescribed. Acetaminophen 600 mg to 650 mg can be co-administered to provide additive analgesia with minimal additional side effects, either at the same time as the NSAID is administered or alternating as long as the maximum recommended dose for each drug is not exceeded. Patients can be provided with a prescription for an opioid combination drug to be filled only if pain is not adequately suppressed by the preventive regimen. Ultracet® combines acetaminophen with tramadol, which is an orally effective opioid drug.

Ultracet has much less potential for misuse than an oxycodone or hydrocodone combination. Prescribe only a 3-day supply of the Ultracet, and provide the patient and a significant other or caregiver with instructions to dispose of the reminder of the prescription when no longer needed and to not share the medication with friends or relatives (Table 3). If severe pain persists beyond 3 days, the patient should be examined for possible postoperative complications, eg, infection or alveolitis, rather than be provided with another opioid-containing prescription.⁴⁰-⁴³

Translating Evidence, Ethics, and Education

Procedures that warrant postoperative analgesic use will likely result in less discomfort for the patient and fewer side effects by shifting from traditional aspirin-opioid or acetaminophen-opioid combination to a preventive strategy. Combining two preventive regimens—the use of preoperative NSAID and long-acting local anesthetic—delays the onset of pain on the day of the procedure, lessens the intensity of pain when it does occur, and attenuates the development of sensitization leading to hyperalgesia over the 48 to 72 hours following a tissue injury-producing procedure. Due to the wide variability in pain and analgesia that exists throughout the patient population, some individuals will report pain that requires intervention. Recommending the administration of acetaminophen in addition of an NSAID results in additive analgesia without opioid-mediated side effects and without exposing the patient to a drug that can be the first step in the development of misuse, with the possible catastrophic consequence of drug dependence. In the minority of cases, not as a routine practice, providing a prescription for an opioid combination that can be filled and administered only if needed provides additional pain relief for individual patients without exposing the population of patients to the risks for opioid abuse and decreases the supply of opioids available for possible diversion. This overall approach permits individualization of analgesic therapy within the limits of currently available medications in an outpatient setting and fulfills the ethical imperative to “do no harm” in the doctor-patient interaction and at the societal level.

DISCLOSURE

The authors had no disclosures to report.

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REFERENCES


TABLE 4

Comparison of Conventional Approach to Targeted Strategies

<table>
<thead>
<tr>
<th></th>
<th>OPIOID COMBINATIONS</th>
<th>PREVENTIVE/ADDITIVE/ADAPTIVE</th>
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<tbody>
<tr>
<td>Analgesia</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>Adverse effects</td>
<td>+++</td>
<td>+</td>
</tr>
<tr>
<td>Abuse potential</td>
<td>+++</td>
<td>0 (without opioid)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>+ (with tramadol)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>++ (with oxycodone or hydrocodone)</td>
</tr>
<tr>
<td>Overdose risk</td>
<td>++</td>
<td>0 (without opioid)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>+ (with tramadol)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>++ (with oxycodone or hydrocodone)</td>
</tr>
</tbody>
</table>

Relative effects based on well-established pharmacology of drug classes and specific agents in Figure 1 ranked on a ranking of 0 to ++++. ⁴⁰-⁴³
May 11, 2016.


Prescribing Opioid Analgesics for Acute Dental Pain: Time to Change Clinical Practices in Response to Evidence and Misperceptions

Raymond A. Dionne, DDS, PhD; Sharon M. Gordon, DDS, MPH, PhD; and Paul A. Moore, DMD, PhD, MPH

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Please complete Answer Form on page 388, including your name and payment information.

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<table>
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<th>QUIZ</th>
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1. How much of opioid prescribing in the United States can be attributed to dentists?
   - A. 5%
   - B. 7%
   - C. 11%
   - D. 15%

2. Which of the following statements is true concerning non-steroidal anti-inflammatory drugs (NSAIDs) and acute pain?
   - A. NSAIDs are almost always more efficacious than opioid combinations.
   - B. NSAIDs produce a lower incidence of adverse events.
   - C. NSAIDs do not contribute to drug abuse.
   - D. all of the above

3. Opioid analgesics are effective for relieving pain by acting at:
   - A. nociceptors that react to damage.
   - B. opioid receptors in the central nervous system.
   - C. tonic receptors, which tend to react slowly.
   - D. phasic receptors, which adapt rapidly.

4. Which of the following within Indiana was identified as the leading predictor of enrollment in drug-abuse treatment programs in the state?
   - A. the prevalence of rural hospitals prescribing opioids
   - B. the ratio of physicians to patients being incongruent
   - C. the prevalence of dental practices and pharmacies
   - D. none of the above

5. Analgesics for acute dental pain:
   - A. should never be orally administered.
   - B. should produce minimal adverse events in ambulatory patients.
   - C. make ambulatory patients feel sedated.
   - D. all of the above

6. How much ibuprofen is superior to 30 mg of dihydrocodeine when evaluated in the oral-surgery model?
   - A. 200 mg
   - B. 400 mg
   - C. 600 mg
   - D. 800 mg

7. Which opioid analgesic is less likely to contribute to abuse, especially if a limited 3-day supply is prescribed without refills?
   - A. tramadol
   - B. fentanyl
   - C. meperidine
   - D. oxycodone

8. What is the recommended dose of oxycodone in combination with aspirin or acetaminophen?
   - A. 3 mg every 6 hours
   - B. 5 mg every 6 hours
   - C. 6 mg every 8 hours
   - D. 5 mg every 8 hours

9. Which of the following is NOT an alternative therapeutic strategy when managing patients’ expectations?
   - A. individualizing the analgesic drug regimen
   - B. using interventions that attempt to prevent pain rather than “manage” it after onset
   - C. educating patients and their significant others about risks for opioid use
   - D. asking patients to defer taking analgesics until pain intensifies

10. Which of the following can be done to minimize the pain from a simple tooth extraction?
    - A. advise the patient to take the analgesic on an as-needed basis
    - B. consult with the patient’s physician regarding the type of analgesia to provide
    - C. administer the analgesic prior to or immediately following the procedure
    - D. advise the patient to use cold compresses on the area in lieu of analgesic

Course is valid from 6/1/2016 to 6/30/2019. Participants must attain a score of 70% on each quiz to receive credit. Participants receiving a failing grade on any exam will be notified and permitted to take one re-examination. Participants will receive an annual report documenting their accumulated credits, and are urged to contact their own state registry boards for special CE requirements.

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