

Pancreatitis, Acute: Pain Control

What We Know

- ▶ Acute pancreatitis (AP) is a rapidly developing, potentially fatal inflammatory disorder of the pancreas, with diverse involvement of other organ systems; AP can be mild to severe, with a clinical course that varies widely from patient to patient.^(5, 6, 7) (See *Quick Lesson About... Pancreatitis, Acute*)
- The inflammation caused by dysfunctionally activated pancreatic enzymes in AP has a direct effect on sensory nerves at spinal cord level T5–T9, which results in visceral pain^(1, 6, 8)
 - Gradually increasing abdominal pain that plateaus after several hours is the primary characteristic of mild AP; pain that persists more than a few days is associated with the development of complications that characterize severe AP^(1, 4, 5)
 - ▶ Pain may radiate from the abdomen to the back or chest
 - ▶ Pain is exacerbated by eating foods high in fat or drinking alcoholic beverages, or when the patient is in a supine position
 - Although rare, painless mild AP may occur in association with postoperative states, renal transplantation, peritoneal dialysis, diabetic ketoacidosis, and shock of unknown origin
- ▶ Providing adequate pain control is an essential treatment strategy for patients with AP^(3, 6)
 - Narcotic (i.e., opioid) analgesia is usually required because alternatives (i.e., nonopioid analgesia medications) are completely ineffective in alleviating the pain of severe AP^(2, 3, 8)
 - The traditional belief that opioid analgesia causes additional pancreatic dysfunction is unsupported by clinical trial evidence^(6, 8, 9)
 - Pain management with patient-controlled analgesia (PCA) is common because oral intake is restricted; PCA-infused narcotic analgesics typically prescribed for patients with AP are^(2, 6, 8, 9)
 - morphine sulfate (Avinza, Kadian, MS Contin, MSIR, Roxanol, Astramorph PF, Duramorph, Infumorph)
 - fentanyl citrate (Sublimaze)
 - ▶ A transdermal system using a fentanyl patch (Duragesic) may be used
 - hydromorphone hydrochloride (Dilaudid)
 - meperidine hydrochloride (Demerol); although occasionally given for pain in the past, meperidine hydrochloride is currently prescribed less frequently because accumulating evidence indicates a risk of neurotoxicity with its use^(9, 10)
 - Patients with pain not relieved with a strong opioid may be given adjuvant medications (i.e., drugs with analgesic properties that have a primary indication that is not the treatment of pain)⁽¹⁾
 - The most commonly prescribed adjuvant analgesics in the treatment of patients with AP are antidepressants (e.g., selective serotonin reuptake inhibitors [SSRIs]) and anticonvulsants
- ▶ Epidural analgesia is used for pain control in some specialized healthcare facilities^(6, 9)
 - Epidural analgesia also may improve the disordered abdominal microcirculation that characterizes AP, which could reduce the rate or severity of complications (e.g., ileus, or intestinal obstruction)⁽⁶⁾
 - Although the safety and effectiveness of epidural analgesia have been established in AP, reasons for using it cautiously include that^(6, 9)
 - ▶ the procedure requires great technical skill
 - ▶ patients must meet specific criteria to qualify for this level of pain relief; patients fail to qualify if their AP disease process includes
 - coagulation abnormalities
 - systemic infection or significantly increased potential for systemic infection
 - cerebral dysfunction or the need for strong sedatives, since both factors potentially interfere with patient awareness of procedure complications

ICD-9
577.0

ICD-10
K85

ICD-10-CAN
K85

Authors

Darlene A. Strayer, RN, MBA
Tanja Schub, BS

Reviewer

Nursing Practice Council
Glendale Adventist Medical Center
Glendale, California

Editor

Diane Pravikoff, RN, PhD, FAAN
Cinahl Information Systems

September 2, 2011

- ▶ Patients who are clinically asymptomatic, without nausea, vomiting, or abdominal pain for 24 hours, and who have normal bowel sounds, are started on a clear liquid diet and monitored closely for postprandial pain.^(2, 6, 7) (See *Evidence-Based Care Sheet: Pancreatitis, Acute: Nutritional Support*)
- Timing of oral refeeding and the composition of the diet are critical factors to prevent abdominal pain recurrence. Oral feeding is associated with a 21–24 % chance of pain relapse; about half of all cases of pain relapse occur within 2 days after starting oral feeding⁽⁶⁾
 - Risk of pain relapse is higher in patients who⁽⁶⁾
 - ▶ are recovering from severe AP
 - ▶ have necrotic pancreatic tissue
- Pain control should resume immediately if the patient experiences pain after eating, and restriction of all oral intake should be resumed until the patient is again pain-free for 24 hours

What We Can Do

- ▶ Learn about acute pain control in AP so you can accurately assess your patients' personal characteristics and health education needs. Share this information with your colleagues
- ▶ Intensively monitor your AP patients' pain status and provide analgesic medications as ordered^(3, 9)
 - Educate your AP patients about the use of PCA; encourage them to use it instead of tolerating pain
 - Provide written materials, if possible, to enhance their understanding of how AP pain control does not result in addiction and is important to the recovery process
 - Explain in detail the oral restriction requirement and the milestones that must be reached before a regular diet is resumed
 - Update your patients each day to help them understand their physiological status and the rationale for treatment
 - Assess your AP patients' level of anxiety and ability to cope with a life-threatening disease; provide emotional support and request a referral to a mental health clinician for counseling, if appropriate
- ▶ Collaborate with your hospital's continuing medical education department to provide education about pain control in AP for clinicians of all specialties

Coding Matrix

References are rated in order of strength:

M	Published meta-analysis
SR	Published systematic or integrative literature review
RCT	Published research (randomized controlled trial)
R	Published research (not randomized controlled trial)
C	Case histories, case studies
G	Published guidelines
RV	Published review of the literature
RU	Published research utilization report
QI	Published quality improvement report
L	Legislation
PGR	Published government report
PFR	Published funded report
PP	Policies, procedures, protocols
X	Practice exemplars, stories, opinions
GI	General or background information/texts/reports
U	Unpublished research, reviews, poster presentations or other such materials
CP	Conference proceedings, abstracts, presentations

References

- Cruciani, R. A., & Jain, S. (2008). Pancreatic pain: A mini review. *Pancreatology*, 8(3), 230-235. **(RV)**
- Despins, L. A., Kivlahan, C., & Cox, K. R. (2005). Acute pancreatitis: Diagnosis and treatment of a potentially fatal condition. *American Journal of Nursing*, 105(11), 54-57. **(GI)**
- Fitzpatrick, E. R. (2010). Assessment and management of patients with biliary disorders. In S. C. Smeltzer, B. G. Bare, J. L. Hinkle, & K. H. Cheever (Eds.), *Brunner & Suddarth's textbook of medical-surgical nursing* (12th ed., Vol. 2, pp. 1181-1185). Philadelphia: Wolters Kluwer Health/Lippincott Williams & Wilkins. **(GI)**
- Greenberger, N. J., & Toskes, P. P. (2008). Acute and chronic pancreatitis. In A. S. Fauci, E. Braunwald, D. L. Kasper, S. L. Hauser, D. L. Longo, J. L. Jameson, et al. (Eds.), *Harrison's principles of internal medicine* (17th ed., p. 2007). New York: McGraw Hill Medical. **(GI)**
- Jha, R. K., Ma, Q., Sha, H., & Palikhe, M. (2009). Acute pancreatitis: A literature review. *Medical Science Monitor*, 15(7), RA147-RA156. **(RV)**
- Mayerle, J., Hlouschek, V., & Lerch, M. M. (2005). Current management of acute pancreatitis. *Nature Clinical Practice Gastroenterology & Hepatology*, 2(10), 473-483. **(SR)**
- Moraes, J. M. M., Felga, G. E. G., Chebli, L. A., Franco, M. B., Gomes, C. A., Gaburri, P. D. ... Chebli, J. M. F. (2010). A full solid diet as the initial meal in mild acute pancreatitis is safe and result in a shorter length of hospitalization: Results from a prospective, randomized, controlled, double-blind clinical trial. *Journal of Clinical Gastroenterology*, 44(7), 517-522. **(RCT)**
- Peiró, A. M., Martínez, J., Martínez, E., de Madaria, E., Llorens, P., Horga, J. F., & Pérez-Mateo, M. (2008). Efficacy and tolerance of metamizole versus morphine for acute pancreatitis pain. *Pancreatology*, 8(1), 25-29. **(R)**
- Renzulli, P., Jakob, S. M., Tauber, M., Candinas, D., & Gloor, B. (2005). Severe acute pancreatitis: Case-oriented discussion of interdisciplinary management. *Pancreatology*, 5(2-3), 145-156. **(RV)**
- Seifert, C. F., & Kennedy, S. (2004). Meperidine is alive and well in the new millennium. *Pharmacotherapy*, 24(6), 776-783. **(R)**