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Survivorship: Pain Version 1.2014:

Clinical Practice Guidelines in Oncology

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Crystal S. Denlinger, MD, Jennifer A. Ligibel, MD, Madhuri Are, MD, K. Scott Baker, MD, MS, Wendy Demark-Wahnefried, PhD, RD, Debra L. Friedman, MD, MS, Mindy Goldman, MD, Lee Jones, PhD, Allison King, MD, Grace H. Ku, MD, Elizabeth Kvale, MD, Terry S. Langbaum, MAS, Kristin Leonardi-Warren, RN, ND, Mary S. McCabe, RN, BS, MS, Michelle Melisko, MD, Jose G. Montoya, MD, Kathi Mooney, RN, PhD, Mary Ann Morgan, PhD, FNP-BC, Javid J. Moslehi, MD, Tracey O'Connor, MD, Linda Overholser, MD, MPH, Electra D. Paskett, PhD, Muhammad Raza, MD, Karen L. Syrjala, PhD, Susan G. Urba, MD, Mark T. Wakabayashi, MD, MPH, Phyllis Zee, MD, Nicole McMillian, MS, and Deborah Freedman-Cass, PhD

Pain in Survivors

More than one-third of posttreatment cancer survivors experience chronic pain, which often leads to psychological distress; decreased activity, motivation, and personal interactions; and an overall poor quality of life.¹⁻⁵ Pain in survivors is often ineffectively managed. Barriers to optimal pain management in cancer survivors include health care providers' lack of training, fear of side effects and addiction, and reimbursement issues.⁶

Pain has 2 predominant mechanisms: nociceptive and neuropathic.^{7,8} Injury to somatic and visceral structures and the resulting activation of nociceptors present in skin, viscera,

Please Note

Disclosures for the NCCN Survivorship Panel

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Category 2B: Based upon lower-level evidence, there is NCCN consensus that the intervention is appropriate.

Category 3: Based upon any level of evidence, there is major NCCN disagreement that the intervention is appropriate. All recommendations are category 2A unless otherwise noted.

Clinical trials: NCCN believes that the best management for any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged.

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At the beginning of each NCCN Guidelines panel meeting, panel members review all potential conflicts of interest. NCCN, in keeping with its commitment to public transparency, publishes these disclosures for panel members, staff, and NCCN itself. Individual disclosures for the NCCN Survivorship Panel members can be found on page 500. (The most recent version of these guidelines and accompanying disclosures are available on the NCCN Web site at NCCN.org.)

These guidelines are also available on the Internet. For the latest update, visit NCCN.org.

muscles, and connective tissues cause nociceptive pain. Somatic nociceptive pain is often described as sharp, throbbing, or pressure-like, and often occurs after surgical procedures. Visceral nociceptive pain is often diffuse and described as aching or cramping. Neuropathic pain is caused by injury to the peripheral or central nervous system and might be described as burning, sharp, or shooting. Neuropathic pain often occurs as a side effect of chemotherapy or radiation therapy or is caused by surgical injury to the nerves.

Screening for and Assessment of Pain

All cancer survivors should be screened for pain at regular intervals. If pain is present, the intensity should be quantified by the survivor. Because pain is inherently subjective, self-report of pain is the current standard of care for assessment. Intensity of pain should be quantified using a 0 to 10 numeric rating scale, a categoric scale, or a pictorial scale (eg, Wong-Baker FACES Pain Rating Scale).⁹⁻¹² In addition, the survivor should be asked to describe the characteristics of the pain (eg, aching, burning). Severe uncontrolled pain is a medical emergency and should be responded to promptly. An oncologic emergency also should be ruled out in these cases.

A comprehensive evaluation, as outlined in the NCCN Guidelines for Adult Cancer Pain (available at NCCN.org), is essential to ensure proper pain management. The cause and pathophysiology of the pain should be identified to determine the optimal therapeutic strategy. In addition, the survivor's goals for comfort and function should be determined.

Management of Pain

The goals of pain management are to increase comfort, maximize function, and improve quality of life. A multidisciplinary approach is recommended, with a combination of pharmacologic treatments, psychosocial and behavioral interventions, physical therapy and exercise, and interventional procedures.^{2,13,14}

The NCCN Survivorship Panel made recommendations for the management of 8 categories of cancer pain syndromes: neuropathic pain, chronic postoperative pain (ie, pain syndromes after amputation, neck dissection, mastectomy), myalgias/arthralgias, skeletal pain, myofascial pain, gastrointestinal/urinary/pelvic pain, lymphedema, and postradiation pain. Neuropathic pain commonly results from some systemic anticancer agents.¹ The incidence of chronic pain after surgical treatment varies with the type of procedure and is as high as 60% in patients treated with breast surgery and 50% in those treated with lung surgery.¹ Arthralgias, characterized by joint pain and stiffness, occur in roughly half of women taking aromatase inhibitors as adjuvant therapy for breast cancer.¹⁵ Pelvic pain often occurs after pelvic radiation, resulting from fractures, fistulae, proctitis, cystitis, dyspareunia, or enteritis.¹

Pharmacologic interventions, local therapies, psychosocial support and behavioral treatments, physical therapy and exercise, and interventional procedures are discussed. For more information about the management of cancer-related pain, please see the NCCN Guidelines for Adult Cancer Pain (to view the most recent version of these guidelines, visit NCCN.org). These guidelines include information on opioid use and pain treatment

agreements for patients at risk for medication misuse or diversion; adjuvant analgesics; and psychosocial support and behavioral interventions that may be modified to fit the individual survivor's circumstances.

Pharmacologic Interventions

Pharmacologic measures are the foundation of treatment of many of the common pain syndromes in survivors. Pharmacologic recommendations in these guidelines vary depending on the pain syndrome and include opioids, adjuvant analgesics, nonsteroidal anti-inflammatory drugs (NSAIDs), and muscle relaxants.^{2,16-18} Topical medications are discussed in "Local Therapies" (see page 497).

Opioids—Opioids may be recommended for the treatment of neuropathic, postoperative, and skeletal pain. Data on the long-term use of opioids in survivors are lacking.^{14,17,19}

The NCCN Guidelines for Adult Cancer Pain (available at NCCN.org) recommend screening survivors for risk factors of aberrant opioid use or diversion of pain medication, using a detailed patient evaluation or tools such as the Screener and Opioid Assessment for Patients with Pain-Revised (SOAPP-R) or Opioid Risk Tool (ORT), before prescribing.²⁰⁻²⁴ In addition, if opioids are deemed necessary for any survivor (regardless of aberrant use risk level), the NCCN Survivorship Panel recommends using the lowest dose possible and reevaluating the effectiveness and necessity of opioids on a regular basis. Pain treatment agreements can also be considered.²⁵

Adjuvant Analgesics—Adjuvant analgesics include antidepressants (eg, serotoninnorepinephrine reuptake inhibitors [SNRIs], tricyclic antidepressants), anticonvulsants (eg, gabapentin, pregabalin), and corticosteroids. These are recommended for the treatment of neuropathic and postoperative pain in survivors. The term *adjuvant* refers to the fact that they are often coadministered with an opioid to enhance analgesia or reduce the opioid requirement, but they may also be used as sole pain treatment. A recent systematic review found that antidepressants, anticonvulsants, other adjuvant analgesics, and opioids were all effective at reducing neuropathic pain in patients with cancer.¹⁷ Another review found that antidepressants and antiepileptics provide additional neuropathic pain relief when added to opioids in patients with cancer.²⁶

Tricyclic antidepressants have been shown to relieve neuropathic pain in the noncancer setting.^{27,28} In addition, the SNRI duloxetine was recently shown to effectively reduce pain in a multiinstitutional, randomized, double-blind, placebo-controlled crossover trial of 231 patients with painful chemotherapy-induced neuropathy.²⁹

The most commonly used anticonvulsant drugs for the treatment of cancer-related pain, gabapentin and pregabalin, have primarily been studied in noncancer neuropathy syndromes.^{30,31} Only limited data support the effectiveness of corticosteroids for cancer-related pain, and these may also have anti-inflammatory effects.³²⁻³⁴

NSAIDs—NSAIDs are recommended for the treatment of myofascial and skeletal pain and for myalgias and arthralgias. NSAIDs are nonopioid analgesics that block the biosynthesis

of prostaglandins, which are inflammatory mediators that initiate, cause, intensify, or

maintain pain. A recent systematic review found that data supporting the use of NSAIDs for control of pain in patients with advanced cancer are limited and weak, but suggest some efficacy at reducing pain and opioid dose requirement.³⁵

A discussion of contraindications and safety precautions that should be considered before prescribing NSAIDs is provided in the NCCN Guidelines for Adult Cancer Pain (to view the most recent version of these guidelines, visit NCCN.org).

Muscle Relaxants—Muscle relaxants (eg, diazepam, lorazepam, metaxalone) reduce muscle spasm and are recommended for the treatment of skeletal pain, myalgias, and arthralgias. Evidence for their efficacy in providing pain relief in the noncancer settings is limited.^{36,37} No data could be found in the setting of cancer-related pain.

Psychosocial Support and Behavioral Interventions

Cognitive interventions are aimed at enhancing a sense of control over the pain or its underlying cause. Breathing exercises, relaxation, imagery or hypnosis, and other behavioral therapies can be very useful.^{3,38-43} Psychosocial support and education should also be provided.⁴⁴ Some studies in patients with cancer suggest that psychosocial and behavioral interventions such as skills training, education, relaxation training, supportive–expressive therapy, and cognitive behavioral therapy may be effective at reducing pain.^{40,45} Hypnosis can also be considered for treatment of neuropathic pain. Overall, data support the benefit of hypnosis for controlling pain in cancer and other settings, but are lacking in the survivorship population.⁴⁶

In general, studies regarding psychosocial support and behavioral interventions for reducing pain in survivors are limited. A recent systematic review and meta-analysis assessed the efficacy of psychosocial interventions for treating pain in patients with breast cancer and survivors.⁴⁷ Although results suggest an effect, more studies are clearly needed in the survivorship population.

Physical Therapy and Exercise

Physical therapy and general exercise may also be effective for the treatment of pain in survivors, with the main goal of increasing mobility.^{3,13,48,49} Several randomized controlled trials have reported a reduction of neck and shoulder pain associated with exercise or therapy programs.⁵⁰⁻⁵² In one study, 52 survivors of head and neck cancer were randomized to a progressive-resistance exercise training (PRET) program or standard therapeutic exercise for 12 weeks.⁵² Pain scores decreased more dramatically in the PRET group (P=. 001). In another study of 66 survivors of breast cancer, those randomized to an 8-week water exercise program experienced a greater reduction of neck and shoulder pain than those randomized to usual care.⁵⁰ In addition, group exercise in the community, with trainers specifically trained to work with cancer survivors, has been shown to reduce pain and other symptoms.⁵³

Local Therapies

Local therapies, including heat, cold packs, massage, medicated creams, ointments, and patches, are recommended for the treatment of myalgias, arthralgias, and neuropathic and myofascial pain.³ Specifically, topical lidocaine, capsaicin, ketamine, and amitriptyline are recommended for treatment of some of the various cancer pain syndromes. Data are limited on the effectiveness of ketamine and amitriptyline,⁵⁴⁻⁵⁹ but the evidence for the effectiveness of lidocaine and capsaicin is stronger.^{54,56-58} In a randomized trial of 35 patients with non–cancer-related postherpetic, postoperative, or diabetes-related neuropathic pain, pain intensity was reduced with topical lidocaine but not with topical amitriptyline when compared with placebo.⁵⁷ A larger trial with a similar population of 92 patients found no effect of topical amitriptyline, ketamine, or a combination of the two.⁶⁰ Another study found that a higher dose of amitriptyline had some efficacy in reducing peripheral neuropathy, but also showed systemic effects.⁶¹ Lidocaine has been shown to reduce the severity of postherpetic neuropathy and cancer-related pain.^{62,63}

Interventional Procedures

Referral to pain management services for interventional procedures, including transcutaneous electrical nerve stimulation (TENS), intercostal nerve blocks, and dorsal column stimulation, is recommended for refractory pain in survivors. Data on the efficacy of these interventions are mainly from patients with active cancer or the noncancer setting.^{3,64} TENS is a noninvasive procedure with electrodes placed in or around the painful area.³ A recent systematic review found that data supporting the efficacy of TENS for reducing cancer-related pain are inconclusive.⁶⁵

The goal of invasive interventions, such as an intercostal nerve block, is to interrupt nerve conduction by either destroying nerves or interfering with their function.³ The data on these interventions are also limited.³

Acupuncture

Acupuncture is recommended as an option for the treatment of myofascial pain in survivors. Evidence supporting the efficacy of this technique for reducing cancer-related pain is extremely limited.^{66,67}

NCCN Survivorship Panel Members

*,a,cCrystal S. Denlinger, MD/Chair†

Fox Chase Cancer Center

*,c,dJennifer A. Ligibel, MD/Vice Chair†

Dana-Farber/Brigham and Women's Cancer Center

^fMadhuri Are, MD£

Fred & Pamela Buffett Cancer Center at The Nebraska Medical Center

^{b,e}K. Scott Baker, MD, MS€ξ

Fred Hutchinson Cancer Research Center/Seattle Cancer Care Alliance

^cWendy Demark-Wahnefried, PhD, RD \cong

University of Alabama at Birmingham Comprehensive Cancer Center

^{b,d}Debra L. Friedman, MD, MS€‡

Vanderbilt-Ingram Cancer Center

 *,g Mindy Goldman, MD Ω

UCSF Helen Diller Family Comprehensive Cancer Center

 $^{c,d}Lee$ Jones, $PhD\Pi$

Memorial Sloan-Kettering Cancer Center

^bAllison King, MD€₽‡

Siteman Cancer Center at Barnes-Jewish Hospital and Washington University School of Medicine

Grace H. Ku, MDE‡

UC San Diego Moores Cancer Center

^{b,h}Elizabeth Kvale, MD£

University of Alabama at Birmingham Comprehensive Cancer Center

^aTerry S. Langbaum, MAS¥

The Sidney Kimmel Comprehensive Cancer Center at Johns Hopkins

^gKristin Leonardi-Warren, RN, ND#

University of Colorado Cancer Center

^bMary S. McCabe, RN, BS, MS#

Memorial Sloan-Kettering Cancer Center

^{b,c,d,g}Michelle Melisko, MD[†]

UCSF Helen Diller Family Comprehensive Cancer Center

^eJose G. Montoya, $MD\Phi$

Stanford Cancer Institute

^{a,d}Kathi Mooney, RN, PhD#

Huntsman Cancer Institute at the University of Utah

^{c,e}Mary Ann Morgan, PhD, FNP-BC#

Moffitt Cancer Center

Javid J. Moslehi, MDλÞ

Dana-Farber/Brigham and Women's Cancer Center

^{d,h}Tracey O'Connor, MD[†]

Roswell Park Cancer Institute

^cLinda Overholser, MD, MPHÞ

University of Colorado Cancer Center

^cElectra D. Paskett, PhDE

The Ohio State University Comprehensive Cancer Center - James Cancer Hospital and Solove Research Institute

^{f,h}Muhammad Raza, MD‡

St. Jude Children's Research Hospital/The University of Tennessee Health Science Center

^fKaren L. Syrjala, PhDθ

Fred Hutchinson Cancer Research Center/Seattle Cancer Care Alliance

*,fSusan G. Urba, MD†£

University of Michigan Comprehensive Cancer Center

 g Mark T. Wakabayashi, MD, MPH Ω

City of Hope Comprehensive Cancer Center

*,^hPhyllis Zee, MD $\Psi\Pi$

Robert H. Lurie Comprehensive Cancer Center of Northwestern University

NCCN Staff: Nicole McMillian, MS, and Deborah Freedman-Cass, PhD

KEY:

*Writing Committee Member

Subcommittees: ^aAnxiety and Depression; ^bCognitive Function; ^cExercise; ^dFatigue; ^eImmunizations and Infections; ^fPain; ^gSexual Function; ^hSleep Disorders

Specialties: ξ Bone Marrow Transplantation; λ Cardiology; ε Epidemiology; IIExercise/ Physiology; Ω Gynecology/Gynecologic Oncology; \ddagger Hematology/Hematology Oncology; Φ Infectious Diseases; PInternal Medicine; \ddagger Medical Oncology; Ψ Neurology/Neuro-Oncology; #Nursing;; \cong Nutrition Science/Dietician; \ddagger Patient Advocacy; \bigoplus Pediatric Oncology; θ Psychiatry, Psychology, Including Health Behavior; \pounds Supportive Care Including Palliative, Pain Management, Pastoral Care, and Oncology Social Work; \P Surgery/Surgical Oncology; ω Urology

References

- Pachman DR, Barton DL, Swetz KM, Loprinzi CL. Troublesome symptoms in cancer survivors: fatigue, insomnia, neuropathy, and pain. J Clin Oncol. 2012; 30:3687–3696. [PubMed: 23008320]
- 2. Paice JA, Ferrell B. The management of cancer pain. CA Cancer J Clin. 2011; 61:157–182. [PubMed: 21543825]
- 3. Raphael J, Hester J, Ahmedzai S, et al. Cancer pain: part 2: physical, interventional and complimentary therapies; management in the community; acute, treatment-related and complex cancer pain: a perspective from the British Pain Society endorsed by the UK Association of Palliative Medicine and the Royal College of General Practitioners. Pain Med. 2010; 11:872–896. [PubMed: 20456069]
- 4. van den Beuken-van Everdingen MH, de Rijke JM, Kessels AG, et al. Prevalence of pain in patients with cancer: a systematic review of the past 40 years. Ann Oncol. 2007; 18:1437–1449. [PubMed: 17355955]
- 5. Meretoja TJ, Leidenius MH, Tasmuth T, et al. Pain at 12 months after surgery for breast cancer. JAMA. 2014; 311:90–92. [PubMed: 24381969]
- 6. Sun V, Borneman T, Piper B, et al. Barriers to pain assessment and management in cancer survivorship. J Cancer Surviv. 2008; 2:65–71. [PubMed: 18648988]
- Caraceni A, Weinstein SM. Classification of cancer pain syndromes. Oncology (Williston Park). 2001; 15:1627–1640. [PubMed: 11780704]
- Hewitt DJ. The management of pain in the oncology patient. Obstet Gynecol Clin North Am. 2001; 28:819–846. [PubMed: 11766154]
- 9. Hicks CL, von Baeyer CL, Spafford PA, et al. The Faces of Pain Scale-Revised: toward a common metric in pediatric pain measurement. Pain. 2001; 93:173–183. [PubMed: 11427329]
- Serlin RC, Mendoza TR, Nakamura Y, et al. When is cancer pain mild, moderate or severe? Grading pain severity by its interference with function. Pain. 1995; 61:277–284. [PubMed: 7659438]
- Soetenga D, Frank J, Pellino TA. Assessment of the validity and reliability of the University of Wisconsin Children's Hospital Pain scale for Preverbal and Nonverbal Children. Pediatr Nurs. 1999; 25:670–676. [PubMed: 12024390]
- Ware LJ, Epps CD, Herr K, Packard A. Evaluation of the revised faces pain scale, verbal descriptor scale, numeric rating scale, and Iowa pain thermometer in older minority adults. Pain Manag Nurs. 2006; 7:117–125. [PubMed: 16931417]
- Levy MH, Chwistek M, Mehta RS. Management of chronic pain in cancer survivors. Cancer J. 2008; 14:401–409. [PubMed: 19060605]
- Moryl N, Coyle N, Essandoh S, Glare P. Chronic pain management in cancer survivors. J Natl Compr Canc Netw. 2010; 8:1104–1110. [PubMed: 20876547]
- Crew KD, Greenlee H, Capodice J, et al. Prevalence of joint symptoms in postmenopausal women taking aromatase inhibitors for early-stage breast cancer. J Clin Oncol. 2007; 25:3877–3883. [PubMed: 17761973]

- Caraceni A, Zecca E, Bonezzi C, et al. Gabapentin for neuropathic cancer pain: a randomized controlled trial from the Gabapentin Cancer Pain Study Group. J Clin Oncol. 2004; 22:2909–2917. [PubMed: 15254060]
- 17. Jongen JL, Huijsman ML, Jessurun J, et al. The evidence for pharmacologic treatment of neuropathic cancer pain: beneficial and adverse effects. J Pain Symptom Manage. 2013
- Moore RA, Wiffen PJ, Derry S, McQuay HJ. Gabapentin for chronic neuropathic pain and fibromyalgia in adults. Cochrane Database Syst Rev. 2011:CD007938. [PubMed: 21412914]
- 19. Koyyalagunta D, Bruera E, Solanki DR, et al. A systematic review of randomized trials on the effectiveness of opioids for cancer pain. Pain Physician. 2012; 15:ES39–58. [PubMed: 22786461]
- Akbik H, Butler SF, Budman SH, et al. Validation and clinical application of the Screener and Opioid Assessment for Patients with Pain (SOAPP). J Pain Symptom Manage. 2006; 32:287–293. [PubMed: 16939853]
- 21. Butler SF, Fernandez K, Benoit C, et al. Validation of the revised Screener and Opioid Assessment for Patients with Pain (SOAPP-R). J Pain. 2008; 9:360–372. [PubMed: 18203666]
- 22. Chou R, Fanciullo GJ, Fine PG, et al. Opioids for chronic noncancer pain: prediction and identification of aberrant drug-related behaviors: a review of the evidence for an American Pain Society and American Academy of Pain Medicine clinical practice guideline. J Pain. 2009; 10:131–146. [PubMed: 19187890]
- Passik SD, Kirsh KL. The interface between pain and drug abuse and the evolution of strategies to optimize pain management while minimizing drug abuse. Exp Clin Psychopharmacol. 2008; 16:400–404. [PubMed: 18837636]
- 24. Webster LR, Webster RM. Predicting aberrant behaviors in opioid-treated patients: preliminary validation of the Opioid Risk Tool. Pain Med. 2005; 6:432–442. [PubMed: 16336480]
- 25. [Accessed November 21, 2013] Transmucosal immediate release fentanyl (TIRF) risk evaluation and mitigation strateby (REMS). Available at: http://www.fda.gov/downloads/drugs/drugsafety/ postmarketdrugsafetyinformationforpatientsandproviders/ucm289730.pdf
- Bennett MI. Effectiveness of antiepileptic or antidepressant drugs when added to opioids for cancer pain: systematic review. Palliat Med. 2010
- Finnerup NB, Sindrup SH, Jensen TS. The evidence for pharmacological treatment of neuropathic pain. Pain. 2010; 150:573–581. [PubMed: 20705215]
- Saarto T, Wiffen PJ. Antidepressants for neuropathic pain: a Cochrane review. J Neurol Neurosurg Psychiatry. 2010; 81:1372–1373. [PubMed: 20543189]
- 29. Smith EM, Pang H, Cirrincione C, et al. Effect of duloxetine on pain, function, and quality of life among patients with chemotherapy-induced painful peripheral neuropathy: a randomized clinical trial. JAMA. 2013; 309:1359–1367. [PubMed: 23549581]
- Baron R, Brunnmuller U, Brasser M, et al. Efficacy and safety of pregabalin in patients with diabetic peripheral neuropathy or postherpetic neuralgia: open-label, non-comparative, flexibledose study. Eur J Pain. 2008; 12:850–858. [PubMed: 18242109]
- Johannessen Landmark C. Antiepileptic drugs in non-epilepsy disorders: relations between mechanisms of action and clinical efficacy. CNS Drugs. 2008; 22:27–47. [PubMed: 18072813]
- 32. Leppert W, Buss T. The role of corticosteroids in the treatment of pain in cancer patients. Curr Pain Headache Rep. 2012; 16:307–313. [PubMed: 22644902]
- Mercadante SL, Berchovich M, Casuccio A, et al. A prospective randomized study of corticosteroids as adjuvant drugs to opioids in advanced cancer patients. Am J Hosp Palliat Care. 2007; 24:13–19. [PubMed: 17347500]
- Wooldridge JE, Anderson CM, Perry MC. Corticosteroids in advanced cancer. Oncology (Williston Park). 2001; 15:225–234. discussion 234-236. [PubMed: 11252935]
- 35. Nabal M, Librada S, Redondo MJ, et al. The role of paracetamol and nonsteroidal antiinflammatory drugs in addition to WHO Step III opioids in the control of pain in advanced cancer: a systematic review of the literature. Palliat Med. 2012; 26:305–312. [PubMed: 22126843]
- 36. Richards BL, Whittle SL, Buchbinder R. Muscle relaxants for pain management in rheumatoid arthritis. Cochrane Database Syst Rev. 2012; 1:CD008922. [PubMed: 22258993]

- Richards BL, Whittle SL, van der Heijde DM, Buchbinder R. The efficacy and safety of muscle relaxants in inflammatory arthritis: a Cochrane systematic review. J Rheumatol Suppl. 2012; 90:34–39. [PubMed: 22942327]
- Cassileth BR, Keefe FJ. Integrative and behavioral approaches to the treatment of cancer-related neuropathic pain. Oncologist. 2010; 15(Suppl 2):19–23. [PubMed: 20489193]
- Huang ST, Good M, Zauszniewski JA. The effectiveness of music in relieving pain in cancer patients: a randomized controlled trial. Int J Nurs Stud. 2010; 47:1354–1362. [PubMed: 20403600]
- Kwekkeboom KL, Cherwin CH, Lee JW, Wanta B. Mind-body treatments for the pain-fatiguesleep disturbance symptom cluster in persons with cancer. J Pain Symptom Manage. 2010; 39:126–138. [PubMed: 19900778]
- Montgomery GH, Weltz CR, Seltz M, Bovbjerg DH. Brief presurgery hypnosis reduces distress and pain in excisional breast biopsy patients. Int J Clin Exp Hypn. 2002; 50:17–32. [PubMed: 11778705]
- Pfister DG, Cassileth BR, Deng GE, et al. Acupuncture for pain and dysfunction after neck dissection: results of a randomized controlled trial. J Clin Oncol. 2010; 28:2565–2570. [PubMed: 20406930]
- 43. Stoelb BL, Molton IR, Jensen MP, Patterson DR. The efficacy of hypnotic analgesia in adults: a review of the literature. Contemp Hypn. 2009; 26:24–39. [PubMed: 20161034]
- 44. Keefe FJ, Abernethy AP, Campbell LC. Psychological approaches to understanding and treating disease-related pain. Annu Rev Psychol. 2005; 56:601–630. [PubMed: 15709948]
- 45. Sheinfeld Gorin S, Krebs P, Badr H, et al. Meta-analysis of psychosocial interventions to reduce pain in patients with cancer. J Clin Oncol. 2012; 30:539–547. [PubMed: 22253460]
- 46. Montgomery GH, Schnur JB, Kravits K. Hypnosis for cancer care: over 200 years young. CA Cancer J Clin. 2013; 63:31–44. [PubMed: 23168491]
- 47. Johannsen M, Farver I, Beck N, Zachariae R. The efficacy of psychosocial intervention for pain in breast cancer patients and survivors: a systematic review and meta-analysis. Breast Cancer Res Treat. 2013; 138:675–690. [PubMed: 23553565]
- Carvalho AP, Vital FM, Soares BG. Exercise interventions for shoulder dysfunction in patients treated for head and neck cancer. Cochrane Database Syst Rev. 2012; 4:CD008693. [PubMed: 22513964]
- Mishra SI, Scherer RW, Geigle PM, et al. Exercise interventions on health-related quality of life for cancer survivors. Cochrane Database Syst Rev. 2012; 8:CD007566. [PubMed: 22895961]
- Cantarero-Villanueva I, Fernandez-Lao C, Fernandez-de-Las-Penas C, et al. Effectiveness of water physical therapy on pain, pressure pain sensitivity, and myofascial trigger points in breast cancer survivors: a randomized, controlled clinical trial. Pain Med. 2012; 13:1509–1519. [PubMed: 22958507]
- 51. Fernandez-Lao C, Cantarero-Villanueva I, Fernandez-de-Las-Penas C, et al. Effectiveness of a multidimensional physical therapy program on pain, pressure hypersensitivity, and trigger points in breast cancer survivors: a randomized controlled clinical trial. Clin J Pain. 2012; 28:113–121. [PubMed: 21705873]
- McNeely ML, Parliament MB, Seikaly H, et al. Effect of exercise on upper extremity pain and dysfunction in head and neck cancer survivors: a randomized controlled trial. Cancer. 2008; 113:214–222. [PubMed: 18457329]
- Rajotte EJ, Yi JC, Baker KS, et al. Community-based exercise program effectiveness and safety for cancer survivors. J Cancer Surviv. 2012; 6:219–228. [PubMed: 22246463]
- Argoff CE. Topical analgesics in the management of acute and chronic pain. Mayo Clin Proc. 2013; 88:195–205. [PubMed: 23374622]
- 55. Barros GA, Miot HA, Braz AM, et al. Topical (S)-ketamine for pain management of postherpetic neuralgia. An Bras Dermatol. 2012; 87:504–505. [PubMed: 22714779]
- 56. Hempenstall K, Nurmikko TJ, Johnson RW, et al. Analgesic therapy in postherpetic neuralgia: a quantitative systematic review. PLoS Med. 2005; 2:e164. [PubMed: 16013891]
- Ho KY, Huh BK, White WD, et al. Topical amitriptyline versus lidocaine in the treatment of neuropathic pain. Clin J Pain. 2008; 24:51–55. [PubMed: 18180637]

- Lin PL, Fan SZ, Huang CH, et al. Analgesic effect of lidocaine patch 5% in the treatment of acute herpes zoster: a double-blind and vehicle-controlled study. Reg Anesth Pain Med. 2008; 33:320– 325. [PubMed: 18675742]
- 59. Lynch ME, Clark AJ, Sawynok J, Sullivan MJ. Topical amitriptyline and ketamine in neuropathic pain syndromes: an open-label study. J Pain. 2005; 6:644–649. [PubMed: 16202956]
- Lynch ME, Clark AJ, Sawynok J, Sullivan MJ. Topical 2% amitriptyline and 1% ketamine in neuropathic pain syndromes: a randomized, double-blind, placebo-controlled trial. Anesthesiology. 2005; 103:140–146. [PubMed: 15983466]
- 61. Kopsky DJ, Hesselink JM. High doses of topical amitriptyline in neuropathic pain: two cases and literature review. Pain Pract. 2012; 12:148–153. [PubMed: 21676162]
- 62. Fleming JA, O'Connor BD. Use of lidocaine patches for neuropathic pain in a comprehensive cancer centre. Pain Res Manag. 2009; 14:381–388. [PubMed: 19862373]
- 63. Gammaitoni AR, Alvarez NA, Galer BS. Safety and tolerability of the lidocaine patch 5%, a targeted peripheral analgesic: a review of the literature. J Clin Pharmacol. 2003; 43:111–117. [PubMed: 12616661]
- 64. Brogan S, Junkins S. Interventional therapies for the management of cancer pain. J Support Oncol. 2010; 8:52–59. [PubMed: 20464881]
- Hurlow A, Bennett MI, Robb KA, et al. Transcutaneous electric nerve stimulation (TENS) for cancer pain in adults. Cochrane Database Syst Rev. 2012; 3:CD006276. [PubMed: 22419313]
- Choi TY, Lee MS, Kim TH, et al. Acupuncture for the treatment of cancer pain: a systematic review of randomised clinical trials. Support Care Cancer. 2012; 20:1147–1158. [PubMed: 22447366]
- 67. Garcia MK, McQuade J, Haddad R, et al. Systematic review of acupuncture in cancer care: a synthesis of the evidence. J Clin Oncol. 2013; 31:952–960. [PubMed: 23341529]

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CANCER PAIN SYNDROME	TREATMENT				
Neuropathic pain	 General measures: Adjuvant analgesics (See PAIN-G from the NCCN Guidelines for Adult Cancer Pain*) Antidepressants Anticonvulsants Corticosteroids Opioids^a (See PAIN-3, PAIN-4, and PAIN-5 in the NCCN Guidelines for Adult Cancer Pain*) Cognitive behavior therapy and psychosocial support (See PAIN-4, from the NCCN Guidelines for Adult Cancer Pain*) Cognitive behavior therapy and psychosocial support (See PAIN-4, from the NCCN Guidelines for Adult Cancer Pain*) Cognitive behavior therapy and psychosocial support (See PAIN-H from the NCCN Guidelines for Adult Cancer Pain*) Consider hypnosis For refractory pain, consider referral to pain management services, interventional specialist, physical therapy, physical medicine, and/or rehabilitation Consider dorsal column stimulation Local therapies Topical patches (lidoderm, capsaicin) Creams (ketamine and amitriptyline combined) Intercostal nerve blocks Nonpharmacologic therapies Heat Ice Acupuncture 				
*To view the most recent version of these guide	elines, visit NCCN.org.				
See Principles of Opioid Use in Long-Term Survivors (SPAIN-A).					



CANCER PAIN SYNDROME	TREATMENT
Myalgias, arthralgias	 Nonpharmacologic Exercise Heat (parafili wax, hot pack) Cold pack Physical therapy Aquatic therapy Ultrasonic stimulation^c Massage Acupuncture Pharmacologic Nonsteroidal anti-inflammatory drugs (NSAIDs) Muscle relaxants Antiepileptic drugs (gabapentin, pregabalin) Serotonin-norepinephrine reuptake inhibitors (SNRIs) Tricyclic antidepressants (TCAs) Consider referral to pain management services, interventional specialist, physical therapy, physical medicine, and/or rehabilitation
Skeletal pain ^d ————————————————————————————————————	 For vertebral compression: General measures: Vitamin D NSAIDs Muscle relaxants Consider referral to pain management services, interventional specialist, physical therapy, physical medicine, and/or rehabilitation For acute vertebral compression: Opioids^a Bracing (thoracolumbar sacral orthosis [TLSO], Jewett brace) Limited bedrest Weight-bearing exercises when pain improves Physical therapy: thoracic and lumbar stabilization exercises For chronic vertebral compression: Weight-bearing exercises Physical therapy: thoracic and lumbar stabilization exercises For avascular necrosis: Physical therapy: based on weight-bearing and range-of-motion restrictions Opioids^a Muscle relaxants if myofascial component Cement augmentation for fractures For osteonecrosis of the jaw: Anticonvulsants SNRIs Opioids Consider referral to oral surgeon
^a See Principles of Opioid Use in Long-Term Surviv ^c Ultrasonic stimulation is a type of heat treatment th with multiple myeloma or bone metastases. ^d For skeletal metastases and/or bone pain, see PA SPAIN-4, SPAIN-5	ors (SPAIN-A). at can penetrate directly to the bone and should be used with caution. It is not recommended for patients N-D in the NCCN Guidelines for Adult Cancer Pain (to view the most recent version of these guidelines, visit NCCN.org).

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Clinical trials: NCCN believes that the best management of any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged. All recommendations are category 2A unless otherwise indicated

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Individual Disclosures for the NCCN Survivorship Panel

Panel Member	Clinical Research Support/Data Safety Monitoring Board	Advisory Boards, Speakers Bureau, Expert Witness, or Consultant	Patent, Equity, or Royalty	Other	Date Completed
Madhuri Are, MD	None	None	None	None	5/15/13
K. Scott Baker, MD, MS	None	None	None	None	11/22/13
Wendy Demark-Wahnefried, PhD, RD	National Cancer Institute; Harvest for Health Gardening Project for Breast Cancer Survivors; and Nutrigenomic Link between Alpha- Linolenic Acid and Aggressive Prostate Cancer	American Society of Clinical Oncology	None	American Society of Preventive Oncology	11/13/13
Crystal S. Denlinger, MD	Bayer HealthCare; ImClone Systems Incorporated; MedImmune Inc.; OncoMed Pharmaceuticals; Astex Pharmaceuticals; Merrimack Pharmaceuticals; and Pfizer Inc.	Eli Lilly and Company	None	None	1/9/14
Debra L. Friedman, MD, MS	None	None	None	None	5/26/13
Mindy Goldman, MD					Pending
Lee W. Jones, PhD	None	None	None	None	2/2/12
Allison King, MD	None	None	None	None	8/12/13
Grace H. Ku, MD	None	None	None	None	8/13/13
Elizabeth Kvale, MD	None	None	None	None	10/7/13
Terry S. Langbaum, MAS	None	None	None	None	8/13/13
Kristin Leonardi-Warren, RN, ND	None	None	None	None	1/6/14
Jennifer A. Ligibel, MD	None	None	None	None	10/3/13
Mary S. McCabe, RN, BS, MS	None	None	None	None	8/12/13
Michelle Melisko, MD	Celldex Therapeutics; and Galena Biopharma	Agendia BV; Genentech, Inc.; and Novartis Pharmaceuticals Corporation	None	None	10/11/13
Jose G. Montoya, MD	None	None	None	None	12/6/13
Kathi Mooney, RN, PhD	University of Utah	None	None	None	9/30/13
Mary Ann Morgan, PhD, FNP-BC	None	None	None	None	8/19/13
Javid J. Moslehi, MD	None	ARIAD Pharmaceuticals, Inc.; Millennium Pharmaceuticals, Inc.; Novartis Pharmaceuticals Corporation; and Pfizer Inc.	None	None	1/27/14

Panel Member	Clinical Research Support/Data Safety Monitoring Board	Advisory Boards, Speakers Bureau, Expert Witness, or Consultant	Patent, Equity, or Royalty	Other	Date Completed
Tracey O'Connor, MD	None	None	None	None	6/13/13
Linda Overholser, MD, MPH	None	Antigenics Inc.; and Colorado Central Cancer Registry Care Plan Project	None	None	10/10/13
Electra D. Paskett, PhD	Merck & Co., Inc.	None	None	None	6/13/13
Muhammad Raza, MD	None	None	None	None	8/23/12
Karen L. Syrjala, PhD	None	None	None	None	10/3/13
Susan G. Urba, MD	None	Eisai Inc.; and Helsinn Therapeutics (U.S.), Inc.	None	None	10/9/13
Mark T. Wakabayashi, MD, MPH	None	None	None	None	6/19/13
Phyllis Zee, MD	Philips/Respironics	Merck & Co., Inc.; Jazz Pharmaceuticals; Vanda Pharmaceuticals; and Purdue Pharma LP	None	None	3/26/14

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