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**NCCN Clinical Practice Guidelines in Oncology (NCCN Guidelines®)**

# Smoking Cessation

Version 2.2015

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# NCCN Guidelines Version 2.2015 Panel Members

## Smoking Cessation

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\* Peter G. Shields, MD/Chair †  
The Ohio State University Comprehensive  
Cancer Center - James Cancer Hospital  
and Solove Research Institute

\* Roy S. Herbst, MD, PhD/Vice Chair †  
Yale Cancer Center/  
Smilow Cancer Hospital

James Davis, MD/Subcommittee Lead ‡  
Duke Cancer Institute

Brian Tiep, MD/Subcommittee Lead ≡  
City of Hope Comprehensive  
Cancer Center

Douglas Arenberg, MD, MS ≡  
University of Michigan  
Comprehensive Cancer Center

Neal L. Benowitz, MD ‡ Σ  
UCSF Helen Diller Family  
Comprehensive Cancer Center

Laura Bierut, MD θ Σ  
Siteman Cancer Center at Barnes-  
Jewish Hospital and Washington  
University School of Medicine

Julie Bylund Luckart, APRN, AOCN, FNP #  
Huntsman Cancer Institute  
at the University of Utah

Paul Cinciripini, PhD θ  
The University of Texas  
MD Anderson Cancer Center

Bradley Collins, PhD θ  
Fox Chase  
Cancer Center

Sean David, MD, SM, DPhil Σ ‡  
Stanford Cancer Institute

Brian Hitsman, PhD θ  
Robert H. Lurie Comprehensive Cancer  
Center of Northwestern University

Andrew Hyland, PhD θ  
Roswell Park Cancer Institute

Margaret Lang, MSN #  
The Sidney Kimmel Comprehensive  
Cancer Center at Johns Hopkins

Scott Leischow, PhD θ  
Mayo Clinic Cancer Center

Elyse R. Park, PhD, MPH θ  
Massachusetts General Hospital  
Cancer Center

W. Thomas Purcell, MD, MBA †  
University of Colorado  
Cancer Center

Andrea Silber, MD †  
Yale Cancer Center/  
Smilow Cancer Hospital

Sharon Spencer, MD §  
University of Alabama at Birmingham  
Comprehensive Cancer Center

Tawee Tanvetyanon, MD, MPH †  
Moffitt Cancer Center

Hilary Tindle, MD, MPH ‡  
Vanderbilt-Ingram Cancer Center

Reginald Tucker-Seeley, MA, ScM, ScD θ  
Dana-Farber/Brigham and Women's  
Cancer Center

James Urbanic, MD §  
UC San Diego  
Moores Cancer Center

Benny Weksler, MD, MBA ¶  
The University of Tennessee  
Health Science Center

C. Will Wikle, NP, RN #  
Memorial Sloan Kettering  
Cancer Center

Douglas E. Wood, MD ¶  
Fred Hutchinson Cancer Research Center/  
Seattle Cancer Care Alliance

### NCCN

Jennifer Burns  
Jillian Scavone, PhD

‡ Internal medicine  
† Medical oncology  
¶ Surgery/Surgical oncology  
§ Radiotherapy/Radiation oncology  
# Nursing  
θ Psychiatry/Psychology/Behavioral science  
≡ Pulmonary medicine  
Σ Pharmacology/Pharmacogenetics  
\* Discussion Writing Committee Member

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

To find clinical trials online at NCCN Member Institutions, [click here: nccn.org/clinical\\_trials/physician.html](#).

**NCCN Categories of Evidence and Consensus:** All recommendations are category 2A unless otherwise specified.

See [NCCN Categories of Evidence and Consensus](#).

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# NCCN Guidelines Version 2.2015 Updates Smoking Cessation

Updates in Version 2.2015 of the NCCN Guidelines for Smoking Cessation from Version 1.2015 include:

## [SC-A \(2 of 2\)](#)

- References were updated.

## [MS-1](#)

- The Discussion section has been added to support the recommendations in the algorithm.

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### GENERAL PRINCIPLES OF THE SMOKING CESSATION GUIDELINES

These guidelines are focused on smoking cessation recommendations for patients with cancer. There are health benefits to smoking cessation even after a cancer diagnosis, regardless of stage or prognosis, namely improvement in cancer treatment outcomes, disease recurrence, and secondary cancers. It is never too late for patients with cancer to stop smoking cigarettes. Smoking and nicotine addiction is a chronic relapsing disorder. Patients may slip or relapse, which is expected and can be managed. Smokers with cancer often demonstrate high-level nicotine dependence. The NCCN Panel recommends that treatment plans for all smokers with cancer include the following:

1. Evidence-based pharmacotherapy,
2. Behavior therapy (counseling), and
3. Close follow-up with retreatment as needed.

#### Clinical Recommendations:

- Pharmacologic therapy is effective and recommended.
  - ▶ The two most effective pharmacotherapy agents are combination nicotine replacement therapy (NRT) and varenicline. Therapies can be combined as needed.
- Combining pharmacologic therapy and counseling is the most effective and leads to the best results for smoking cessation.
  - ▶ High-intensity behavior therapy with multiple counseling sessions is most effective, but at least a minimum of brief counseling is highly recommended.
- Smoking status should be documented in the patient health record. Patient health records should be updated at regular intervals to indicate changes in smoking status, quit attempts made, and interventions utilized.
- Smoking relapse and brief slips are common. Providers should discuss this and provide guidance and support to encourage continued smoking cessation attempts. Smoking slips are not necessarily an indication to try an alternative method. It may take more than one quit attempt with the same therapy to achieve long term cessation.

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# NCCN Guidelines Version 2.2015 Smoking Cessation

## EVALUATION AND ASSESSMENT OF PATIENT SMOKING<sup>a</sup>

### INITIAL EVALUATION<sup>b</sup>

### STATUS

Assess current cigarette smoking status of all patients with cancer:<sup>b,c,d</sup>

- Have you ever smoked cigarettes?
- Do you currently smoke cigarettes or have you smoked in the last 30 days?

Current smoker and/or those who have smoked within the last 30 days

→ [See Assessment of Current Smokers \(SC-2\)](#)

Former smoker or recently quit (>30 days since patient last smoked)

→ [See Assessment of Former Smokers \(SC-3\)](#)

Never smoked or long-term former smoker

→ Encourage patient to remain smoke-free

<sup>a</sup>For the purposes of this guideline, "smoking" refers to cigarette use.

<sup>b</sup>Initial evaluation and assessment of patient smoking may be completed by any member of the health care team, including physicians, nurses, medical assistants, health educators, or other dedicated staff.

<sup>c</sup>Smoking status should be documented in the patient health record and assessment should be repeated at every visit (less often for patients with remote smoking histories).

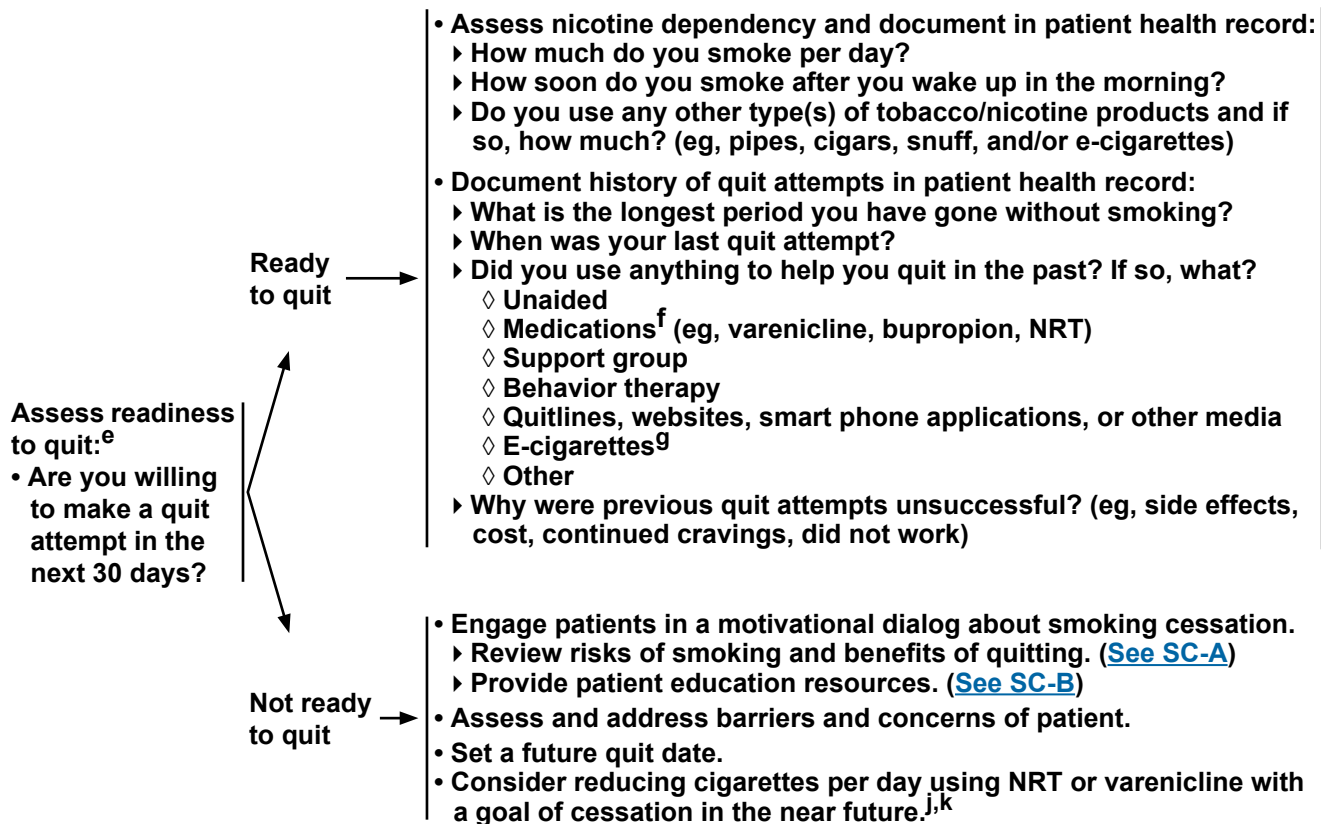
<sup>d</sup>Smoking cessation should be offered to all smokers with cancer regardless of cancer prognosis. [See Smoking-Associated Risks for Patients With Cancer \(SC-A\)](#).

**Note:** All recommendations are category 2A unless otherwise indicated.

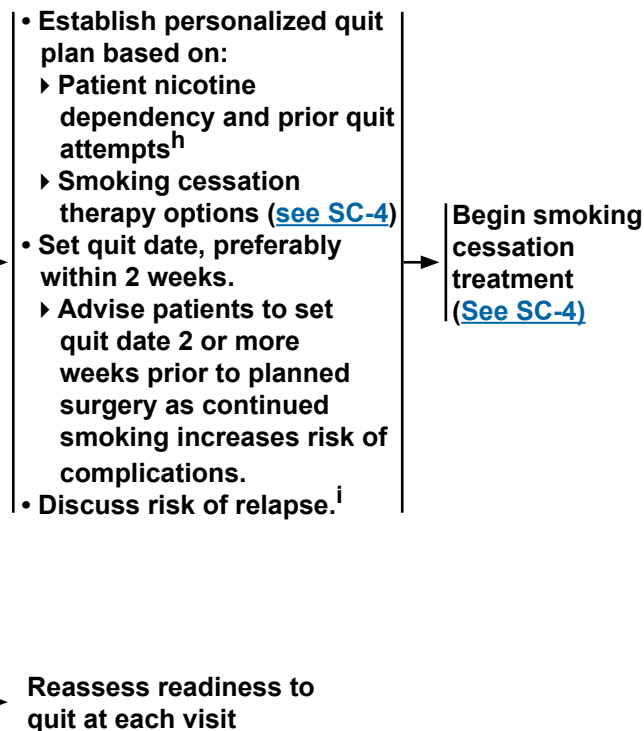
**Clinical Trials:** NCCN believes that the best management of any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged.

### CURRENT SMOKERS (Smoked Within Last 30 Days) EVALUATION AND ASSESSMENT

#### EVALUATION



#### MANAGEMENT



<sup>e</sup>Physicians and members of the health care team should discuss potential risks and benefits of quitting with each patient. Readiness to quit is to be determined by both physician and patient.

<sup>f</sup>Document type and dose of medications used during previous quit attempts.

<sup>g</sup>There is currently insufficient evidence to support the use of electronic nicotine delivery systems (e-cigarettes) in smoking cessation for patients with cancer.

<sup>h</sup>Adjustments to therapy length, intensity, and surveillance may be considered, as clinically indicated, for patients with high nicotine dependency and/or prior unsuccessful quit attempts.

<sup>i</sup>Providers should discuss risk of relapse and smoking slips and provide guidance and support to encourage continued smoking cessation attempts.

<sup>j</sup>Making an immediate quit attempt is preferred but smoking reduction may be considered with a goal of cessation. Setting a future quit date is preferred (ie. 1-3 mo).

<sup>k</sup>[See Principles of Smoking Cessation Pharmacotherapy \(SC-C\)](#)

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# NCCN Guidelines Version 2.2015

## Smoking Cessation

### FORMER SMOKERS AND RECENT QUITTERS (>30 Days Since Last Smoked) EVALUATION AND ASSESSMENT

#### EVALUATION

- Evaluate patient for risk of smoking relapse. Patients meeting 1 or more of the following criteria may be considered high risk for relapse:
  - ▶ Frequent/intense cravings
  - ▶ Elevated stress/depression<sup>l</sup>
  - ▶ Living/working with smokers
  - ▶ Time since quitting (<1 year)
  - ▶ Currently using a smoking cessation treatment (ie, pharmacotherapy, NRT)
  - ▶ Drug use/abuse (ie, marijuana, narcotics, stimulants)
- Document responses to assessment questions in patient health record.

#### STATUS

High risk for relapse<sup>i</sup>

Low risk for relapse<sup>i</sup>

#### MANAGEMENT

- For patients concerned about ability to maintain abstinence, suggest pharmacotherapy (ie, short-acting NRT)<sup>k</sup> and behavior therapy<sup>m</sup>
- Review smoking-associated risks and benefits of remaining abstinent from smoking ([See SC-A](#))
- Brief counseling for relapse risk factors and preventing relapse<sup>i</sup>
- Offer patient support resources ([See SC-B](#))
- Refer for behavior support<sup>m</sup>
- Document management and counseling plans in patient health record.

- Reinforce success and importance of remaining abstinent
- Reevaluate risk of relapse at each visit

#### RE-EVALUATION

Regularly re-evaluate smoking status and risk of relapse in person or by phone

If relapse:

- [See Assessment of Current Smokers \(SC-2\)](#)
- Refer for smoking cessation pharmacotherapy<sup>k</sup> and counseling<sup>m</sup> ([See SC-4](#))

<sup>i</sup>Providers should discuss risk of relapse and smoking slips and provide guidance and support to encourage continued smoking cessation attempts.

<sup>k</sup>[See Principles of Smoking Cessation Pharmacotherapy \(SC-C\).](#)

<sup>l</sup>Evaluate patient for psychiatric comorbidities and refer to specialist if indicated.

<sup>m</sup>[See Principles of Behavior Therapy \(SC-D\).](#)

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# NCCN Guidelines Version 2.2015

## Smoking Cessation

### GENERAL APPROACH TO SMOKING CESSATION DURING CANCER TREATMENT

#### FIRST-LINE THERAPY<sup>n,o,p</sup>

##### Pharmacotherapy<sup>k</sup> + Behavior Therapy<sup>m</sup>

- Combination NRT<sup>k,q</sup> (12 wk) + individual/group therapy (4 or more sessions)<sup>m</sup> or
- Varenicline<sup>k,r,s</sup> (12 wk) + individual/group therapy (4 or more sessions)<sup>m</sup>

#### FOLLOW-UP

- Follow-up in person or by phone within 2 weeks to assess efficacy and toxicity of pharmacotherapy.
  - ▶ May be done during individual/group therapy
- Assess risk for relapse of recent quitters and consider adjustments to dose and/or type of pharmacotherapy.<sup>k</sup>
- Encourage continued therapy and provide support for brief slips; adjusting therapy may or may not be needed.
- Additional/close follow-up during remaining therapy.

#### SURVEILLANCE

Assess smoking status in person or by phone at 12 weeks, and at the end of pharmacotherapy if longer than 12 weeks.<sup>k,p</sup>

Smoke-free

Relapse

#### ADDITIONAL THERAPY AND/OR FOLLOW-UP

- Second-line therapy:<sup>k</sup>
  - ▶ Varenicline<sup>r,s</sup> + combination NRT<sup>q</sup> or
  - ▶ Bupropion<sup>s</sup> + combination NRT<sup>q</sup>
- Continue individual/group therapy<sup>m</sup>

Additional follow-up in person or by phone at 6 and 12 months

- As clinically indicated, consider:
  - ▶ Extended use of pharmacotherapy for more than 12 weeks<sup>k,p</sup>
  - ▶ Third-line therapy<sup>k</sup>
  - ▶ Addition of more intensive or extended behavior therapy<sup>m</sup>
- Assess smoking status in person or by phone at the end of pharmacotherapy
- Additional follow-up at 6 and 12 months

<sup>k</sup>See Principles of Smoking Cessation Pharmacotherapy (SC-C).

<sup>m</sup>See Principles of Behavior Therapy (SC-D).

<sup>n</sup>Efficacy data are lacking for the use of e-cigarettes and alternative therapies (eg, hypnosis, acupuncture, nutritional supplements). Patients should be encouraged to use evidence-based cessation methods to avoid delay in achieving smoking abstinence. See SC-C (2 of 2).

<sup>o</sup>The use of marijuana, or other substances associated with smoking relapse, is discouraged for those attempting to quit smoking.

<sup>p</sup>Therapy may be extended to promote continued cessation (ie, 6 mo–1 yr) while attempting to avoid extended therapy if possible.

<sup>q</sup>Combination NRT is defined as the use of nicotine patch + short-acting NRT (gum/lozenge/inhaler/nasal spray).

<sup>r</sup>Nausea is a common side effect of varenicline and may need to be managed for patients with cancer, especially during chemotherapy.

<sup>s</sup>If prescribing varenicline or bupropion, document patient's history of mental illness or suicidal ideation.

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### SMOKING-ASSOCIATED RISKS FOR PATIENTS WITH CANCER (1 of 2)

- The 2014 Surgeon General's Report<sup>1</sup> stated that:
  - ▶ Sufficient evidence exists to support a causal relationship between smoking and adverse health outcomes, increased all-cause mortality and cancer-specific mortality, and increased risk for secondary primary cancers.
  - ▶ Existing evidence is suggestive of a link between smoking and increased risk of cancer recurrence, poor treatment response, and increased treatment-related toxicity.
- Providers should:
  - ▶ Inform patients of the potential benefits of smoking cessation, including improved survival, treatment outcomes, and health-related quality of life, as well as decreased treatment-related toxicity, drug side effects, and surgical complications.
  - ▶ Educate patients on the specific risks of smoking during treatment for their particular cancer.
  - ▶ Encourage smoking cessation as far in advance as possible before initiating cancer treatment.
  - ▶ Consider patient smoking status, prior to initiating treatment, when making decisions regarding treatment selection, dosage, and timing of initiation.

#### Treatment-Specific Risks (see [Discussion](#) for additional information)

- Smoking can impact the metabolism of chemotherapy and targeted therapy.
  - ▶ Smoking effects on cytochrome P450 enzymes may include altered drug clearance time and plasma concentration, potentially impacting the efficacy of certain drugs for patients who smoke. Providers should consider whether patients are at risk for altered drug metabolism due to smoking and determine if medication or dose adjustments may be required. Drugs whose metabolisms are known to be affected include erlotinib and irinotecan.<sup>2-4</sup>
- Smoking increases risk of radiation therapy (RT)-associated treatment complications during RT and may decrease treatment response.<sup>5-7</sup>
- Smoking is associated with increased rates of postoperative complications and mortality after cancer surgery.
  - ▶ Compared with nonsmokers, patients who smoke may experience decreased health-related quality of life after cancer surgery (eg, dyspnea, fatigue, pain).<sup>8-10</sup>
  - ▶ Smoking may impair wound healing following surgery for cancer.<sup>11,12</sup>
  - ▶ Increased infection rates, pulmonary complications, and longer postoperative hospital stays are more commonly observed in patients who smoke.<sup>13</sup>
  - ▶ Postoperative mortality rates are higher among patients who smoke.<sup>14</sup>

#### Potential Nicotine Effects on Cancer and Cardiovascular Risks (see [Discussion](#) for additional information)

- Blood nicotine levels from NRT, including combination NRT, are significantly less than from smoking cigarettes. Therefore, providers and smokers should not be dissuaded from using NRT to foster quitting and long-term cessation. The use of combination NRT as one type of pharmacotherapy is recommended.<sup>15-17</sup>
- There is insufficient evidence that NRT causes cancer in humans.<sup>18-22</sup>
- While myocardial infarction has rarely been reported in NRT users, there is insufficient evidence that NRT increases the risk of myocardial infarction or cardiovascular disease.<sup>23-25</sup>

[References on next page](#)

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**SMOKING-ASSOCIATED RISKS FOR PATIENTS WITH CANCER (2 of 2)****REFERENCES**

- <sup>1</sup>The Health Consequences of Smoking-50 Years of Progress: A Report of the Surgeon General. Atlanta (GA); 2014. Available at: <http://www.surgeongeneral.gov/library/reports/50-years-of-progress/>.
- <sup>2</sup>O'Malley M, King AN, Conte M, et al. Effects of cigarette smoking on metabolism and effectiveness of systemic therapy for lung cancer. *J Thorac Oncol* 2014;9:917-926.
- <sup>3</sup>Hamilton M, Wolf JL, Rusk J, et al. Effects of smoking on the pharmacokinetics of erlotinib. *Clin Cancer Res* 2006;12:2166-2171.
- <sup>4</sup>van der Bol JM, Mathijssen RH, Loos WJ, et al. Cigarette smoking and irinotecan treatment: pharmacokinetic interaction and effects on neutropenia. *J Clin Oncol* 2007;25:2719-2726.
- <sup>5</sup>Eifel PJ, Jhingran A, Bodurka DC, et al. Correlation of smoking history and other patient characteristics with major complications of pelvic radiation therapy for cervical cancer. *J Clin Oncol* 2002;20:3651-3657.
- <sup>6</sup>Browman GP, Wong G, Hodson I, et al. Influence of cigarette smoking on the efficacy of radiation therapy in head and neck cancer. *N Engl J Med* 1993;328:159-163.
- <sup>7</sup>Zevallos JP, Mallen MJ, Lam CY, et al. Complications of radiotherapy in laryngopharyngeal cancer: effects of a prospective smoking cessation program. *Cancer* 2009;115:4636-4644.
- <sup>8</sup>Balduyck B, Sardari Nia P, Cogen A, et al. The effect of smoking cessation on quality of life after lung cancer surgery. *Eur J Cardiothorac Surg* 2011;40:1432-1437; discussion 1437-1438.
- <sup>9</sup>Erhunmwunsee L, Onaitis MW. Smoking cessation and the success of lung cancer surgery. *Curr Oncol Rep* 2009;11:269-274.
- <sup>10</sup>Mason DP, Subramanian S, Nowicki ER, et al. Impact of smoking cessation before resection of lung cancer: a Society of Thoracic Surgeons General Thoracic Surgery Database study. *Ann Thorac Surg* 2009;88:362-370; discussion 370-361.
- <sup>11</sup>Chang DW, Reece GP, Wang B, et al. Effect of smoking on complications in patients undergoing free TRAM flap breast reconstruction. *Plast Reconstr Surg* 2000;105:2374-2380.
- <sup>12</sup>Kuri M, Nakagawa M, Tanaka H, et al. Determination of the duration of preoperative smoking cessation to improve wound healing after head and neck surgery. *Anesthesiology* 2005;102:892-896.
- <sup>13</sup>Ehlers SL, Gastineau DA, Patten CA, et al. The impact of smoking on outcomes among patients undergoing hematopoietic SCT for the treatment of acute leukemia. *Bone Marrow Transplant* 2011;46:285-290.
- <sup>14</sup>Sharma A, Deeb AP, Iannuzzi JC, et al. Tobacco smoking and postoperative outcomes after colorectal surgery. *Ann Surg* 2013;258:296-300.
- <sup>15</sup>Benowitz NL, Jacob P, 3rd, Savanapridi C. Determinants of nicotine intake while chewing nicotine polacrilex gum. *Clin Pharmacol Ther*. 1987;41(4):467-473.
- <sup>16</sup>Benowitz NL, Jacob P, 3rd, Fong I, Gupta S. Nicotine metabolic profile in man: comparison of cigarette smoking and transdermal nicotine. *J Pharmacol Exp Ther*. 1994;268(1):296-303.
- <sup>17</sup>Benowitz NL, Porchet H, Sheiner L, Jacob P., 3rd Nicotine absorption and cardiovascular effects with smokeless tobacco use: comparison with cigarettes and nicotine gum. *Clin Pharmacol Ther*. 1988;44(1):23-28.
- <sup>18</sup>Stepanov I, Carmella SG, Briggs A, et al. Presence of the carcinogen N'-nitrosonornicotine in the urine of some users of oral nicotine replacement therapy products. *Cancer Res* 2009;69:8236-8240.
- <sup>19</sup>Murray RP, Connett JE, Zapawa LM. Does nicotine replacement therapy cause cancer? Evidence from the Lung Health Study. *Nicotine Tob Res* 2009;11:1076-1082.
- <sup>20</sup>Murphy SE, von Weyarn LB, Schutten MM, et al. Chronic nicotine consumption does not influence 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone-induced lung tumorigenesis. *Cancer Prev Res (Phila)* 2011;4:1752-1760.
- <sup>21</sup>Maier CR, Hollander MC, Hobbs EA, et al. Nicotine does not enhance tumorigenesis in mutant K-ras-driven mouse models of lung cancer. *Cancer Prev Res (Phila)* 2011;4:1743-1751.
- <sup>22</sup>Shields PG. Long-term nicotine replacement therapy: cancer risk in context. *Cancer Prev Res (Phila)* 2011;4:1719-1723.
- <sup>23</sup>Hubbard R, Lewis S, Smith C, et al. Use of nicotine replacement therapy and the risk of acute myocardial infarction, stroke, and death. *Tobacco Control* 2005;14(6), 416-421.
- <sup>24</sup>Trip J. Meine, Manesh R. Patel, et al; Safety and effectiveness of transdermal nicotine patch in smokers admitted with acute coronary syndromes, *The American Journal of Cardiology* 2005; 95(8), 976-978.
- <sup>25</sup>Kimmel, JA Berlin, CM Miles, et al. Risk of acute first myocardial infarction and use of nicotine patches in a general population. *J Am Coll Cardiol*, 2001;37, 1297-1302.

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### SMOKING CESSATION RESOURCES FOR HEALTHCARE PROVIDERS AND PATIENTS (1 of 2)

Quitlines/Online Support/Mobile Apps	
American Lung Association	<ul style="list-style-type: none"> <li>• 1-800-LUNGUSA (1-800-586-4872)</li> <li>• Live Help (Online Chat)- <a href="http://www.lung.org/about-us/lung-helpline.html">http://www.lung.org/about-us/lung-helpline.html</a></li> </ul>
National Network of Tobacco Cessation Quitlines	1-800-QUIT-NOW (1-800-784-8669)
National Cancer Institute (NCI)	<ul style="list-style-type: none"> <li>• 1-877-448-7848</li> <li>• Live Help (Online Chat)- <a href="https://livehelp.cancer.gov/app/chat/chat_launch">https://livehelp.cancer.gov/app/chat/chat_launch</a></li> </ul>
Smokefree.gov	<ul style="list-style-type: none"> <li>• SmokefreeTXT (Text messaging support)- <a href="http://smokefree.gov/smokefreetxt">http://smokefree.gov/smokefreetxt</a></li> <li>• Smokefree Apps (for smartphones)- <a href="http://smokefree.gov/apps-quitstart">http://smokefree.gov/apps-quitstart</a></li> </ul>
TRICARE (For military service members and their families)	<ul style="list-style-type: none"> <li>• Quitlines: North: 1-866-459-8766; South: 1-877-414-9949; West: 1-888-713-4597</li> <li>• <a href="http://www.tricare.mil/HealthWellness/Tobacco.aspx">http://www.tricare.mil/HealthWellness/Tobacco.aspx</a></li> </ul>
Quit Tobacco: UCANQUIT2.org	<ul style="list-style-type: none"> <li>• Live chat with quit coach: <a href="http://www.ucanquit2.org">http://www.ucanquit2.org</a></li> <li>• SmokefreeMIL text message support: <a href="http://www.ucanquit2.org/en/HowToQuit/SmokefreeMIL.aspx">http://www.ucanquit2.org/en/HowToQuit/SmokefreeMIL.aspx</a></li> </ul>
General Information Online	
American Heart Association	<a href="http://www.heart.org/HEARTORG/GettingHealthy/QuitSmoking/Quit-Smoking_UCM_001085_SubHomePage.jsp">http://www.heart.org/HEARTORG/GettingHealthy/QuitSmoking/Quit-Smoking_UCM_001085_SubHomePage.jsp</a>
American Lung Association	<a href="http://www.lung.org/stop-smoking/">http://www.lung.org/stop-smoking/</a>
Centers for Disease Control and Prevention (CDC)	<a href="http://www.cdc.gov/tobacco/quit_smoking/how_to_quit/resources/index.htm">http://www.cdc.gov/tobacco/quit_smoking/how_to_quit/resources/index.htm</a>
NCI	<a href="http://www.cancer.gov/cancertopics/tobacco/smoking">http://www.cancer.gov/cancertopics/tobacco/smoking</a>
SmokeFree.gov	<a href="http://smokefree.gov">http://smokefree.gov</a>
Smoking Cessation Programs	
American Lung Association	<a href="http://www.lung.org/stop-smoking/how-to-quit/freedom-from-smoking/">http://www.lung.org/stop-smoking/how-to-quit/freedom-from-smoking/</a>
Ex: A New Way To Think About Quitting Smoking	<a href="http://www.becomeanex.org/">http://www.becomeanex.org/</a>
Guides to Quitting	
American Cancer Society (ACS)	<a href="http://www.cancer.org/healthy/stayawayfromtobacco/guidetoquittingsmoking/index">http://www.cancer.org/healthy/stayawayfromtobacco/guidetoquittingsmoking/index</a>
NCI: "Clearing the Air: Quit Smoking Today"	<a href="http://smokefree.gov/sites/default/files/pdf/clearing-the-air-accessible.pdf">http://smokefree.gov/sites/default/files/pdf/clearing-the-air-accessible.pdf</a>

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### SMOKING CESSATION RESOURCES FOR HEALTHCARE PROVIDERS AND PATIENTS (2 OF 2)

Additional Resources for Health Professionals	
American Association for Cancer Research (AACR)	<a href="http://www.aacr.org/AdvocacyPolicy/GovernmentAffairs/Documents/AACRStatementTobaccoUseCancerPatients_2013_CCR_f3f578.pdf">http://www.aacr.org/AdvocacyPolicy/GovernmentAffairs/Documents/AACRStatementTobaccoUseCancerPatients_2013_CCR_f3f578.pdf</a>
American College of Chest Physicians (ACCP)	Tobacco dependence treatment toolkit: <a href="http://tobaccodependence.chestnet.org/">http://tobaccodependence.chestnet.org/</a>
American Society of Clinical Oncology (ASCO)	Tobacco cessation and control resources: <a href="http://www.asco.org/practice-research/tobacco-cessation-and-control-resources">http://www.asco.org/practice-research/tobacco-cessation-and-control-resources</a>
Association for the Treatment of Tobacco Use and Dependence (ATTUD)	<ul style="list-style-type: none"> <li>• <a href="http://www.attud.org/">http://www.attud.org/</a></li> <li>• List of ATTUD accredited training programs: <a href="http://attudaccred.org/programs">http://attudaccred.org/programs</a></li> </ul>
NCCN Guidelines for Lung Cancer Screening	<a href="http://www.nccn.org/professionals/physician_gls/pdf/lung_screening.pdf">http://www.nccn.org/professionals/physician_gls/pdf/lung_screening.pdf</a>
NCCN Guidelines for Survivorship	<a href="http://www.nccn.org/professionals/physician_gls/pdf/survivorship.pdf">http://www.nccn.org/professionals/physician_gls/pdf/survivorship.pdf</a>
NCI- Physician Data Query: "Smoking In Cancer Care"	<a href="http://www.cancer.gov/cancertopics/pdq/supportivecare/smokingcessation/HealthProfessional">http://www.cancer.gov/cancertopics/pdq/supportivecare/smokingcessation/HealthProfessional</a>
Smokefree.gov	<a href="http://smokefree.gov/health-care-professionals">http://smokefree.gov/health-care-professionals</a>
Treatobacco.net	<a href="http://www.treatobacco.net/en/index.php">http://www.treatobacco.net/en/index.php</a>
U.S. Department of Health and Human Services- Surgeon General Reports	<a href="http://www.surgeongeneral.gov/priorities/tobacco">http://www.surgeongeneral.gov/priorities/tobacco</a>

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# NCCN Guidelines Version 2.2015

## Smoking Cessation

### PRINCIPLES OF SMOKING CESSATION PHARMACOTHERAPY (1 of 2)

#### Pharmacotherapy Options

<b>First-Line</b>	<ul style="list-style-type: none"> <li>• Combination NRT <ul style="list-style-type: none"> <li>▸ Nicotine patch + short-acting NRT (lozenge/gum/inhaler/nasal spray)</li> </ul> </li> <li>• Varenicline<sup>a</sup></li> </ul>
<b>Second-Line</b>	<ul style="list-style-type: none"> <li>• Varenicline + combination NRT</li> <li>• Bupropion + combination NRT</li> </ul>
<b>Third-Line</b>	<ul style="list-style-type: none"> <li>• Varenicline + bupropion ± NRT</li> <li>• Nortriptyline (tricyclic antidepressant)</li> <li>• Clonidine (antihypertensive, alpha-2 adrenergic receptor agonist)</li> </ul>

#### Standard Dosing Information

	Standard Dose <sup>b</sup>	Duration	Drug Warnings and Contraindications
<b>Varenicline</b>	<ul style="list-style-type: none"> <li>• Initiate dosing 1-2 wk prior to quitting</li> <li>• 0.5 mg orally, once daily on days 1–3</li> <li>• 0.5 mg orally, twice daily on days 4–7</li> <li>• 1 mg orally, twice daily from week 2–12, if tolerated</li> </ul>	12 weeks <sup>c</sup>	Providers should monitor for the development or worsening of serious neuropsychiatric issues, including those without a previous history, and discontinue use if these signs occur. See Manufacturer Black Box Warning, and weigh the substantial benefits of immediate smoking cessation versus risks of increased hostility, depression, or suicidal behavior. <sup>1</sup>
<b>Bupropion</b>	<ul style="list-style-type: none"> <li>• Initiate dosing 1-2 wk prior to quitting</li> <li>• 150 mg orally, once daily on days 1–3<sup>d</sup></li> <li>• 150 mg orally, twice daily (300 mg daily) starting on day 4, if tolerated</li> <li>• Maximum 300 mg per day</li> </ul>	7–12 weeks <sup>c</sup>	Providers should monitor for the development or worsening of serious neuropsychiatric issues, including those without a previous history, and discontinue use if these signs occur. See Manufacturer Black Box Warning, and weigh the substantial benefits of immediate smoking cessation versus risks of increased hostility, depression, or suicidal behavior. <sup>2</sup> • Contraindicated for patients with seizure risks (ie, stroke, brain metastases), those taking MOA inhibitors (increased risk of hypertensive reactions) or tamoxifen, those with closed-angle glaucoma.
<b>Combination NRT</b>	<ul style="list-style-type: none"> <li>• 21 mg patch + short-acting NRT</li> <li>• If 21 mg patch is not effective, consider increasing patch dose to 35 or 42 mg, as clinically indicated</li> </ul>	12 weeks <sup>c</sup>	Blood nicotine levels from NRT, including combination NRT, are significantly less than from smoking cigarettes. NRT is well tolerated and nicotine toxicity is rare and transient, even when used with smoking.

[Continued on next page](#)

<sup>a</sup>Nausea is a common side effect of varenicline and may need to be managed for patients with cancer, especially during chemotherapy.

<sup>b</sup>Dose adjustments may be considered, if clinically indicated.

<sup>c</sup>Therapy may be extended to promote continued cessation (ie, 6 months– 1 year) while attempting to avoid longer periods of time if possible.

<sup>d</sup>Dose adjustment for hepatic or renal insufficiency.

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### PRINCIPLES OF SMOKING CESSATION PHARMACOTHERAPY (2 of 2)

#### Side Effects of Smoking Cessation Medications:

In most circumstances the side effects related to all first-line smoking cessation medications are minimal and are considered an acceptable risk compared to smoking. Serious side effects are extremely rare. Providers should refer to manufacturer inserts for exhaustive lists of potential side effects and warnings.<sup>1,2</sup>

#### Use of E-Cigarettes and Complementary/Alternative Medicine:

- There is currently insufficient evidence to support the use of e-cigarettes in smoking cessation, because efficacy data are lacking for the use of e-cigarettes and alternative therapies (eg, hypnosis, acupuncture, nutritional supplements) alone or in combination with standard smoking cessation methods. Therefore, the use of specific alternative therapies is not recommended.
- Patients should be encouraged to use evidence-based cessation methods to avoid delay in achieving smoking abstinence. Prior unsuccessful quit attempts with conventional therapies do not justify the use of unproven alternative cessation methods. Relapse and smoking slips are common, so repeated attempts are frequently needed.
- When considering alternative therapies, providers should counsel patients on potential interactions with evidence-based cessation methods and/or cancer treatments.

<sup>1</sup>National Institutes of Health. Varenicline (Chantix) drug label and full prescribing information. Available at: <http://dailymed.nlm.nih.gov/dailymed/drugInfo.cfm?setid=d52bc40b-db7b-4243-888c-9ee95bbc6545> Accessed November 10, 2015.

<sup>2</sup>National Institutes of Health. Bupropion hydrochloride (Zyban) Drug label and full prescribing information. Available at: <http://dailymed.nlm.nih.gov/dailymed/drugInfo.cfm?setid=a3327c31-d987-40ec-b3b5-097bbf2f4f8c> Accessed November 10, 2015.

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# NCCN Guidelines Version 2.2015

## Smoking Cessation

### PRINCIPLES OF BEHAVIOR THERAPY (1 of 2)

- Pharmacotherapy is most effective when combined with behavior therapy.<sup>1</sup> In addition to the benefits of enhancing motivation and knowledge of the addiction process, behavior therapy assists patients with medication use and strategies since adherence to tobacco treatment medication recommendations is often inadequate. More intensive therapy, with a dose-response effect, is preferred over brief advice; however, brief advice, at minimum, should be delivered.
- Counseling can be in-person and/or by phone, or within a group, and include print or web-based materials. Behavior therapy, tailored somewhat to the patient's nicotine dependence and previous quit attempts, provides strategies for coping with nicotine withdrawal, identifying smoking triggers, coping with stressful and difficult situations in which smoking is likely, avoiding high-risk situations as well as addressing other patient specific barriers to and facilitators of smoking behavior change.
  - ▶ Population-level studies of real-world effectiveness of smoking cessation treatment modalities indicate that counseling by a smoking cessation specialist plus medication results in a significant improvement in cessation rates relative to no counseling or medication (OR =3.25; CI, 2.05–5.15).<sup>2</sup> Therefore, OTC medication alone, without some form of counseling may not be better than unaided cessation.
  - ▶ Intensive therapy is 4 or more behavior intervention sessions (individual or group) that are each at least 10 minutes and usually 30 minutes or more in duration. They are provided by a trained tobacco treatment specialist, with skills training, social support, and motivational interviewing as interventional components.
  - ▶ Brief advice of about 3 minutes by physicians or other health care providers results in a small but important increase in quit rates.<sup>3</sup>
  - ▶ Refer to a smoking cessation quitline, in addition to providing brief counseling from a health care provider, if face-to-face or group intervention is not available. ([See SC-B](#))

#### Motivational counseling for patients unwilling to quit:<sup>3,4,5</sup>

- Motivational counseling includes exploring the smoker's feelings, beliefs, ideas and values in order to identify areas for change towards willingness to quit. Provide reasons, ideas and needs for cessation, with encouragement. It is important to be directive with a smoker, while using an empathic approach to help the smoker understand his/her reasons for smoking and build his/her confidence to quit. The four general principles to follow are: (1) express empathy, (2) develop discrepancy, (3) roll with resistance, and (4) support self-efficacy.<sup>4</sup>

#### Behavior therapy for smoking cessation:<sup>3,5</sup>

- Through behavior therapy, smokers are provided with problem solving skills, support and encouragement. The elements include identifying risky situations (e.g., triggers for stress, drinking alcohol, being around other smokers, triggers for urges and other cues) and develop coping skills (avoid risky situations, provide cognitive strategies, short-acting NRT). Follow the same four general principles above. Note that nicotine withdrawal symptoms typically peak within 1-2 weeks after quitting and then subside.
- In smokers with cancer, there is a high incidence of depression, anxiety, and stress, all of which are common causes of relapse. It may be optimal to enroll patients in a behavior therapy program with specific interventions designed to ameliorate these conditions and other cancer-related relapse challenges. This may require referral to specialized smoking cessation programs that have staff trained to treat mental health disorders, or referral to behavior therapists who have expertise in treating co-morbid substance dependence and mental health disorders.
- Specialized treatment centers may consider providing smoking cessation therapy targeted specifically to patients with cancer (eg, individual therapy and group support that focuses on challenges specific to cancer survival and treatment) with access to counselors or group leaders experienced in the treatment of patients with cancer.

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[References on next page](#)





### PRINCIPLES OF BEHAVIOR THERAPY (2 of 2)

#### REFERENCES

- <sup>1</sup>Stead LF, Lancaster T. Combined pharmacotherapy and behavioural interventions for smoking cessation. Cochrane Database Syst Rev 2012;10:Cd008286.
- <sup>2</sup>Kotz D, Brown J, West R. 'Real-world' effectiveness of smoking cessation treatments: a population study. Addiction. 2014;109:491-499.
- <sup>3</sup>Fiore MC, Jaen CR, Baker TB, et al. Treating Tobacco Use and Dependence: 2008 Update, Clinical Practice Guideline. Rockville, MD: U.S. Department of Health and Human Services. Public Health Service, 2008. (Treating Tobacco Use and Dependence. April 2013. Agency for Healthcare Research and Quality, Rockville, MD. <http://www.ahrq.gov/professionals/clinicians-providers/guidelines-recommendations/tobacco/clinicians/update/index.html>.)
- <sup>4</sup>Miller WR, Rollnick S. Motivational Interviewing: Preparing People for Change. New York, NY: Guilford Press; 2002.
- <sup>5</sup>Lindson-Hawley N, Thompson TP, Begh R. Motivational interviewing for smoking cessation. Cochrane Database Syst Rev 2015, in press;3:Cd006936.

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## Discussion

### NCCN Categories of Evidence and Consensus

**Category 1:** Based upon high-level evidence, there is uniform NCCN consensus that the intervention is appropriate.

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

**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.**

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# NCCN Guidelines Version 2.2015

## Smoking Cessation

### Overview

An estimated 42.1 million adults in the United States currently smoke cigarettes, accounting for approximately 18% of the adult population.<sup>1</sup> Cigarette smoking results in over 480,000 premature deaths yearly, and 1 in every 5 deaths is smoking-related.<sup>2</sup> Tobacco smoking has been implicated in causing cancers of mouth, lips, nose, sinuses, larynx, pharynx, esophagus, stomach, pancreas, kidney, bladder, uterus, cervix, colon/rectum, ovary, and myeloid leukemia.<sup>3</sup> Moreover, tobacco use causes at least 30% of all cancer deaths.<sup>3</sup> These guidelines emphasize the importance of smoking cessation in all patients with cancer and seek to establish evidence-based, standard-of-care recommendations tailored to the unique needs and concerns of patients with cancer.

In a 2014 report from the Surgeon General, “The Health Consequences of Smoking—50 Years of Progress,” a comprehensive review of the evidence revealed the following important findings for patients with cancer and cancer survivors:<sup>2</sup>

- Sufficient evidence exists to infer a causal relationship between cigarette smoking and adverse health outcomes, increased all-cause mortality (ACM), and cancer-specific mortality.
- Sufficient evidence exists to infer a causal relationship between cigarette smoking and second primary cancer.
- Evidence is suggestive of a causal relationship between cigarette smoking and risk of recurrence, poorer response to treatment, and increased treatment-related toxicity.

Although the harmful effects of smoking after a cancer diagnosis have been clearly demonstrated, many patients continue to smoke cigarettes during treatment and beyond. The prevalence of continued smoking

among those who have received a cancer diagnosis has been examined in recent studies. The rate of smoking post-diagnosis varies widely by cancer type and with other factors such as gender, race, and age.

### Smoking Prevalence in Patients with Cancer

Using data from the National Health and Nutrition Examination Survey (NHANES), of the 566 cancer survivors who regularly smoked prior to their diagnosis, 64% continued to smoke post-diagnosis. Those identified at higher risk for continued smoking included female, younger, and Hispanic individuals.<sup>4</sup> In the Cancer Care Outcomes Research and Surveillance (CanCORS) cohort of patients with lung (n = 2456) and colorectal (n = 3063) cancers, 90% of patients with lung cancers and 55% of patients with colorectal cancer reported a history of smoking. At diagnosis, 39% of those with lung and 14% of those with colorectal cancer were current smokers, and of these individuals, 14% of patients with lung and 9% of patients with colorectal cancer continued to smoke at 5 months post-diagnosis.<sup>5</sup>

Smoking often persists beyond cancer treatment and well into survivorship. In a population-based study using the Behavioral Risk Factor Surveillance System, survivors of tobacco-related cancers had a smoking prevalence of 27% compared with 16% and 18% for other cancer survivors and those without a history of cancer, respectively.<sup>6</sup> Based on prospectively collected data from 772 individuals with cancer in the Cancer Prevention Study-II Nutrition Cohort, persistent smoking was observed in 68.7% and 57% of the cohort at 2 and 4 years post-diagnosis, respectively.<sup>7</sup> One study revealed smoking prevalence to be highest among survivors of bladder, lung, and ovarian cancers.<sup>8</sup> Other studies have found high prevalence of continued smoking in survivors of cervical cancers.<sup>9</sup>

### Health Care Community Response

Given the adverse health effects and prevalence of smoking in patients with cancer and survivors, several leading national organizations have called upon the oncology community for improved smoking cessation efforts. In 2013, the American Association for Cancer Research (AACR) released a policy statement calling for provision of evidence-based smoking cessation assistance to all patients with cancer along, outlining the following objectives:

- “Universal assessment and documentation of tobacco use by cancer patients in all clinical settings;
- Development of universal standards for measurement of tobacco use and exposure in clinical and research settings;
- Incorporation of evidence-based tobacco interventions into review criteria used by research and health care quality and accreditation bodies; and
- Recognition and support of the value of tobacco cessation interventions by health systems, payers, and research funders through provision of appropriate incentives for infrastructure development and intervention delivery.”<sup>10</sup>

Additionally, in a recent policy statement update, the American Society for Clinical Oncology (ASCO) called upon oncology professionals to treat tobacco dependence as aggressively and compassionately as cancer and to advocate for the wide availability of tobacco cessation services.<sup>11</sup>

However, despite general consensus on the importance of smoking cessation, particularly for patients with cancer, many cancer centers and oncology practices report that they fall short of providing consistent, high-quality smoking cessation services. In a survey of 58 National

Cancer Institute (NCI)-designated cancer centers, 20% reported offering no smoking cessation services for their patients, 38% did not routinely provide tobacco education materials to patients, and only half reported that they effectively identified tobacco use in their patients.<sup>12</sup> The American Association for Cancer Research (AACR) Taskforce on Tobacco found that few cancer care institutions utilize systematic and consistent mechanisms to foster cessation among patients with cancer.<sup>10</sup>

Data from large surveys of oncologists practicing in academic medical centers, non-academic hospitals, and oncology practices depict generally high rates of smoking assessment and provision of initial advice to quit.<sup>13-16</sup> However, smoking assessment rates were weaker outside of the academic/university setting (ie, for those practicing in a hospital-based, nonacademic or private setting).<sup>16</sup> Regardless of work setting, only 30% to 44% of respondents reported discussing specific interventions or providing subsequent follow-up. Moreover, the majority of respondents report inadequate training and/or a lack of confidence in ability to provide effective smoking cessation counseling and intervention.<sup>13-16</sup> A dearth of smoking assessment and documentation has also been demonstrated in oncology trials.<sup>17,18</sup>

Issues regarding insurance coverage and provider reimbursement for smoking cessation assessment, counseling, and cessation aids have also presented a challenge for the oncology community in the past. However, implementation of the Affordable Care Act has led to changes designed to increase access to smoking cessation interventions.<sup>19</sup>

### Barriers to Smoking Cessation in Oncology Patients

Although over 68.8% of current smokers in the United States express a desire to quit and 52.5% report making a quit attempt within the past year, only 6.2% report recent smoking cessation.<sup>20</sup> In the general



population, individuals who smoke report a number of different barriers to quitting, including stress; dependence; home, work, and social environmental factors; and lack of resources and support for quitting.<sup>21</sup> Importantly, patients and providers in the oncology setting face additional life challenges that can amplify the magnitude of these barriers.

Surveys of oncology providers have identified common themes among barriers to smoking cessation for patients with cancer. Inadequate provider training and lack of time are often cited by oncology providers as barriers to successful intervention.<sup>14</sup> Providers have also cited patient-related factors such as inability to quit, lack of motivation, or resistance to treatment.<sup>13,14</sup> However, guilt over smoking, fear of stigmatization, and fatalism regarding disease may also represent obstacles unique to oncology patients, particularly those with advanced disease.<sup>22-25</sup>

Notably, clinical trial research on smoking cessation for patients with cancer is limited, particularly for patients thought to have non-tobacco-related cancers. Barriers that limit or prevent enrollment in smoking cessation trials include smoking rate, medical history, contraindicated medications, lack of interest, and language barriers.<sup>26</sup>

Given the complexity of smoking cessation interventions for patients with cancer, there is a great need for resources that provide guidance on smoking cessation specifically for this patient population. The inaugural NCCN Guidelines for Smoking Cessation have been created to establish a standard of care for smoking cessation in patients with cancer. The NCCN Guidelines Panel has developed these guidelines in order to facilitate implementation of this standard, to allow for quality control monitoring, to fill a gap among existing treatment guidelines, and ultimately, to improve the health and outcomes for patients with cancer.

## Literature Search Criteria and Guidelines Update Methodology

Prior to the development of this inaugural version of the NCCN Guidelines® for Smoking Cessation, an electronic search of the PubMed database was performed to obtain key literature in smoking cessation for patients with cancer, published through May 2015, using the following search terms: (cancer[Title/Abstract] OR cancer patient[Title/Abstract]) AND (tobacco[Title/Abstract] OR smoking[Title/Abstract]) AND (cessation[Title/Abstract] OR quitting[Title/Abstract]). The PubMed database was chosen as it remains the most widely used resource for medical literature and indexes only peer-reviewed biomedical literature.<sup>27</sup>

The search results were narrowed by selecting studies in humans published in English. Results were confined to the following article types: Clinical Trial; Practice Guideline; Randomized Controlled Trial; Meta-Analysis; Systematic Reviews; and Validation Studies. The PubMed search resulted in **345** citations and their potential relevance was examined. The data from key PubMed articles selected by the panel for review during the Guidelines update meeting as well as articles from additional sources deemed as relevant to these Guidelines and discussed by the panel have been included in this version of the Discussion section (eg, e-publications ahead of print, meeting abstracts). Recommendations for which high-level evidence is lacking are based on the panel's review of lower-level evidence and expert opinion.

The complete details of the Development and Update of the NCCN Guidelines are available on the NCCN [webpage](#).



## General Principles of the Smoking Cessation Guidelines

These Guidelines are focused on smoking cessation recommendations for patients with cancer. Smoking cessation has health benefits even after a cancer diagnosis, regardless of stage or prognosis, namely improvement in cancer treatment outcomes, disease recurrence, and secondary cancers. Importantly, a diagnosis of cancer may present a teachable moment and valuable opportunity for providers to encourage smoking cessation.<sup>7,28</sup> It is the view of the NCCN Guidelines Panel that it is never too late for patients with cancer at any stage to stop smoking cigarettes.

Because smokers with cancer often demonstrate high-level nicotine dependence, the panel recommends a multimodal approach to cessation therapy. The NCCN Guidelines Panel recommends that treatment plans for all smokers with cancer include the following 3 tenets: behavior therapy (counseling), which can be brief; evidence-based pharmacotherapy; and close follow-up with retreatment as needed.

The panel asserts that a smoking cessation approach combining pharmacologic therapy and counseling is the most effective and leads to the best results for smoking cessation. The two most effective pharmacotherapies are combination nicotine replacement therapy (NRT) and varenicline. There is a dose-response relationship for the success of counseling; high-intensity behavior therapy with multiple counseling sessions is most effective, but at least a minimum of brief counseling is needed and effective.

The panel also emphasizes that importance of documenting smoking status and treatment plans in the patient health record. Patient health

records should be updated at regular intervals to indicate changes in smoking status, quit attempts made, and interventions utilized.

Finally, the panel emphasizes that smoking relapse and brief slips are common. Providers should discuss this with patients and provide guidance and support to encourage continued smoking cessation attempts despite slips. Additionally, providers should be aware that smoking slips do not necessarily indicate a need for an alternative intervention. More than one quit attempt with the same therapy may be necessary to achieve long-term cessation.

## Smoking-Associated Risks for Patients with Cancer

Per the Surgeon General's Report, *The Health Consequences of Smoking—50 Years of Progress*<sup>2</sup>, sufficient evidence exists to support a causal relationship between smoking and adverse health outcomes, increased ACM and cancer-specific mortality, and increased risk for secondary primary cancers. Additionally, existing evidence is suggestive of a link between smoking and increased risk of cancer recurrence, poor treatment response, and increased treatment-related toxicity.

NCCN recommends that providers should inform patients of the potential benefits of smoking cessation, including improved survival, treatment outcomes, and health-related quality of life, as well as decreased treatment-related toxicity, drug side effects, and surgical complications. Patients should receive education on the specific risks of smoking during treatment for their particular cancer and should be encouraged to stop smoking as far in advance as possible before initiating cancer treatment. Prior to initiating treatment, when making decisions regarding treatment selection, dosage, and timing of initiation, providers should consider patient smoking status and potential smoking-related effects.

### Overall Survival and Mortality

Smoking has been linked not only to the development of disease in tobacco-related cancers, but also to prognosis upon diagnosis and risk of death during treatment. Evidence suggests that current smoking increases risk of death and negatively impacts survival for patients with cancer in a variety of disease sites, including bladder,<sup>29,30</sup> breast,<sup>31</sup> cervix,<sup>32,33</sup> colon/rectum,<sup>34</sup> endometrium,<sup>35</sup> esophagus,<sup>36,37</sup> head and neck,<sup>38,39</sup> kidney,<sup>40,41</sup> lung,<sup>42-45</sup> pancreas,<sup>46</sup> and prostate,<sup>47</sup> as well as hematologic malignancies.<sup>48</sup>

Prospective studies of smoking at cancer diagnosis offer insight into the negative effects of smoking on overall survival (OS), disease-specific mortality (DSM), and ACM. In the 2014 Surgeon General's Report, a comprehensive review of the data assessing smoking and ACM in patients with cancer revealed that 87% of the studies (139/159) indicated increased risk, while 62% of all studies (99/159) demonstrated a statistically significant risk increase.<sup>2</sup> Additionally, over half of the reviewed studies found at least a 50% increase in risk of death.<sup>2</sup> Additionally, among the studies examining OS in patients with cancer who smoke, 77% (48/62) were indicative of shortened survival, with 42% (26/62) revealing statistical significance.<sup>2</sup> Finally, smoking in patients with cancer was associated with higher DSM in 79% of studies reviewed (46/58), with a statistically significant link between cancer-related mortality and patients' smoking status in 59% (34/59).<sup>2</sup>

### Risk of Recurrence or Secondary Primary Tumor

A number of studies have linked cigarette smoking and heightened risk of recurrence (ie, recurrent cancer in the same anatomic location as the original primary cancer). The 2014 Surgeon General's Report identified a positive association between smoking and risk of recurrence in 82% of the reviewed studies (42/51), with 53% of studies revealing

significantly increased risk.<sup>2</sup> Among the studies that compared relative risk (RR) of recurrence between never smokers, former smokers, and current smokers, the median RR was 1.42 and 1.15 for current and former smokers, respectively.<sup>2</sup>

Disease sites with data linking current patient smoking to increased risk of recurrence include the anus,<sup>49</sup> bladder,<sup>29,50</sup> breast,<sup>51</sup> lung,<sup>42</sup> stomach,<sup>52</sup> and prostate.<sup>47,53,54</sup>

Studies have also examined the impact of continued smoking in patients with cancer on the risk of second primary tumor formation. The 2014 Surgeon General's Report identified a positive association between smoking and risk of second primary tumor in all studies examined (n = 26). The association was strongest when considering the effects of smoking on RR of developing a smoking-related second primary cancer (eg, lung cancer). Among 5 studies classifying smoking status into "never," "former," and "current," the median elevated RR of a second primary tumor was 1.20 and 2.20 for former and current smokers, respectively. Additionally, data also suggest that smoking interacts synergistically with radiation therapy (RT) to elevate the risk of second primary cancers.<sup>2,55,56</sup>

### Smoking-Related Effects on Treatment Efficacy, Side Effects, and Outcomes

A majority of the existing data establishes and supports the detrimental impact of persistent smoking during cancer treatment. In a 2014 report from the Surgeon General, 80% of the evaluated studies (66/82) demonstrated a statistically significant association between active smoking and increased anticancer treatment-related toxicity.<sup>2</sup> Smoking has implications across the spectrum of cancer treatment, including surgical outcomes, RT efficacy and toxicity, chemotherapy metabolism

and side effects, and overall symptom burden. This discussion also addresses the developing evidence base for the benefits of smoking cessation after receiving a cancer diagnosis.

### **Smoking-Associated Risks**

#### *Surgery*

Smoking has been shown to negatively impact outcomes from cancer surgery, affecting postoperative complications, quality of life, length of hospital stay, and mortality risk.

In lung cancer, studies show that smoking impacts the success of surgical resection, decreases postoperative quality of life, and increases persistent dyspnea and thoracic pain at 12 months postoperatively.<sup>57,58</sup> Analysis of data from 7990 patients who had primary resections for lung cancer (Society of Thoracic Surgeons General Thoracic Surgery Database) revealed increased risk of hospital death and pulmonary complications associated with smoking.<sup>59</sup>

The adverse effects of smoking on postoperative outcomes was examined in over 20,000 patients with gastrointestinal (n = 12,432), lung (n = 4490), and urinary tract cancers (n = 3491) using the Veterans Health Administration Surgical Quality Improvement Program (VASQIP) database for 2002–2008.<sup>60</sup> Surgical complications examined included surgical site infections, vascular complications (ie, venous thromboembolism, stroke/cerebrovascular accident, myocardial infarction), and composite pulmonary outcomes (CPO: pneumonia, failure to wean from ventilator >48 hours, or re-intubation for cardio-respiratory failure). Across all three cohorts, never smokers had fewer complications than former and current smokers. Compared with prior smokers, current smokers in the gastrointestinal cancer cohort had higher postoperative rates of pneumonia, failure to wean from ventilator,

reintubation, and CPO. In the lung cancer cohort, current smokers had higher rates of pneumonia, failure to wean from the ventilator, reintubation, CPO, and return to surgery compared with former smokers. Current smoking status was associated with an increased length of hospital stay across all cancer sites when compared with never smokers; never smokers and prior smokers did not differ on this measure.

Postsurgical outcomes (ie, incisional infections, infectious and major complications, mortality at 30 days) were compared between cohorts of never smokers, former smokers, and current smokers using data from over 26,000 patients with colorectal cancer in the American College of Surgeon's National Surgical Quality Improvement Program database (2005–2010). Postoperative morbidity and mortality rates were higher among current smokers and a significant dose-dependent effect was observed when stratifying risk of major complications by pack-years of smoking.<sup>61</sup>

In patients undergoing hematopoietic stem cell transplantation to treat acute leukemias, pulmonary complications and longer postoperative hospital stays were more commonly observed in patients who smoked.<sup>62</sup>

Smoking can also impair wound healing and predispose patients to surgical complications for those undergoing reconstructive surgeries after cancer treatment. Among patients with breast cancer who underwent transverse rectus abdominis myocutaneous (TRAM) flap breast reconstruction surgery, smoking was associated with significantly higher risk of flap complications and delayed healing,<sup>63,64</sup> and evidence suggested that complication risk was reduced by smoking cessation of at least 4 weeks prior to surgery.<sup>63</sup> In patients with stage III or IV squamous cell carcinoma of the head and neck, serum cotinine





concentration was dose-dependently linked to increased risk of wound complications following reconstructive head and neck surgery.<sup>65</sup>

### *Radiation*

Studies have shown that prior smoking and active smoking during RT may decrease treatment response and increase complication rates, particularly for patients with head and neck cancers, but also in cervical, lung, breast, or prostate cancers.

In patients with head and neck cancer receiving RT, current smokers had poorer rates of locoregional control.<sup>38,67</sup> In another cohort, patients with head and neck cancer who continued to smoke during RT had lower rates of complete response and worse survival times than nonsmokers or those who quit prior to treatment.<sup>68</sup> Continued smoking during RT in patients with head and neck cancer has also been shown to increase the rates of treatment-related complications. In patients with laryngopharyngeal cancers, smoking during treatment was associated with significantly elevated incidence of osteoradionecrosis and hospitalization during treatment.<sup>69</sup> Another study demonstrated a significantly greater decline in several health-related quality-of-life measures in patients who continued to smoke during therapy compared with patients who quit beforehand.<sup>70</sup>

Among 3,489 patients receiving RT as part of treatment for stage I or II cervical cancer, heavy smoking (defined as at least 1 pack/day) was the strongest independent factor in predicting long-term major bladder, rectal, or small bowel complications, with even light/moderate smoking (less than 1 pack/day) predisposing patients to small bowel complications.<sup>71</sup> In another study of 565 patients with cervical cancer who were receiving primary RT, patients who smoked during treatment had lower cure rates, higher frequency of RT side effects, and higher rates of severe, irreversible complications.<sup>72</sup>

Smoking during RT for non-small cell lung cancer was associated with significantly decreased locoregional control.<sup>73</sup> Active smoking may also decrease the efficacy of RT for prostate cancers,<sup>74</sup> and increase the prevalence of long-lasting treatment-related effects on the bowel and anal sphincter.<sup>75</sup> Concurrent smoking and RT increased the risk of cardiovascular disease in 4414 10-year survivors of breast cancer.<sup>76</sup>

### *Chemotherapy/Systemic Therapy*

Data on the impact of smoking on chemotherapy are much more limited than that for surgery and RT, in part because smoking quantity during treatment is often left out of the medical record.<sup>17</sup> Many of the purported effects of smoking during chemotherapy are extrapolated from what is known about the impact of chemotherapy and smoking as individual factors of health. Smoking has the potential to exacerbate the risk of anticancer drug-related pulmonary and cardiac toxicities such as cardiomyopathy and pulmonary fibrosis.<sup>77</sup> Combining neoplastic agents with radiation while smoking may present further toxicity. Additionally, cancer drug side effects such as weight loss, cachexia, and fatigue may also be increased by smoking during treatment.<sup>77,78</sup>

Many systemic anticancer agents result in some degree of immune suppression/compromise, and smoking during chemotherapy may further compromise immune function in an already vulnerable patient population.<sup>17,77</sup> Preclinical and clinical studies suggest that smoking and nicotine exposure can be detrimental to the function of both the adaptive<sup>79-82</sup> and innate immune system.<sup>83-87</sup> Similarly, cigarette smoking may increase the incidence of infection, particularly for smoking-related infectious diseases such as pneumonia and influenza.<sup>88</sup>

Preclinical studies also suggest a potential link between nicotine exposure from smoking and the development of chemoresistance,<sup>89-93</sup>

although no clinical data are currently available to support these findings.

Smoking can also impact the metabolism of certain cytotoxic chemotherapies and other systemic therapy. Smoking effects on cytochrome P450 enzymes may alter drug clearance time and plasma concentration, potentially impacting the efficacy of certain drugs for patients who smoke. Providers should consider whether patients are at risk for altered drug metabolism due to smoking and determine if medication or dose adjustments may be required.

Drugs with metabolisms that are known to be affected include erlotinib and irinotecan. Rapid drug clearance has been observed in smokers who were receiving erlotinib therapy, such that higher doses may be required to achieve equivalent systemic exposure to standard dosing in nonsmokers.<sup>94,95</sup> Similarly, smoking increases the clearance time of irinotecan, potentially lessening systemic exposure.<sup>94,96</sup> Given the narrow therapeutic index of systemic therapy for lung cancer, small changes in drug exposure due to smoking could affect treatment efficacy and patient outcomes.<sup>94</sup> However, smoking does not appear to alter the pharmacokinetic properties of taxane chemotherapeutics (eg, docetaxel, paclitaxel) despite its paradoxical protective effects on drug-induced neutropenia and leukopenia.<sup>97</sup>

### *Symptom Burden*

In a study of 947 patients who were undergoing chemotherapy and/or RT, smoking during treatment was linked to a higher overall burden of symptoms commonly experienced among patients with cancer. In analyses that controlled for age, gender, race, education, occupation, treatment, cancer site, and Karnofsky performance score, current smokers had a significantly higher symptom burden compared with nonsmokers, both during treatment and 6 months afterwards.<sup>98</sup>

Additional studies suggest that current smokers with cancer may experience more severe or frequent pain than nonsmoking counterparts.<sup>99-102</sup>

### ***Benefits of Smoking Cessation for Patients with a Cancer Diagnosis***

For many smokers, the benefits of smoking cessation can be appreciated immediately through reduced blood carbon monoxide levels, decreased irritative respiratory symptoms (eg, cough, shortness of breath), and improved lung function. In the long term, cessation is associated with reduced risk of smoking-related disease, development of malignancy, and smoking-related mortality.<sup>103</sup> Although the deleterious effects of smoking after a cancer diagnosis are well documented and understood, research on the benefits of cessation post-diagnosis is much more limited.<sup>10,104</sup> For patients with cancer, the potential benefits and risk reductions associated with cessation are of critical importance.

Studies have begun to assess the impact of smoking cessation at or near the time of a cancer diagnosis by comparing outcomes of patients who continue to smoke during cancer treatment to those who quit prior to treatment (“recent quitters”). Studies generally show that recent quitters have survival outcomes intermediate to that of never-smokers and current-smokers, suggesting a measurable benefit of cessation post-diagnosis. The data to support this survival pattern are derived primarily from cohorts of patients with lung or head and neck cancers,<sup>42,44,68,105,106</sup> but similar patterns have been observed for other disease sites.<sup>44,107</sup>

A prospective longitudinal study examined the impact of smoking at diagnosis in 5185 patients with cancer across 13 disease sites over the course of at least 12 years.<sup>44</sup> In this study, recent quitters were examined as a specific subset of former smokers who quit within 1 year

of the study's structured smoking assessment, allowing for comparisons to individuals who actively smoked during cancer treatment. For disease sites with larger recent quit cohorts (ie, lung, head /neck cancers), recent quitters had lower overall mortality risk compared with continued smokers (current smoker vs. recent quitter: lung cancer HR= 1.38–1.42; head/neck cancer HR= 2.11–2.15).<sup>44</sup> Similarly, a systematic review and meta-analysis of 10 observational studies pointed to a 5-year survival benefit for patients with lung cancer who quit smoking compared to patients who continued smoking (70% vs. 33%).<sup>42</sup> In a comprehensive cancer center study that controlled for disease characteristics, smoking history, and patient demographics, 250 patients with lung cancer who quit smoking had statistically improved survival time of 9 months over those who continued to smoke through treatment and beyond.<sup>108</sup>

Smoking cessation is linked to reduced risk of recurrence and second primary tumor formation. Data from patients with lung and head and neck cancers showed that rates of second primary cancers were lower for patients who quit smoking than for those who continued to smoke after diagnosis.<sup>106,109,110</sup>

Cessation at or near cancer diagnosis appears to reduce treatment-related complications compared to patients who continued smoking. In patients undergoing lung cancer resection, preoperative cessation mitigated the risk of pulmonary complications and in-hospital mortality. Risk-adjusted odds ratios (ORs) for mortality and pulmonary complications decreased as preoperative cessation time increased from 14 days to 1 month, to 1 to 12 months, and to more than 12 months.<sup>59</sup> A retrospective study of 188 patients undergoing reconstructive surgery after treatment for head and neck cancer revealed that preoperative smoking cessation of at least 3 weeks led to lower incidence of wound healing complications than patients who continued smoking.<sup>66</sup>

Cessation has been shown to lead to improvements in various measures of general health and well-being for patients with cancer. Smoking cessation improved performance status at 6 and 12 months post-lung cancer diagnosis over that of continued smokers when adjusting for disease stage, patient demographics, therapy, and comorbidity.<sup>111</sup> Additionally, patients with cancer who quit smoking benefited from lower rates of smoking-related cardiovascular and pulmonary disease.<sup>112</sup>

### Evaluation and Assessment of Patient Smoking

These guidelines highlight the importance of evaluating and assessing smoking status and history in patients with cancer. In a recent policy statement, the AACR emphasized the need for universal assessment and documentation of tobacco use by patients with cancer both in the standard clinical setting and in oncology clinical trials.<sup>10</sup> However, current practice is suboptimal, as inadequate or inconsistent assessment and documentation of smoking status has been reported both in the care setting and in the context of clinical trials.<sup>12,18</sup>

Despite the demonstrated adverse effects of smoking during cancer treatment, a large proportion of cancer clinical trials do not collect adequate, up-to-date information regarding patient smoking status and history, particularly for malignancies other than well-known tobacco-related cancers (eg, lung, head and neck cancers).<sup>17,18</sup> Such assessments are needed to make evidence-based determinations of the impact of smoking on patients, treatment efficacy, and side effects.

In a large study conducted at a comprehensive cancer center, a smoking assessment questionnaire was integrated into the electronic health record (EHR) in order to automatically identify and refer appropriate candidates for onsite cessation services. The smoking assessment items incorporated into the EHR were refined based on

analysis of responses from an initial patient screen containing 23 items. Response analysis revealed that the most effective questions for generating referrals included whether: 1) patients smoked cigarettes every day, some days, or not at all; and 2) if/what other types of tobacco products were used. For former smokers, it was important to assess the last time a patient smoked a cigarette, “even a puff,” and for established enrollees to the cessation program, what type(s) of cessation aids were being employed.<sup>113</sup> The study revealed that just 3 assessment questions made it possible to efficiently and accurately identify the vast majority (over 98%) of current smokers or those at risk for smoking relapse.

### Determining Smoking Status

The NCCN Guidelines for Smoking Cessation advocate for smoking status to be updated in the patient’s health record at regular intervals to indicate any status changes or quit attempts. To do so, the panel recommends that providers initially ascertain: 1) whether the patient has ever smoked, and if so, initially and at regular intervals; 2) whether the patient is a current smoker; and 3) whether the patient has smoked within the past 30 days. All information should be recorded in the medical record. As a follow-up to the initial evaluation, these guidelines direct providers to a tailored patient assessment based on smoking status and history. Specific algorithms for current smokers (patient smoked within the last 30 days) and recent quitters/former smokers (greater than 30 days since patient last smoked) are included. For never smokers or longer-term former smokers, providers should urge patients to remain smoke-free, providing them with the benefits for remaining smoke-free.

### Assessing Smokers

In patients who are current smokers, assessment begins by determining readiness to quit within the next 30 days. For patients who are ready to quit, providers should assess nicotine dependency to understand the

chances for success and risk of relapse, and document the findings in the patient’s health record. To assess nicotine dependency, providers should query patients regarding the amount of cigarettes smoked per day, how soon the patient smokes after waking up in the morning, whether the patient uses other forms of tobacco (eg, pipes, cigars, snuff, e-cigarettes), and if so, what quantity. In order to best tailor the personalized quit plan, providers should also gather information regarding the patient’s history of quit attempts and why they were or weren’t successful. Specifically, providers should ascertain the longest period of abstinence achieved, the date of the most recent quit attempt, what cessation aids were employed, and why these failed. It is important to document the patient’s previous experience with smoking cessation aids, including any medications, behavior therapy, e-cigarettes, quitlines, websites, smart phone applications, or other media aids. The patient’s perspective on why these aids were unsuccessful—such as medication side effects, continued cravings, or inefficacy—are important pieces of information.

When possible, providers should work with patients to set a near future quit date and/or consider smoking reduction with the goal of cessation in the near future. A meta-analysis of 10 randomized trials in 3760 patients with cancer found quit rates to be comparable when comparing abrupt cessation to gradual smoking reduction.<sup>114,115</sup> Therefore, both options can be used after discussions with the patient. Trial data since then have continued to mirror this trend of comparable success rates with abrupt cessation and gradual reduction.<sup>116</sup> Pharmacotherapy with NRT or varenicline can be initiated in active smokers to promote smoking reduction as an intermediate step to eventual cessation. At each visit, providers should reassess readiness to quit and engage in motivational dialog as indicated.



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## Smoking Cessation

### *Motivational Interviewing*

For patients who do not express a readiness to quit, providers are encouraged to engage patients in a motivation dialog about smoking and to ensure patients are aware of the disease-specific risks of smoking and benefits of quitting. Educational resources should be provided. Providers should assess and address patient-reported barriers and concerns regarding cessation. The panel recommends that clinicians provide patients with reasons, ideas, and needs for smoking cessation, emphasizing the importance of both encouragement and directness with patients who smoke. When incorporating MI to promote willingness to quit, the panel emphasized the importance of the following general principles: 1) express empathy; 2) develop discrepancy; 3) roll with resistance; and 4) support self-efficacy.<sup>117,118</sup> For a summary of the methods and data on motivational interviewing (MI) for smoking cessation, see *Principles of Behavior Therapy*.

### *Assessing Former Smokers/Recent Quitters*

To evaluate patients who recently quit at least 30 days prior, providers should assess the patient's risk of relapse. The panel suggests the following characteristics to identify patients at high risk for relapse: frequent/intense cravings; elevated work stress or depression; cohabitating or working with smokers; quitting within the past year; use of ongoing smoking cessation treatment; or substance use or abuse. The panel considers patients demonstrating at least one of these characteristics to be at higher risk for relapse and recommends a management plan tailored to prevent relapse. Providers should discuss risk of relapse with patients and provide guidance and support to promote continued smoking cessation attempts.

Management of patients who demonstrate an elevated risk of relapse includes discussion of pharmacotherapy (eg, short-acting NRT) to promote maintenance of abstinence. Providers should review smoking-

associated risks for patients with cancer as well as the health benefits of abstinence. If needed, brief counseling on relapse risk factors and relapse prevention can be offered, with referrals for behavior therapy as indicated. All management plans and counseling should be documented in the patient health record. For patients deemed to be at low risk for relapse, providers should reinforce success and highlight the importance of continued abstinence. Risk of relapse should be reevaluated at subsequent visits.

It is important to regularly reevaluate patients' smoking status and risk of relapse, which can be accomplished in person or by phone. If relapse occurs, referral for smoking cessation pharmacotherapy and counseling are recommended. Additionally, providers should remain aware that patient self-report of smoking status may underestimate the rate of active smoking among patients with cancer, as is evidenced by research comparing self-reported and objective measures.<sup>119-121</sup>

## Smoking Cessation During Cancer Treatment

### Devising a Treatment Plan

Following assessments, providers should establish a personalized quit plan for each patient that takes into account the patient's nicotine dependency, prior quit attempts and any cessation aids used, and smoking cessation therapy options. Providers should work with patients to set a quit date, preferably within 2 weeks and at least 2 or more weeks prior to any planned surgical procedures. Preoperative smoking cessation interventions that combine pharmacotherapy with behavioral therapy may be most effective.<sup>122-125</sup> Risk of relapse and smoking slips should be discussed with the patient along with reassurance and support for continued cessation efforts should slips occurs.

### First-line Therapy Recommendations

Based on clinical trial data of smoking cessation in patients with cancer, the panel recommends a combination frontline approach including pharmacotherapy and behavior therapy for smoking cessation for patients with cancer. Population studies and meta-analyses of randomized or quasi-randomized trial data support the addition of behavior therapy to pharmacotherapy to enhance the rate of success.<sup>126-128</sup> First-line therapy recommendations in these guidelines include combination NRT (combined long- and short-acting NRT) or varenicline (typical initial duration of 12 weeks), and include behavioral counseling as indicated below. The combination NRT approach includes the use of nicotine patch plus a short-acting NRT (eg, for break-through craving) such as nicotine gum, lozenge, inhaler, or nasal spray. Providers should note that nausea is a common side effect of varenicline and may need to be managed in patients with cancer who are receiving chemotherapy. For discussion of the evidence for individual pharmacotherapeutic regimens, see *Principles of Pharmacotherapy*.

### Follow-up and Surveillance

Initial follow-up within 2 weeks of initiating smoking cessation therapy is important in order to assess efficacy and toxicity of pharmacotherapy. When possible, follow-up can occur during planned clinical visits or individual/group therapy sessions. Alternatives include phone contact. During follow-up, providers should assess risk of relapse and, as indicated, consider adjusting the dose and or type of pharmacotherapeutic. Patients may slip or relapse, which is expected and can be managed. Brief slips may or may not require medication adjustments. Maintain close follow-up through the duration of therapy. At 12 weeks, assessment of smoking status should be made in person or by phone. For pharmacotherapy courses exceeding 12 weeks

duration, assessment should be repeated at the end of the course of therapy. If patients remain smoke free, additional follow-up should take place at 6 and 12 months, either in person or by phone. Patients who experience smoking relapse can be considered for second-line therapy.

### Treatment for Relapse

For patients who experience relapse, the panel recommends continued behavior therapy with either alternative first-, second-, or third-line pharmacotherapy as indicated. Second-line pharmacotherapy includes varenicline with combination NRT or bupropion with combination NRT. For further relapse, extended duration of pharmacotherapy can be considered. Additional or more intensive behavior therapy is also an option. Third-line pharmacotherapy options include a combination regimen with varenicline, bupropion, and combination NRT, or nortriptyline or clonidine. Smoking status should be re-evaluated at the end of each course of prescribed pharmacotherapy, with additional follow-up at 6 and 12 months after successful quitting.

## Principles of Pharmacotherapy

### General

Recommended initial duration of treatment is 12 weeks for varenicline and combination NRT, and 7 to 12 weeks for bupropion. Research suggests that longer courses of certain cessation agents may be associated with higher rates of 7-day point-prevalence abstinence.<sup>129</sup> Duration of therapy can be extended to promote continued cessation, but providers should attempt to avoid unnecessarily long treatment duration when possible. Dose adjustments should be considered as clinically indicated.

In most circumstances the side effects related to all first-line smoking cessation medications are minimal and are considered an acceptable



risk compared to smoking. A review of post-marketing case reports on adverse neuropsychiatric effects from smoking cessation medications have generated some safety concerns in the past,<sup>130</sup> but recent large-scale analyses of the data support the safety of these agents.<sup>131,132</sup> Although serious side effects of first-line cessation agents are extremely rare, providers should refer to manufacturer inserts for exhaustive lists of potential side effects and warnings.

Adherence to pharmacotherapy is important to promote optimal outcomes and success, and numerous studies have tested interventions designed to promote and improve medication adherence.<sup>133</sup>

Below, data from various clinical trials are discussed. Included in this discussion are findings from a 2013 Cochrane network meta-analysis that included data on pharmacologic interventions across 267 individual studies in 101,804 participants.<sup>134</sup> The authors characterized positive treatment outcome as continuous or prolonged abstinence at least 6 months from the start of smoking cessation therapy. Harm outcomes were measured by the incidence of serious adverse events associated with treatment.

### Front-line Therapy Recommendations

For patients with cancer, the guidelines recommend front-line therapy with either combination NRT or varenicline. The efficacy and safety data on these agents are summarized below.

#### **Varenicline**

##### *Efficacy*

Varenicline is a non-nicotinic partial agonist of the alpha4beta2 subtype of the nicotinic acetylcholine receptor. Varenicline partially mimics the

effects of nicotine in the brain's reward center and competitively inhibits the binding of nicotine from cigarettes.<sup>135</sup> Systematic reviews/meta-analyses have identified varenicline as the most effective single pharmacotherapy option for smoking cessation.<sup>134,136,137</sup> Cochrane network meta-analysis data report that varenicline increases the odds of smoking cessation by almost three-fold compared with placebo (OR, 2.88; 95% CI, 2.40–3.47).<sup>134</sup> Direct comparison of the cumulative data suggest that varenicline was more efficacious than bupropion (OR, 1.59; 95% CI, 1.29–1.96) and single forms of NRT such as nicotine patch, nicotine gum, and other formulations (OR, 1.57; 95% CI, 1.29–1.91).<sup>134</sup> Varenicline appeared to be equally as likely to promote smoking cessation as combined treatment with more than one form of NRT (OR, 1.06; 95% CI, 0.75–1.48), so that both may be offered depending on patient circumstances.<sup>134</sup>

A recent clinical trial (n = 1510) revealed that a 24-week course of varenicline effectively promoted smoking cessation in patients who were unwilling to quit but willing to gradually reduce cigarette consumption.<sup>138</sup> Therefore, this agent provides an alternative for patients who cannot or will not attempt abrupt cessation. A clinical trial of 1236 smokers showed that an additional 12 weeks of varenicline maintenance therapy helped to sustain continued abstinence in those who successfully quit during initial treatment.<sup>139</sup>

A recent study examined whether varenicline dose increases would boost treatment efficacy in patients who had a low or no response to standard dosing. A double-blind randomized controlled trial (RCT) of 503 smokers found no evidence to suggest that gradual dose titration beyond the standard 2 mg dose (up to a maximum 5 mg/day) lessened frequency of urges and nicotine withdrawal symptoms, or increased cessation rates.<sup>140</sup> However, nausea and vomiting were increased in the treatment group receiving more than 2 mg/day. Additionally, another



study showed that varenicline was effective and well-tolerated for retreating patients who had previously received this agent (RCT, n = 498).<sup>141</sup>

### *Safety*

Varenicline safety has been extensively examined to determine the risk of adverse effects, particularly serious cardiovascular events and neuropsychiatric changes. Initial phase III studies found varenicline to be safe and generally well-tolerated compared with bupropion or placebo; common side effects included nausea, insomnia, and abnormal dreams with rates of approximately 28% to 29%, 14%, and 10% to 13%, respectively.<sup>142,143</sup>

Importantly, recent systematic reviews and meta-analyses of RCT data have not identified a significant link between varenicline and increased risk of serious cardiovascular adverse events.<sup>134,144-146</sup> However, the cardiovascular safety of varenicline has remained a topic of interest and concern.<sup>147-149</sup> Based on the current evidence base for cardiovascular risks, the panel considers varenicline to be safe and to have a favorable risk/benefit ratio for use in patients with cancer who smoke.

Concerns regarding neuropsychiatric adverse effects of varenicline have also been extensively investigated in smokers with comorbid mental illness.<sup>134</sup> Despite reviews of case reports that raised concern,<sup>130</sup> a 2015 systematic review and meta-analysis of 39 randomized controlled smoking cessation trials identified no evidence to suggest that varenicline increases risk of suicide or suicide attempts, suicidal ideation, depression, or death.<sup>132</sup> Another trial showed that varenicline increased smoking cessation rates without exacerbating anxiety and depression symptoms in adults with stably treated current or past depression.<sup>150</sup>

Varenicline maintenance therapy was examined in a trial that enrolled patients with bipolar disorder or schizophrenia who were abstinent following an initial treatment course with 12 weeks of varenicline and cognitive behavioral (CBT) therapy.<sup>151</sup> At 6 months and 1 year following maintenance treatment, the cohort receiving maintenance therapy with varenicline therapy along with CBT had higher cessation rates than the cohort receiving maintenance CBT alone.

### ***Combination Nicotine Replacement Therapy***

#### *Efficacy*

NRT offers an alternative nicotine delivery method and can be used to ameliorate nicotine withdrawal symptoms during cessation attempts. Combination NRT incorporating long-term and fast-acting NRT offer the greatest potential benefits for smokers.<sup>122,134,152,153</sup> Cochrane network meta-analysis data published in 2013 support the superiority of combination NRT over single forms of NRT such as nicotine patch (OR, 1.43; 95% CI, 1.08–1.91), nicotine gum (OR, 1.63; 95% CI, 1.21–2.2), and various other forms that collectively include inhaler, lozenge, spray, or tablets (OR, 1.34; 95% CI, 1.0–1.8).<sup>134</sup> All forms of NRT were superior to placebo, but smokers using combination NRT were almost three times as likely to succeed (OR, 2.73; 95% CI, 2.07–3.65).<sup>134</sup> Compared with single forms of NRT, combination NRT using a patch plus short-acting NRT improved the odds of quitting (OR, 1.34; 95% CI, 1.18–1.51).<sup>134,154</sup>

The success of NRT may be contingent on concurrent behavior therapy to support cessation. In a large population study, over-the-counter NRT resulted in similar rates of cessation to those who used no aid. The addition of behavior therapy to NRT increased the odds of success nearly three-fold.<sup>126</sup>



### Safety

The safety of combination NRT for use in humans has been demonstrated and benefits are considered to outweigh potential risks.<sup>155</sup> Importantly, providers should be aware that blood nicotine levels from NRT, including combination NRT, are significantly less than that from smoking cigarettes. Therefore, providers and smokers should not be dissuaded from using NRT to foster quitting and long-term cessation. Recent reviews of the data suggest that NRT is not linked to increased serious cardiovascular adverse events when used for smoking cessation.<sup>146</sup> While myocardial infarction has rarely been reported in NRT users, there is insufficient evidence that NRT increases the risk of myocardial infarction or cardiovascular disease.<sup>154</sup>

In the past, the safety of NRT has been evaluated in light of the bioactivity of nicotine and evidence that this drug can promote cell growth in certain types of cancer cells.<sup>156</sup> Some *in vitro* data suggested that nicotine increased the malignant potential of small cell lung cancer cells<sup>157</sup>; induced chemoresistance in models using lung cancer cells<sup>89-91</sup> and nasal epithelial cells<sup>92</sup>; and promoted chemoresistance and metastasis in pancreatic cancer cell and mouse models.<sup>93</sup>

However, other studies suggested no effects of physiological levels of nicotine exposure on tumorigenesis in mouse lung cancer models.<sup>158,159</sup> Moreover, there is insufficient evidence that NRT causes cancer in humans.<sup>158-162</sup> Evaluation of data from 3320 participants in the Lung Health Study, which recorded in-study NRT use and smoking exposure, found that NRT was not a significant predictor of lung cancer, while smoking was.<sup>161</sup>

### Second- and Third-line Therapy Recommendations

Second-line therapy recommendations include varenicline with combination NRT or bupropion with combination NRT. In the third-line

setting, recommendations include combination therapy with varenicline, bupropion, and NRT. Single-agent third-line options include nortriptyline and clonidine. The data for various regimens are summarized below.

### Bupropion

#### Efficacy

Bupropion was first approved to treat depression but its efficacy as a cessation aid also became apparent. In addition to its effects on the dopaminergic and adrenergic systems, this agent also acts as an inhibitor of nicotinic acetylcholine receptors. A 2014 Cochrane review of 44 trials examined bupropion efficacy, revealing an RR of 1.62 (95% CI, 1.49–1.76).<sup>163</sup> Some evidence suggests that bupropion may be particularly beneficial as a smoking cessation agent for persons with depression.<sup>164,165</sup> Additionally, longer duration of bupropion treatment may help to prevent relapse in those who have successfully quit.<sup>166</sup>

#### Safety

Bupropion reduces the seizure threshold and meta-analyses of trial data have found a 0.1% seizure risk among those receiving the drug for smoking cessation.<sup>163</sup> Neuropsychiatric effects have also been identified as a safety concern with bupropion, although to a lesser extent than varenicline.<sup>130</sup> However, recent systematic reviews of the data have found that serious neuropsychiatric adverse events were rarely associated with bupropion prescribed for smoking cessation, including studies of bupropion in patients with mental illness.<sup>134,167</sup> Similarly, regarding risk of serious adverse cardiovascular effects, recent meta-analyses do not show elevated risk as a result of bupropion use for smoking cessation.<sup>134,146,163</sup>



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## Smoking Cessation

### **Varenicline + NRT**

A study in 435 smokers found that the addition of nicotine patch to varenicline therapy significantly increased the cessation rates at the end of treatment (12 weeks), at 24 weeks, and at 6-months follow-up.<sup>168</sup> No significant differences were noted for side effect incidence between varenicline/NRT and varenicline/placebo with the exception of skin reactions, which were increased with combination therapy (14.4% vs. 7.8%;  $P = .03$ ). However, one RCT of 341 smokers did not find enhanced cessation rates at 12 and 24 weeks follow-up among individuals receiving a combination of varenicline and nicotine patch, versus varenicline alone.<sup>169</sup> The addition of nicotine patch to varenicline did not cause significant changes in side effect profiles. Similarly, a trial in 117 participants did not find evidence that the addition of NRT to varenicline increased abstinence rates at 1, 4, or 12 weeks after the targeted quit date, and no between-group differences in adverse effects were found.<sup>170</sup>

### **Bupropion + NRT**

A large trial in the United Kingdom (n = 1071) examined the efficacy of NRT alone, bupropion alone, and NRT plus bupropion.<sup>164</sup> All participants received 7 weeks of behavior therapy support in addition to the pharmacologic interventions. Abstinence rates at 6-month follow-up ranged from 24.2% to 27.9% and did not differ significantly between cohorts. Several unwanted side effects were more common with bupropion than NRT (eg, disturbed sleep, dry mouth, headaches, nausea), and side effects of combination therapy were not significantly different versus bupropion alone. Five serious adverse events occurred in the bupropion group, including allergic reaction (n = 3), neuropsychiatric symptoms (n = 1), and chest pain (n = 1). A trend towards improved efficacy of bupropion in patients with a history of depression was noted ( $\chi^2 = 2.86$ ,  $P = .091$ ). A smaller RCT studying the addition of bupropion to combination NRT and behavior therapy in

patients with schizophrenia suggested that combination pharmacotherapy promoted smoking reduction and cessation, but also demonstrated a high relapse rate after discontinuation of treatment.<sup>171</sup> A 2014 meta-analysis of 12 trials examining this combination revealed a nonsignificant trend in improved cessation with the addition of NRT to bupropion.<sup>163</sup>

### **Varenicline + Bupropion**

In an RCT of smokers who demonstrated an inadequate response to first-line nicotine patch treatment (n = 222), combination therapy with varenicline and bupropion appeared to be more efficacious than varenicline alone as a second-line therapy option.<sup>172</sup> This observation was more pronounced among male smokers and those with a high level of nicotine dependency. Although no significant differences in side effects were observed between varenicline and bupropion versus varenicline alone, dose reductions were required for 11.5% and 24.8% of patients, respectively. Common side effects were vivid dreams, change in taste perception, thirst, insomnia, and irritability. Another study of varenicline plus bupropion therapy versus varenicline alone (n = 316) demonstrated that combination therapy increased prolonged abstinence but did not affect 7-day point prevalence at 12 and 26 weeks follow-up, and no significant differences were observed between the groups at 52 weeks.<sup>173</sup> In this study, anxiety (7.2% vs. 3.1%;  $P = .04$ ) and depressive symptoms (3.6% vs. 0.8%;  $P = .03$ ) occurred more frequently in patients receiving combination therapy versus varenicline alone.

### **Nortriptyline and Clonidine**

Studies have also suggested some efficacy of off-label use of nortriptyline, a tricyclic antidepressant, as well as the antihypertensive agent, clonidine.

A Cochrane network review identified 6 trials comparing nortriptyline with placebo, finding a pooled RR of 2.03 (95% CI, 1.48–2.78).<sup>102,133</sup> However, as an adjunct to NRT, clear-cut benefits were not observed.<sup>134,174</sup>

Clonidine is recommended as a third-line smoking cessation option. Although several studies have produced data in favor of clonidine as a cessation aid versus placebo, not all study data reveal a statistically significant effect. Additionally, clonidine's benefits can be counteracted by dose-dependent increases in adverse side effects. A 2013 Cochrane network review of data from 6 studies resulted in a pooled RR of 1.63 (95% CI, 1.22–2.18).<sup>102</sup>

## Principles of Behavior Therapy

### General Principles

The guidelines provide the following guiding principles on behavior therapy, which have been developed in consideration of the existing evidence base, clinical practice guidelines, and expert consensus.<sup>127,128,155</sup>

The panel recommends a combination of behavior therapy with pharmacotherapy for best outcomes. A 2012 systematic review of 41 studies provided support for the efficacy of this approach.<sup>127</sup> The “real world effectiveness” of adding a behavior therapy component to smoking cessation therapy was further supported by a large population study published in 2014.<sup>126</sup> Behavior therapy may enhance motivation and support optimal medication strategies and adherence to pharmacotherapy.

As a general principle, more intensive behavior therapy is preferred over brief advice or counseling.<sup>128</sup> The evidence supports a measurable dose-response effect of behavior therapy with more numerous and/or

longer sessions delivering improved outcomes. The panel defines intensive behavioral therapy as at least 4 sessions (in person; group or individual) lasting at least 10 minutes but typically 30 minutes or longer.

The most successful behavior therapy strategies employ practical counseling, which addresses problem solving and skills training, as well as social support and MI (see section on Motivational Interviewing below) as elements of the treatment plan.<sup>118,155</sup> Optimally, behavior therapy plans should take into account a patient's nicotine dependence levels, previous quit attempts, and cessation aids utilized. In doing so, patients can be equipped with tailored strategies to cope with nicotine withdrawal symptoms, environmental smoking triggers, and stressful situations. For instance, the addition of a CBT program designed to improve stress management improved cessation rates over controls receiving standard smoking cessation therapy.<sup>175</sup>

Providers should prepare patients for nicotine withdrawal symptoms, which typically peak at several days to 2 weeks post-cessation before gradually subsiding.<sup>176-178</sup>

A number of modalities can be employed to deliver behavior therapy to patients. Counseling can take place in a variety of settings such as in-person, remotely by telephone,<sup>179</sup> or through web-based interventions.<sup>180</sup> Effective in-person counseling can occur as an individual session or in the group therapy setting.<sup>181,182</sup> Additionally, print materials<sup>183</sup> and mobile telephone “apps”<sup>184-186</sup> can be used to deliver behavior therapy. However, providers should be aware that media-based behavioral interventions, particularly those using mobile telephones, may vary in the degree to which they comply with clinical practice guidelines.<sup>187</sup> Selection of a particular modality or modalities should be guided by patient preference, medical history, and resource availability.

If specialized resources are limited, effective behavior counseling can still be provided. For instance, brief counseling by providers has been shown to generate a small but important increase in quit rates.<sup>188-190</sup> Additionally, quitlines can provide essential behavioral support in the absence of in-person counseling resources.<sup>179</sup> For instance, the addition of combination NRT to quitline counseling improved cessation outcomes.<sup>191</sup>

### Tailoring Behavior Therapy for Patients with Cancer

The prevalence of mental disorders or serious emotional issues in patients with cancer is high, with several large studies reporting rates between 30% and 40%.<sup>192-194</sup> The high rates of anxiety, depression, and stress can present a significant challenge for patients with cancer who attempt to quit smoking in the face of these common smoking/relapse triggers. Patients with cancer, particularly those experiencing psychiatric comorbidity, may benefit significantly from behavior therapy programs tailored to manage cancer-related issues that predispose patients to relapse. Referral to specialized smoking cessation programs may be necessary so that these patients have access to staff trained to treat co-morbid substance dependence and mental health disorders.

As resources allow, specialized treatment centers should provide tailored smoking cessation therapy programs that address the unique needs of patients with cancer. Beneficial services might include individual and group therapy focusing on the challenges specific to cancer treatment and survival, which would ideally be provided by clinicians experienced in working with patients with cancer.

### Motivational Interviewing for Patients Unwilling to Quit

MI can be employed for smokers who are currently unwilling to quit. MI is typically performed by a provider trained in strategic questioning and listening, and uses a series of directive and nondirective

approaches.<sup>117,118</sup> Through this approach to counseling, clinicians can explore the smoker's feelings, beliefs, and values in order to identify opportunities for change and facilitate willingness to quit.

A 2015 Cochrane Database review of 28 studies examined the efficacy of MI for smoking cessation, revealing a modest but significant increase in chance of quitting with MI versus brief advice or usual care.<sup>195</sup> MI by a primary care physician appeared to be somewhat more successful than that administered by counselors, although both were effective. Notably, one-time short MI sessions of less than 20 minutes had demonstrated efficacy.<sup>195</sup>

In order to promote willingness to quit smoking, the U.S. Preventive Services Task Force (USPSTF) recommends a model of MI that employs the “5 R's” of relevance, risks, rewards, roadblocks, and repetition.<sup>155</sup> This model encourages that motivational information be *relevant* to the individual patient, and that clinicians and patients work together to identify personalized *risks* of smoking and potential *rewards* of cessation. By having the patient identify perceived *roadblocks* to quitting, providers can suggest tailored treatments to address patient-reported concerns. Finally, this model recommends *repetition* of MI at each patient visit, coupled with reminders that repeated quit attempts may be necessary to achieve long-term cessation.

### Alternative Treatment Approaches

The panel has reviewed the available evidence for alternative smoking cessation treatment approaches. Particular attention has been paid to the discussion of e-cigarettes for smoking cessation given increasing popularity and widespread use. Limited data are available on the safety and efficacy of these approaches, specifically for patients with cancer; data have been drawn primarily from studies in the general population.



The panel has found insufficient evidence to support the use of alternative therapies alone or in combination with standard smoking cessation methods, and use of alternative therapies is not recommended. The guidelines recommend that patients use evidence-based cessation methods to avoid any delay in achieving smoking abstinence. Smoking slips and relapse are common, and prior unsuccessful quit attempts with conventional therapies do not justify the use of unproven alternative cessation methods. When discussing alternative therapies, providers should counsel patients on potential interactions with evidence-based cessation methods and/or cancer treatments.

### Electronic Cigarettes

The popularity of e-cigarettes, and their various derivatives, is a recent phenomenon, and, as such, the available literature is new and relatively limited, particularly within specific subpopulations such as patients with cancer. At the present time, e-cigarettes are not recommended, and instead known effective methods for smoking cessation should be offered. Below, we discuss the current data and expert opinions on e-cigarettes for smoking cessation.

In a recent draft recommendation statement, the USPSTF “concludes that the current evidence is insufficient to recommend electronic nicotine delivery systems (ENDS) for tobacco cessation. The USPSTF recommends that clinicians direct patients who smoke tobacco to other cessation interventions with established effectiveness and safety.”<sup>196</sup> Several health care organizations have released similar policy statements concerning ENDS, highlighting the urgent need for research on the safety of these devices and efficacy as a cessation aid. The American Heart Association, American Association for Cancer Research, and American Society of Clinical Oncology recognize the

potential for ENDS to alter existing smoking behaviors, as well as the lack of definitive data regarding associated benefits and harms.<sup>197,198</sup> Experts in the field generally acknowledge that ENDS may offer an attractive approach for smoking reduction and/or cessation in certain populations. However, these policy statements also highlight the unknown potential for ENDS to affect nicotine addiction, combustible tobacco product use, and renormalization of smoking behaviors.

One study examined e-cigarette use in 1074 patients with cancer who enrolled in a tobacco treatment program at a comprehensive cancer center.<sup>199</sup> The study revealed a marked increase in e-cigarette use from 10.6% to 38.5% between 2012 and 2013. E-cigarette users, most often patients with thoracic or head and neck cancers, were more nicotine dependent and had greater numbers of prior quit attempts. At follow-up (6–12 months after intake), e-cigarette users were no more likely to have quit than non-users (OR, 1.0; 95% CI, 0.5–1.7), calling into question the potential benefits of e-cigarettes as a cessation agent for patients with cancer.

Systematic reviews have analyzed the data from the general population to determine the potential efficacy of e-cigarettes as a smoking cessation aid, noting a limited overall pool of data and heterogeneous measures. A 2014 Cochrane Database review of RCT data from 662 participants showed that participants using nicotine e-cigarettes were significantly more likely to achieve cessation for at least 6 months compared to those using placebo e-cigarettes (RR, 2.29; 95% CI, 1.05–4.96).<sup>200</sup> A greater number of participants were able to reduce cigarette consumption with nicotine e-cigarettes vs. placebo (RR, 1.31; 95% CI, 1.02–1.68), and one study suggested that nicotine e-cigarettes improved reduction over nicotine patch (RR, 1.41; 95% CI, 1.20–1.670).<sup>200</sup> Across all 13 studies examined, no serious adverse events were reported.<sup>200</sup> Similarly, a 2015 systematic review and meta-analysis



of data from 1242 participants suggested that nicotine-containing e-cigarettes were more effective cessation aids than non-nicotine-containing e-cigarettes (RR, 2.29; 95% CI, 1.05–4.97).<sup>201</sup> E-cigarette use over a minimum of 6 months was associated with an 18% reported smoking cessation rate (effect size, 0.20; 95% CI, 0.11–0.28), and e-cigarette use was also associated with smoking reduction.<sup>201</sup>

A large cross-sectional study of 5863 adults in the United Kingdom assessed the “real-world effectiveness” of e-cigarettes for smoking cessation compared to NRT and unaided quitting, revealing that e-cigarette users were more likely to report abstinence compared with the other cohorts (e-cigarettes vs. NRT: OR, 2.23; 95% CI, 1.70–2.93; e-cigarette vs. no aid OR, 1.38; 95% CI, 1.08–1.76).<sup>202</sup> These observations persisted when adjusting for measures of nicotine dependence across the cohorts.

### Other Alternative Approaches

Very limited data exist to support exercise-based interventions; small study size, inadequate controls, and insufficient exercise intensity limit the ability to make conclusions based on the existing evidence.<sup>203</sup>

Sufficient efficacy data are also lacking to support the use of alternative therapies such as acupuncture, hypnosis, and nutritional supplements.

A 2014 systematic review of the data on acupuncture, acupressure, and laser therapy revealed no consistent, bias-free evidence to support these methods for smoking cessation, although pooled evidence was suggestive of possible short-term benefits.<sup>204</sup> Acupuncture was less effective than NRT and there was no evidence to support electrostimulation for smoking cessation. Similarly, systematic reviews of the data on hypnosis for smoking cessation revealed inadequate high-quality evidence to support this approach.<sup>205,206</sup> Claims of efficacy data for hypnosis from several studies were not substantiated by the

review of RCT data. Controlled studies are needed to provide higher quality evidence on these interventions both in the general population and among patients with cancer.

### References

- Jamal A, Agaku IT, O'Connor E, et al. Current Cigarette Smoking Among Adults — United States, 2005–2013. Morbidity and Mortality Weekly Report (MMWR). Vol. 63: Centers for Disease Control and Prevention; 2014:1108-1112. Available at: <http://www.cdc.gov/mmwr/preview/mmwrhtml/mm6347a4.htm>.
- The Health Consequences of Smoking—50 Years of Progress: A Report of the Surgeon General. Atlanta, GA: U.S. Department of Health and Human Services 2014. Available at: <http://www.surgeongeneral.gov/library/reports/50-years-of-progress/>.
- Siegel RL, Miller KD, Jemal A. Cancer statistics, 2015. CA Cancer J Clin 2015;65:5-29. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25559415>.
- Tseng TS, Lin HY, Moody-Thomas S, et al. Who tended to continue smoking after cancer diagnosis: the national health and nutrition examination survey 1999-2008. BMC Public Health 2012;12:784. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22974404>.
- Park ER, Japuntich SJ, Rigotti NA, et al. A snapshot of smokers after lung and colorectal cancer diagnosis. Cancer 2012;118:3153-3164. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22271645>.
- Underwood JM, Townsend JS, Tai E, et al. Persistent cigarette smoking and other tobacco use after a tobacco-related cancer diagnosis. J Cancer Surviv 2012;6:333-344. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22706885>.
- Westmaas JL, Newton CC, Stevens VL, et al. Does a Recent Cancer Diagnosis Predict Smoking Cessation? An Analysis From a Large Prospective US Cohort. J Clin Oncol 2015. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25897151>.
- Westmaas JL, Alcaraz KI, Berg CJ, Stein KD. Prevalence and correlates of smoking and cessation-related behavior among survivors of ten cancers: findings from a nationwide survey nine years after diagnosis. Cancer Epidemiol Biomarkers Prev 2014;23:1783-1792. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25100826>.
- Mayer DK, Carlson J. Smoking patterns in cancer survivors. Nicotine Tob Res 2011;13:34-40. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/21097514>.
- Toll BA, Brandon TH, Gritz ER, et al. Assessing tobacco use by cancer patients and facilitating cessation: an American Association for Cancer Research policy statement. Clin Cancer Res 2013;19:1941-1948. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23570694>.
- Hanna N, Mulshine J, Wollins DS, et al. Tobacco cessation and control a decade later: American society of clinical oncology policy statement update. J Clin Oncol 2013;31:3147-3157. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23897958>.
- Goldstein AO, Ripley-Moffitt CE, Pathman DE, Patsakham KM. Tobacco use treatment at the U.S. National Cancer Institute's designated Cancer Centers. Nicotine Tob Res 2013;15:52-58. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22499079>.
- Weaver KE, Danhauer SC, Tooze JA, et al. Smoking cessation counseling beliefs and behaviors of outpatient oncology providers. Oncologist 2012;17:455-462. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22334454>.
- Warren GW, Marshall JR, Cummings KM, et al. Addressing tobacco use in patients with cancer: a survey of American Society of Clinical Oncology members. J Oncol Pract 2013;9:258-262. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23943904>.
- Warren GW, Marshall JR, Cummings KM, et al. Practice patterns and perceptions of thoracic oncology providers on tobacco use and cessation in cancer patients. J Thorac Oncol 2013;8:543-548. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23529191>.



16. Warren GW, Dibaj S, Hutson A, et al. Identifying Targeted Strategies to Improve Smoking Cessation Support for Cancer Patients. J Thorac Oncol 2015. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/26317914>.

17. Gritz ER, Dresler C, Sarna L. Smoking, the missing drug interaction in clinical trials: ignoring the obvious. Cancer Epidemiol Biomarkers Prev 2005;14:2287-2293. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/16214906>.

18. Peters EN, Torres E, Toll BA, et al. Tobacco assessment in actively accruing National Cancer Institute Cooperative Group Program Clinical Trials. J Clin Oncol 2012;30:2869-2875. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22689794>.

19. McAfee T, Babb S, McNabb S, Fiore MC. Helping smokers quit--opportunities created by the Affordable Care Act. N Engl J Med 2015;372:5-7. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25409263>.

20. Malarcher A, Dube S, L. S, et al. Quitting Smoking Among Adults - United States, 2001-2010. Morbidity and Mortality Weekly Report. Vol. 60: Centers for Disease Control and Prevention; 2011:1513-1519. Available at: <http://www.cdc.gov/mmwr/preview/mmwrhtml/mm6044a2.htm#tab1>.

21. Twyman L, Bonevski B, Paul C, Bryant J. Perceived barriers to smoking cessation in selected vulnerable groups: a systematic review of the qualitative and quantitative literature. BMJ Open 2014;4:e006414. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25534212>.

22. Chapple A, Ziebland S, McPherson A. Stigma, shame, and blame experienced by patients with lung cancer: qualitative study. BMJ 2004;328:1470. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/15194599>.

23. Schnoll RA, Malstrom M, James C, et al. Correlates of tobacco use among smokers and recent quitters diagnosed with cancer. Patient

Educ Couns 2002;46:137-145. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/11867244>.

24. Morgan G, Schnoll RA, Alfano CM, et al. National cancer institute conference on treating tobacco dependence at cancer centers. J Oncol Pract 2011;7:178-182. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/21886500>.

25. Mazza R, Lina M, Boffi R, et al. Taking care of smoker cancer patients: a review and some recommendations. Ann Oncol 2010;21:1404-1409. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/20089563>.

26. Martinez E, Tatum KL, Weber DM, et al. Issues related to implementing a smoking cessation clinical trial for cancer patients. Cancer Causes Control 2009;20:97-104. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/18758971>.

27. U.S. National Library of Medicine. Key MEDLINE® Indicators. Available at: [http://www.nlm.nih.gov/bsd/bsd\\_key.html](http://www.nlm.nih.gov/bsd/bsd_key.html). Accessed May 2015.

28. Gritz ER, Fingeret MC, Vidrine DJ, et al. Successes and failures of the teachable moment: smoking cessation in cancer patients. Cancer 2006;106:17-27. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/16311986>.

29. Rink M, Zabor EC, Furberg H, et al. Impact of smoking and smoking cessation on outcomes in bladder cancer patients treated with radical cystectomy. Eur Urol 2013;64:456-464. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23206854>.

30. Ehdaie B, Furberg H, Zabor EC, et al. Impact of smoking status at diagnosis on disease recurrence and death in upper tract urothelial carcinoma. BJU Int 2013;111:589-595. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22642265>.



31. Braithwaite D, Izano M, Moore DH, et al. Smoking and survival after breast cancer diagnosis: a prospective observational study and systematic review. *Breast Cancer Res Treat* 2012;136:521-533. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23053660>.
32. Waggoner SE, Darcy KM, Fuhrman B, et al. Association between cigarette smoking and prognosis in locally advanced cervical carcinoma treated with chemoradiation: a Gynecologic Oncology Group study. *Gynecol Oncol* 2006;103:853-858. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/16815535>.
33. Gillison ML, Zhang Q, Jordan R, et al. Tobacco smoking and increased risk of death and progression for patients with p16-positive and p16-negative oropharyngeal cancer. *J Clin Oncol* 2012;30:2102-2111. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22565003>.
34. Walter V, Jansen L, Hoffmeister M, Brenner H. Smoking and survival of colorectal cancer patients: systematic review and meta-analysis. *Ann Oncol* 2014;25:1517-1525. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24692581>.
35. Modesitt SC, Huang B, Shelton BJ, Wyatt S. Endometrial cancer in Kentucky: the impact of age, smoking status, and rural residence. *Gynecol Oncol* 2006;103:300-306. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/16631234>.
36. Kountourakis P, Correa AM, Hofstetter WL, et al. Combined modality therapy of cT2N0M0 esophageal cancer: the University of Texas M. D. Anderson Cancer Center experience. *Cancer* 2011;117:925-930. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/20960497>.
37. Zheng Y, Cao X, Wen J, et al. Smoking affects treatment outcome in patients with resected esophageal squamous cell carcinoma who received chemotherapy. *PLoS One* 2015;10:e0123246. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25874561>.
38. Hoff CM, Grau C, Overgaard J. Effect of smoking on oxygen delivery and outcome in patients treated with radiotherapy for head and neck squamous cell carcinoma--a prospective study. *Radiother Oncol* 2012;103:38-44. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22385797>.
39. Sharp L, McDevitt J, Carsin AE, et al. Smoking at diagnosis is an independent prognostic factor for cancer-specific survival in head and neck cancer: findings from a large, population-based study. *Cancer Epidemiol Biomarkers Prev* 2014;23:2579-2590. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25128401>.
40. Ehdaie B, Furberg H, Zabor EC, et al. Comprehensive assessment of the impact of cigarette smoking on survival of clear cell kidney cancer. *J Urol* 2014;191:597-602. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24018238>.
41. Keizman D, Gottfried M, Ish-Shalom M, et al. Active smoking may negatively affect response rate, progression-free survival, and overall survival of patients with metastatic renal cell carcinoma treated with sunitinib. *Oncologist* 2014;19:51-60. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24309979>.
42. Parsons A, Daley A, Begh R, Aveyard P. Influence of smoking cessation after diagnosis of early stage lung cancer on prognosis: systematic review of observational studies with meta-analysis. *BMJ* 2010;340:b5569. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/20093278>.
43. Tao L, Wang R, Gao YT, Yuan JM. Impact of postdiagnosis smoking on long-term survival of cancer patients: the Shanghai cohort study. *Cancer Epidemiol Biomarkers Prev* 2013;22:2404-2411. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24319070>.
44. Warren GW, Kasza KA, Reid ME, et al. Smoking at diagnosis and survival in cancer patients. *Int J Cancer* 2013;132:401-410. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22539012>.



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45. Tammemagi CM, Neslund-Dudas C, Simoff M, Kvale P. Smoking and lung cancer survival: the role of comorbidity and treatment. *Chest* 2004;125:27-37. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/14718417>.

46. Lin Y, Yagyu K, Ueda J, et al. Active and passive smoking and risk of death from pancreatic cancer: findings from the Japan Collaborative Cohort Study. *Pancreatology* 2013;13:279-284. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/23719601>.

47. Kenfield SA, Stampfer MJ, Chan JM, Giovannucci E. Smoking and prostate cancer survival and recurrence. *JAMA* 2011;305:2548-2555.

Available at: <http://www.ncbi.nlm.nih.gov/pubmed/21693743>.

48. Chelghoum Y, Danaila C, Belhabri A, et al. Influence of cigarette smoking on the presentation and course of acute myeloid leukemia. *Ann Oncol* 2002;13:1621-1627. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/12377652>.

49. Ramamoorthy S, Luo L, Luo E, Carethers JM. Tobacco smoking and risk of recurrence for squamous cell cancer of the anus. *Cancer Detect Prev* 2008;32:116-120. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/18639388>.

50. Wyszynski A, Tanyos SA, Rees JR, et al. Body mass and smoking are modifiable risk factors for recurrent bladder cancer. *Cancer* 2014;120:408-414. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/24122218>.

51. Bishop JD, Killelea BK, Chagpar AB, et al. Smoking and breast cancer recurrence after breast conservation therapy. *Int J Breast Cancer* 2014;2014:327081. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/24693439>.

52. Han MA, Kim YW, Choi IJ, et al. Association of smoking history with cancer recurrence and survival in stage III-IV male gastric cancer patients. *Cancer Epidemiol Biomarkers Prev* 2013;22:1805-1812.

Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23904463>.

53. Joshu CE, Mondul AM, Meinhold CL, et al. Cigarette smoking and prostate cancer recurrence after prostatectomy. *J Natl Cancer Inst* 2011;103:835-838. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/21498781>.

54. Moreira DM, Aronson WJ, Terris MK, et al. Cigarette smoking is associated with an increased risk of biochemical disease recurrence, metastasis, castration-resistant prostate cancer, and mortality after radical prostatectomy: results from the SEARCH database. *Cancer* 2014;120:197-204. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/24127391>.

55. Ford MB, Sigurdson AJ, Petrulis ES, et al. Effects of smoking and radiotherapy on lung carcinoma in breast carcinoma survivors. *Cancer* 2003;98:1457-1464. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/14508833>.

56. van Leeuwen FE, Klokman WJ, Stovall M, et al. Roles of radiotherapy and smoking in lung cancer following Hodgkin's disease. *J Natl Cancer Inst* 1995;87:1530-1537. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/7563187>.

57. Balduyck B, Sardari Nia P, Cogen A, et al. The effect of smoking cessation on quality of life after lung cancer surgery. *Eur J Cardiothorac Surg* 2011;40:1432-1437; discussion 1437-1438. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/21498082>.

58. Erhunmwunsee L, Onaitis MW. Smoking cessation and the success of lung cancer surgery. *Curr Oncol Rep* 2009;11:269-274. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/19508831>.

59. Mason DP, Subramanian S, Nowicki ER, et al. Impact of smoking cessation before resection of lung cancer: a Society of Thoracic Surgeons General Thoracic Surgery Database study. *Ann Thorac Surg* 2009;88:362-370; discussion 370-361. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/19632374>.



60. Gajdos C, Hawn MT, Campagna EJ, et al. Adverse effects of smoking on postoperative outcomes in cancer patients. *Ann Surg Oncol* 2012;19:1430-1438. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22065194>.
61. Sharma A, Deeb AP, Iannuzzi JC, et al. Tobacco smoking and postoperative outcomes after colorectal surgery. *Ann Surg* 2013;258:296-300. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23059503>.
62. Ehlers SL, Gastineau DA, Patten CA, et al. The impact of smoking on outcomes among patients undergoing hematopoietic SCT for the treatment of acute leukemia. *Bone Marrow Transplant* 2011;46:285-290. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/20479707>.
63. Chang DW, Reece GP, Wang B, et al. Effect of smoking on complications in patients undergoing free TRAM flap breast reconstruction. *Plast Reconstr Surg* 2000;105:2374-2380. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/10845289>.
64. Spear SL, Ducic I, Cuoco F, Hannan C. The effect of smoking on flap and donor-site complications in pedicled TRAM breast reconstruction. *Plast Reconstr Surg* 2005;116:1873-1880. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/16327598>.
65. Marin VP, Pytynia KB, Langstein HN, et al. Serum cotinine concentration and wound complications in head and neck reconstruction. *Plast Reconstr Surg* 2008;121:451-457. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/18300961>.
66. Kuri M, Nakagawa M, Tanaka H, et al. Determination of the duration of preoperative smoking cessation to improve wound healing after head and neck surgery. *Anesthesiology* 2005;102:892-896. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/15851873>.
67. Chen AM, Chen LM, Vaughan A, et al. Tobacco smoking during radiation therapy for head-and-neck cancer is associated with unfavorable outcome. *Int J Radiat Oncol Biol Phys* 2011;79:414-419. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/20399030>.
68. Browman GP, Wong G, Hodson I, et al. Influence of cigarette smoking on the efficacy of radiation therapy in head and neck cancer. *N Engl J Med* 1993;328:159-163. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/8417381>.
69. Zevallos JP, Mallen MJ, Lam CY, et al. Complications of radiotherapy in laryngopharyngeal cancer: effects of a prospective smoking cessation program. *Cancer* 2009;115:4636-4644. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/19569250>.
70. Egestad H, Emaus N. Changes in health related quality of life in women and men undergoing radiation treatment for head and neck cancer and the impact of smoking status in the radiation treatment period. *Eur J Oncol Nurs* 2014;18:339-346. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24877857>.
71. Eifel PJ, Jhingran A, Bodurka DC, et al. Correlation of smoking history and other patient characteristics with major complications of pelvic radiation therapy for cervical cancer. *J Clin Oncol* 2002;20:3651-3657. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/12202666>.
72. Kucera H, Enzelsberger H, Eppel W, Weghaupt K. The influence of nicotine abuse and diabetes mellitus on the results of primary irradiation in the treatment of carcinoma of the cervix. *Cancer* 1987;60:1-4. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/3581022>.
73. Rades D, Setter C, Schild SE, Dunst J. Effect of smoking during radiotherapy, respiratory insufficiency, and hemoglobin levels on outcome in patients irradiated for non-small-cell lung cancer. *Int J Radiat Oncol Biol Phys* 2008;71:1134-1142. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/18258387>.
74. Pickles T, Liu M, Berthelet E, et al. The effect of smoking on outcome following external radiation for localized prostate cancer. *J Urol*



2004;171:1543-1546. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/15017216>.

75. Alsadius D, Hedelin M, Johansson KA, et al. Tobacco smoking and long-lasting symptoms from the bowel and the anal-sphincter region after radiotherapy for prostate cancer. *Radiother Oncol* 2011;101:495-501. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/21737169>.

76. Hooning MJ, Botma A, Aleman BM, et al. Long-term risk of cardiovascular disease in 10-year survivors of breast cancer. *J Natl Cancer Inst* 2007;99:365-375. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/17341728>.

77. Tyc VL, Hudson MM, Hinds P, et al. Tobacco use among pediatric cancer patients: recommendations for developing clinical smoking interventions. *J Clin Oncol* 1997;15:2194-2204. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/9196131>.

78. Tobacco Use During Cancer Treatment. American Society for Clinical Oncology; 2012. Available at: <http://www.cancer.net/navigating-cancer-care/prevention-and-healthy-living/tobacco-use/tobacco-use-during-cancer-treatment>. Accessed September 23, 2015.

79. Geng Y, Savage SM, Razani-Boroujerdi S, Sopori ML. Effects of nicotine on the immune response. II. Chronic nicotine treatment induces T cell anergy. *J Immunol* 1996;156:2384-2390. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/8786295>.

80. Kalra R, Singh SP, Savage SM, et al. Effects of cigarette smoke on immune response: chronic exposure to cigarette smoke impairs antigen-mediated signaling in T cells and depletes IP3-sensitive Ca(2+) stores. *J Pharmacol Exp Ther* 2000;293:166-171. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/10734166>.

81. Geng Y, Savage SM, Johnson LJ, et al. Effects of nicotine on the immune response. I. Chronic exposure to nicotine impairs antigen receptor-mediated signal transduction in lymphocytes. *Toxicol Appl*

*Pharmacol* 1995;135:268-278. Available at:

<http://www.ncbi.nlm.nih.gov/pubmed/8545837>.

82. Singh SP, Kalra R, Puttfarcken P, et al. Acute and chronic nicotine exposures modulate the immune system through different pathways. *Toxicol Appl Pharmacol* 2000;164:65-72. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/10739745>.

83. Birrell MA, Wong S, Catley MC, Belvisi MG. Impact of tobacco-smoke on key signaling pathways in the innate immune response in lung macrophages. *J Cell Physiol* 2008;214:27-37. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/17541958>.

84. Mehta H, Nazzal K, Sadikot RT. Cigarette smoking and innate immunity. *Inflamm Res* 2008;57:497-503. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/19109742>.

85. Kimura T, Shibata Y, Yamauchi K, et al. Oxidized phospholipid, 1-palmitoyl-2-(9'-oxo-nonanoyl)-glycerophosphocholine (PON-GPC), produced in the lung due to cigarette smoking, impairs immune function in macrophages. *Lung* 2012;190:169-182. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/21986851>.

86. Ferson M, Edwards A, Lind A, et al. Low natural killer-cell activity and immunoglobulin levels associated with smoking in human subjects. *Int J Cancer* 1979;23:603-609. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/457307>.

87. Tollerud DJ, Clark JW, Brown LM, et al. Association of cigarette smoking with decreased numbers of circulating natural killer cells. *Am Rev Respir Dis* 1989;139:194-198. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/2912340>.

88. Arcavi L, Benowitz NL. Cigarette smoking and infection. *Arch Intern Med* 2004;164:2206-2216. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/15534156>.



89. Nishioka T, Luo LY, Shen L, et al. Nicotine increases the resistance of lung cancer cells to cisplatin through enhancing Bcl-2 stability. *Br J Cancer* 2014;110:1785-1792. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24548862>.
90. Zhang J, Kamdar O, Le W, et al. Nicotine induces resistance to chemotherapy by modulating mitochondrial signaling in lung cancer. *Am J Respir Cell Mol Biol* 2009;40:135-146. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/18676776>.
91. Zhao J, Xin M, Wang T, et al. Nicotine enhances the antiapoptotic function of Mcl-1 through phosphorylation. *Mol Cancer Res* 2009;7:1954-1961. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/19903766>.
92. Shen T, Le W, Yee A, et al. Nicotine induces resistance to chemotherapy in nasal epithelial cancer. *Am J Rhinol Allergy* 2010;24:e73-77. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/20338106>.
93. Trevino JG, Pillai S, Kunigal S, et al. Nicotine induces inhibitor of differentiation-1 in a Src-dependent pathway promoting metastasis and chemoresistance in pancreatic adenocarcinoma. *Neoplasia* 2012;14:1102-1114. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23308043>.
94. O'Malley M, King AN, Conte M, et al. Effects of cigarette smoking on metabolism and effectiveness of systemic therapy for lung cancer. *J Thorac Oncol* 2014;9:917-926. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24926542>.
95. Hamilton M, Wolf JL, Rusk J, et al. Effects of smoking on the pharmacokinetics of erlotinib. *Clin Cancer Res* 2006;12:2166-2171. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/16609030>.
96. van der Bol JM, Mathijssen RH, Loos WJ, et al. Cigarette smoking and irinotecan treatment: pharmacokinetic interaction and effects on neutropenia. *J Clin Oncol* 2007;25:2719-2726. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/17563393>.
97. de Graan AJ, Loos WJ, Friberg LE, et al. Influence of smoking on the pharmacokinetics and toxicity profiles of taxane therapy. *Clin Cancer Res* 2012;18:4425-4432. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22645049>.
98. Peppone LJ, Mustian KM, Morrow GR, et al. The effect of cigarette smoking on cancer treatment-related side effects. *Oncologist* 2011;16:1784-1792. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22135122>.
99. Ditre JW, Gonzalez BD, Simmons VN, et al. Associations between pain and current smoking status among cancer patients. *Pain* 2011;152:60-65. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/21168758>.
100. Gonzalez A, Japuntich S, Keating NL, et al. Pain experiences among a population-based cohort of current, former, and never regular smokers with lung and colorectal cancer. *Cancer* 2014;120:3554-3561. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25043285>.
101. Logan HL, Fillingim RB, Bartoshuk LM, et al. Smoking status and pain level among head and neck cancer patients. *J Pain* 2010;11:528-534. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/20015696>.
102. Novy DM, Lam C, Gritz ER, et al. Distinguishing features of cancer patients who smoke: pain, symptom burden, and risk for opioid misuse. *J Pain* 2012;13:1058-1067. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23010143>.
103. The Health Benefits of Smoking Cessation: A Report of the Surgeon General. Atlanta, GA: U.S. Department of Health and Human Services 1990. Available at: <http://profiles.nlm.nih.gov/NN/B/B/C/T/>.



104. Sitas F, Weber MF, Egger S, et al. Smoking cessation after cancer. *J Clin Oncol* 2014;32:3593-3595. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25267760>.

105. Johnston-Early A, Cohen MH, Minna JD, et al. Smoking abstinence and small cell lung cancer survival. An association. *JAMA* 1980;244:2175-2179. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/6252357>.

106. Khuri FR, Kim ES, Lee JJ, et al. The impact of smoking status, disease stage, and index tumor site on second primary tumor incidence and tumor recurrence in the head and neck retinoid chemoprevention trial. *Cancer Epidemiol Biomarkers Prev* 2001;10:823-829. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/11489748>.

107. Fleshner N, Garland J, Moadel A, et al. Influence of smoking status on the disease-related outcomes of patients with tobacco-associated superficial transitional cell carcinoma of the bladder. *Cancer* 1999;86:2337-2345. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/10590376>.

108. Dobson Amato KA, Hyland A, Reed R, et al. Tobacco Cessation May Improve Lung Cancer Patient Survival. *J Thorac Oncol* 2015;10:1014-1019. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/26102442>.

109. Richardson GE, Tucker MA, Venzon DJ, et al. Smoking cessation after successful treatment of small-cell lung cancer is associated with fewer smoking-related second primary cancers. *Ann Intern Med* 1993;119:383-390. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/8393311>.

110. Garces YI, Schroeder DR, Nirelli LM, et al. Second primary tumors following tobacco dependence treatments among head and neck cancer patients. *Am J Clin Oncol* 2007;30:531-539. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/17921716>.

111. Baser S, Shannon VR, Eapen GA, et al. Smoking cessation after diagnosis of lung cancer is associated with a beneficial effect on performance status. *Chest* 2006;130:1784-1790. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/17166997>.

112. Garces YI, Hays JT. Tobacco dependence: why should an oncologist care? *J Clin Oncol* 2003;21:1884-1886. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/12721267>.

113. Warren GW, Marshall JR, Cummings KM, et al. Automated tobacco assessment and cessation support for cancer patients. *Cancer* 2014;120:562-569. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24496870>.

114. Lindson-Hawley N, Aveyard P, Hughes JR. Reduction versus abrupt cessation in smokers who want to quit. *Cochrane Database Syst Rev* 2012;11:CD008033. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23152252>.

115. Lindson-Hawley N, Aveyard P, Hughes JR. Gradual reduction vs abrupt cessation as a smoking cessation strategy in smokers who want to quit. *JAMA* 2013;310:91-92. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23821093>.

116. Ostroff JS, Burkhalter JE, Cinciripini PM, et al. Randomized trial of a presurgical scheduled reduced smoking intervention for patients newly diagnosed with cancer. *Health Psychol* 2014;33:737-747. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23895203>.

117. Miller WR, Rollnick S. *Motivational Interviewing: Preparing People for Change* (ed 2). New York, NY: The Guilford Press; 2002.

118. Rollnick S, Miller WR, Butler CC. *Motivational Interviewing in Health Care: Helping Patients Change Behavior* (ed 1). New York, NY: The Guilford Press; 2007.

119. Ehlers SL, Bronars CA, Patten CA, et al. Accuracy of self-reported tobacco use status among hematopoietic SCT patients. *Bone Marrow*



Transplant 2014;49:961-965. Available at:  
<http://www.ncbi.nlm.nih.gov/pubmed/24732958>.

120. Hald J, Overgaard J, Grau C. Evaluation of objective measures of smoking status--a prospective clinical study in a group of head and neck cancer patients treated with radiotherapy. *Acta Oncol* 2003;42:154-159. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/12801134>.

121. Warren GW, Arnold SM, Valentino JP, et al. Accuracy of self-reported tobacco assessments in a head and neck cancer treatment population. *Radiother Oncol* 2012;103:45-48. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22119370>.

122. Thomsen T, Villebro N, Moller AM. Interventions for preoperative smoking cessation. *Cochrane Database Syst Rev* 2014;3:CD002294. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24671929>.

123. Wong J, Abrishami A, Yang Y, et al. A perioperative smoking cessation intervention with varenicline: a double-blind, randomized, placebo-controlled trial. *Anesthesiology* 2012;117:755-764. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22890119>.

124. Moller AM, Villebro N, Pedersen T, Tonnesen H. Effect of preoperative smoking intervention on postoperative complications: a randomised clinical trial. *Lancet* 2002;359:114-117. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/11809253>.

125. Lindstrom D, Sadr Azodi O, Wladis A, et al. Effects of a perioperative smoking cessation intervention on postoperative complications: a randomized trial. *Ann Surg* 2008;248:739-745. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/18948800>.

126. Kotz D, Brown J, West R. 'Real-world' effectiveness of smoking cessation treatments: a population study. *Addiction* 2014;109:491-499. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24372901>.

127. Stead LF, Lancaster T. Combined pharmacotherapy and behavioural interventions for smoking cessation. *Cochrane Database*

*Syst Rev* 2012;10:CD008286. Available at:  
<http://www.ncbi.nlm.nih.gov/pubmed/23076944>.

128. Stead LF, Lancaster T. Behavioural interventions as adjuncts to pharmacotherapy for smoking cessation. *Cochrane Database Syst Rev* 2012;12:CD009670. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23235680>.

129. Hughes JR. Dependence potential and abuse liability of nicotine replacement therapies. *Biomed Pharmacother* 1989;43:11-17. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/2659095>.

130. Moore TJ, Furberg CD, Glenmullen J, et al. Suicidal behavior and depression in smoking cessation treatments. *PLoS One* 2011;6:e27016. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22073240>.

131. Thomas KH, Martin RM, Davies NM, et al. Smoking cessation treatment and risk of depression, suicide, and self harm in the Clinical Practice Research Datalink: prospective cohort study. *BMJ* 2013;347:f5704. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24124105>.

132. Thomas KH, Martin RM, Knipe DW, et al. Risk of neuropsychiatric adverse events associated with varenicline: systematic review and meta-analysis. *BMJ* 2015;350:h1109. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25767129>.

133. Hollands GJ, McDermott MS, Lindson-Hawley N, et al. Interventions to increase adherence to medications for tobacco dependence. *Cochrane Database Syst Rev* 2015;2:CD009164. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25914910>.

134. Cahill K, Stevens S, Perera R, Lancaster T. Pharmacological interventions for smoking cessation: an overview and network meta-analysis. *Cochrane Database Syst Rev* 2013;5:CD009329. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23728690>.



135. Coe JW, Brooks PR, Vetelino MG, et al. Varenicline: an alpha4beta2 nicotinic receptor partial agonist for smoking cessation. *J Med Chem* 2005;48:3474-3477. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/15887955>.

136. Mills EJ, Wu P, Lockhart I, et al. Comparisons of high-dose and combination nicotine replacement therapy, varenicline, and bupropion for smoking cessation: a systematic review and multiple treatment meta-analysis. *Ann Med* 2012;44:588-597. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22860882>.

137. Cahill K, Stead LF, Lancaster T. Nicotine receptor partial agonists for smoking cessation. *Cochrane Database Syst Rev* 2012;4:CD006103. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22513936>.

138. Ebbert JO, Hughes JR, West RJ, et al. Effect of varenicline on smoking cessation through smoking reduction: a randomized clinical trial. *JAMA* 2015;313:687-694. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25688780>.

139. Tonstad S, Tonnesen P, Hajek P, et al. Effect of maintenance therapy with varenicline on smoking cessation: a randomized controlled trial. *JAMA* 2006;296:64-71. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/16820548>.

140. Hajek P, McRobbie H, Myers Smith K, et al. Increasing varenicline dose in smokers who do not respond to the standard dosage: a randomized clinical trial. *JAMA Intern Med* 2015;175:266-271. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25545858>.

141. Gonzales D, Hajek P, Pliamm L, et al. Retreatment with varenicline for smoking cessation in smokers who have previously taken varenicline: a randomized, placebo-controlled trial. *Clin Pharmacol Ther* 2014;96:390-396. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24911368>.

142. Jorenby DE, Hays JT, Rigotti NA, et al. Efficacy of varenicline, an alpha4beta2 nicotinic acetylcholine receptor partial agonist, vs placebo or sustained-release bupropion for smoking cessation: a randomized controlled trial. *JAMA* 2006;296:56-63. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/16820547>.

143. Williams KE, Reeves KR, Billing CB, Jr., et al. A double-blind study evaluating the long-term safety of varenicline for smoking cessation. *Curr Med Res Opin* 2007;23:793-801. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/17407636>.

144. Prochaska JJ, Hilton JF. Risk of cardiovascular serious adverse events associated with varenicline use for tobacco cessation: systematic review and meta-analysis. *BMJ* 2012;344:e2856. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22563098>.

145. Svanstrom H, Pasternak B, Hviid A. Use of varenicline for smoking cessation and risk of serious cardiovascular events: nationwide cohort study. *BMJ* 2012;345:e7176. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23138033>.

146. Mills EJ, Thorlund K, Eapen S, et al. Cardiovascular events associated with smoking cessation pharmacotherapies: a network meta-analysis. *Circulation* 2014;129:28-41. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24323793>.

147. Sharma A, Thakar S, Lavie CJ, et al. Cardiovascular adverse events associated with smoking-cessation pharmacotherapies. *Curr Cardiol Rep* 2015;17:554. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25410148>.

148. Chelladurai Y, Singh S. Varenicline and cardiovascular adverse events: a perspective review. *Ther Adv Drug Saf* 2014;5:167-172. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25083271>.

149. Singh S, Loke YK, Spangler JG, Furberg CD. Risk of serious adverse cardiovascular events associated with varenicline: a systematic



review and meta-analysis. CMAJ 2011;183:1359-1366. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/21727225>.

150. Anthenelli RM, Morris C, Ramey TS, et al. Effects of varenicline on smoking cessation in adults with stably treated current or past major depression: a randomized trial. Ann Intern Med 2013;159:390-400. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24042367>.

151. Evins AE, Cather C, Pratt SA, et al. Maintenance treatment with varenicline for smoking cessation in patients with schizophrenia and bipolar disorder: a randomized clinical trial. JAMA 2014;311:145-154. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24399553>.

152. Smith SS, McCarthy DE, Japuntich SJ, et al. Comparative effectiveness of 5 smoking cessation pharmacotherapies in primary care clinics. Arch Intern Med 2009;169:2148-2155. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/20008701>.

153. Cahill K, Stevens S, Lancaster T. Pharmacological treatments for smoking cessation. JAMA 2014;311:193-194. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24399558>.

154. Stead LF, Perera R, Bullen C, et al. Nicotine replacement therapy for smoking cessation. Cochrane Database Syst Rev 2012;11:CD000146. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23152200>.

155. Fiore MC, Jaen CR, Baker TB, et al. Treating Tobacco Use and Dependence: 2008 Update. In: Panel TUaDG ed. A Public Health Service-Sponsored Clinical Practice Guideline. Rockville, MD; 2008. Available at: <http://www.ncbi.nlm.nih.gov/books/NBK63952/>.

156. Warren GW, Singh AK. Nicotine and lung cancer. J Carcinog 2013;12:1. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23599683>.

157. Martinez-Garcia E, Irigoyen M, Gonzalez-Moreno O, et al. Repetitive nicotine exposure leads to a more malignant and metastasis-prone phenotype of SCLC: a molecular insight into the importance of

quitting smoking during treatment. Toxicol Sci 2010;116:467-476. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/20457658>.

158. Maier CR, Hollander MC, Hobbs EA, et al. Nicotine does not enhance tumorigenesis in mutant K-ras-driven mouse models of lung cancer. Cancer Prev Res (Phila) 2011;4:1743-1751. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22027685>.

159. Murphy SE, von Weyarn LB, Schutten MM, et al. Chronic nicotine consumption does not influence 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone-induced lung tumorigenesis. Cancer Prev Res (Phila) 2011;4:1752-1760. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22027684>.

160. Stepanov I, Carmella SG, Briggs A, et al. Presence of the carcinogen N'-nitrosonornicotine in the urine of some users of oral nicotine replacement therapy products. Cancer Res 2009;69:8236-8240. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/19843845>.

161. Murray RP, Connett JE, Zapawa LM. Does nicotine replacement therapy cause cancer? Evidence from the Lung Health Study. Nicotine Tob Res 2009;11:1076-1082. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/19571249>.

162. Shields PG. Long-term nicotine replacement therapy: cancer risk in context. Cancer Prev Res (Phila) 2011;4:1719-1723. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22052338>.

163. Hughes JR, Stead LF, Hartmann-Boyce J, et al. Antidepressants for smoking cessation. Cochrane Database Syst Rev 2014;1:CD000031. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24402784>.

164. Stapleton J, West R, Hajek P, et al. Randomized trial of nicotine replacement therapy (NRT), bupropion and NRT plus bupropion for smoking cessation: effectiveness in clinical practice. Addiction 2013;108:2193-2201. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23859696>.



165. van der Meer RM, Willemsen MC, Smit F, Cuijpers P. Smoking cessation interventions for smokers with current or past depression. *Cochrane Database Syst Rev* 2013;8:CD006102. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23963776>.

166. Hays JT, Hurt RD, Rigotti NA, et al. Sustained-release bupropion for pharmacologic relapse prevention after smoking cessation: a randomized, controlled trial. *Ann Intern Med* 2001;135:423-433. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/11560455>.

167. Englisch S, Morgen K, Meyer-Lindenberg A, Zink M. Risks and benefits of bupropion treatment in schizophrenia: a systematic review of the current literature. *Clin Neuropharmacol* 2013;36:203-215. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24201231>.

168. Koegelenberg CF, Noor F, Bateman ED, et al. Efficacy of varenicline combined with nicotine replacement therapy vs varenicline alone for smoking cessation: a randomized clinical trial. *JAMA* 2014;312:155-161. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25005652>.

169. Ramon JM, Morchon S, Baena A, Masuet-Aumatell C. Combining varenicline and nicotine patches: a randomized controlled trial study in smoking cessation. *BMC Med* 2014;12:172. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25296623>.

170. Hajek P, Smith KM, Dhanji AR, McRobbie H. Is a combination of varenicline and nicotine patch more effective in helping smokers quit than varenicline alone? A randomised controlled trial. *BMC Med* 2013;11:140. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23718718>.

171. Evins AE, Cather C, Culhane MA, et al. A 12-week double-blind, placebo-controlled study of bupropion sr added to high-dose dual nicotine replacement therapy for smoking cessation or reduction in schizophrenia. *J Clin Psychopharmacol* 2007;27:380-386. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/17632223>.

172. Rose JE, Behm FM. Combination treatment with varenicline and bupropion in an adaptive smoking cessation paradigm. *Am J Psychiatry* 2014;171:1199-1205. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24934962>.

173. Ebbert JO, Hatsukami DK, Croghan IT, et al. Combination varenicline and bupropion SR for tobacco-dependence treatment in cigarette smokers: a randomized trial. *JAMA* 2014;311:155-163. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24399554>.

174. Aveyard P, Johnson C, Fillingham S, et al. Nortriptyline plus nicotine replacement versus placebo plus nicotine replacement for smoking cessation: pragmatic randomised controlled trial. *BMJ* 2008;336:1223-1227. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/18441375>.

175. Yalcin BM, Unal M, Pirdal H, Karahan TF. Effects of an anger management and stress control program on smoking cessation: a randomized controlled trial. *J Am Board Fam Med* 2014;27:645-660. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25201934>.

176. Hughes JR. Effects of abstinence from tobacco: valid symptoms and time course. *Nicotine Tob Res* 2007;9:315-327. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/17365764>.

177. McLaughlin I, Dani JA, De Biasi M. Nicotine withdrawal. *Curr Top Behav Neurosci* 2015;24:99-123. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25638335>.

178. Shiffman S, Patten C, Gwaltney C, et al. Natural history of nicotine withdrawal. *Addiction* 2006;101:1822-1832. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/17156182>.

179. Stead LF, Hartmann-Boyce J, Perera R, Lancaster T. Telephone counselling for smoking cessation. *Cochrane Database Syst Rev* 2013;8:CD002850. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23934971>.



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180. Civljak M, Stead LF, Hartmann-Boyce J, et al. Internet-based interventions for smoking cessation. Cochrane Database Syst Rev 2013;7:CD007078. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23839868>.

181. Stead LF, Lancaster T. Group behaviour therapy programmes for smoking cessation. Cochrane Database Syst Rev 2005:CD001007. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/15846610>.

182. Lancaster T, Stead LF. Individual behavioural counselling for smoking cessation. Cochrane Database Syst Rev 2005:CD001292. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/15846616>.

183. Hartmann-Boyce J, Lancaster T, Stead LF. Print-based self-help interventions for smoking cessation. Cochrane Database Syst Rev 2014;6:CD001118. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24888233>.

184. Whittaker R, McRobbie H, Bullen C, et al. Mobile phone-based interventions for smoking cessation. Cochrane Database Syst Rev 2012;11:CD006611. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23152238>.

185. Free C, Knight R, Robertson S, et al. Smoking cessation support delivered via mobile phone text messaging (txt2stop): a single-blind, randomised trial. Lancet 2011;378:49-55. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/21722952>.

186. Abrams LC, Boal AL, Simmens SJ, et al. A randomized trial of Text2Quit: a text messaging program for smoking cessation. Am J Prev Med 2014;47:242-250. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24913220>.

187. Abrams LC, Padmanabhan N, Thaweethai L, Phillips T. iPhone apps for smoking cessation: a content analysis. Am J Prev Med 2011;40:279-285. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/21335258>.

188. Stead LF, Buitrago D, Preciado N, et al. Physician advice for smoking cessation. Cochrane Database Syst Rev 2013;5:CD000165. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/23728631>.

189. Coleman T. ABC of smoking cessation. Use of simple advice and behavioural support. BMJ 2004;328:397-399. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/14962878>.

190. Aveyard P, Begh R, Parsons A, West R. Brief opportunistic smoking cessation interventions: a systematic review and meta-analysis to compare advice to quit and offer of assistance. Addiction 2012;107:1066-1073. Available at:

191. Smith SS, Keller PA, Kobinsky KH, et al. Enhancing tobacco quitline effectiveness: identifying a superior pharmacotherapy adjuvant. Nicotine Tob Res 2013;15:718-728. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/22992296>.

192. Mehnert A, Kuhnt S, Braehler E, et al. Twelve-month prevalence of mental disorders in cancer patients across major tumor entities [abstract]. ASCO Meeting Abstracts 2015;33:9552. Available at: [http://meeting.ascopubs.org/cgi/content/abstract/33/15\\_suppl/9552](http://meeting.ascopubs.org/cgi/content/abstract/33/15_suppl/9552).

193. Singer S, Das-Munshi J, Braehler E. Prevalence of mental health conditions in cancer patients in acute care--a meta-analysis. Ann Oncol 2010;21:925-930. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/19887467>.

194. Mehnert A, Braehler E, Faller H, et al. Four-week prevalence of mental disorders in patients with cancer across major tumor entities. J Clin Oncol 2014;32:3540-3546. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25287821>.

195. Lindson-Hawley N, Thompson TP, Begh R. Motivational interviewing for smoking cessation. Cochrane Database Syst Rev 2015;3:CD006936. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25726920>.



## NCCN Guidelines Version 2.2015 Smoking Cessation

196. Draft Recommendation Statement: Tobacco Smoking Cessation in Adults and Pregnant Women: Behavioral and Pharmacotherapy Interventions: U.S. Preventative Services Task Force; 2015. Available at: <http://www.uspreventiveservicestaskforce.org/Page/Document/draft-recommendation-statement147/tobacco-use-in-adults-and-pregnant-women-counseling-and-interventions1#Pod9>.

197. Brandon TH, Goniewicz ML, Hanna NH, et al. Electronic nicotine delivery systems: a policy statement from the American Association for Cancer Research and the American Society of Clinical Oncology. *J Clin Oncol* 2015;33:952-963. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25572671>.

198. Bhatnagar A, Whitsel LP, Ribisl KM, et al. Electronic cigarettes: a policy statement from the American Heart Association. *Circulation* 2014;130:1418-1436. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25156991>.

199. Borderud SP, Li Y, Burkhalter JE, et al. Electronic cigarette use among patients with cancer: characteristics of electronic cigarette users and their smoking cessation outcomes. *Cancer* 2014;120:3527-3535. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25252116>.

200. McRobbie H, Bullen C, Hartmann-Boyce J, Hajek P. Electronic cigarettes for smoking cessation and reduction. *Cochrane Database Syst Rev* 2014;12:CD010216. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25515689>.

201. Rahman MA, Hann N, Wilson A, et al. E-cigarettes and smoking cessation: evidence from a systematic review and meta-analysis. *PLoS One* 2015;10:e0122544. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25822251>.

202. Brown J, Beard E, Kotz D, et al. Real-world effectiveness of e-cigarettes when used to aid smoking cessation: a cross-sectional population study. *Addiction* 2014;109:1531-1540. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24846453>.

203. Ussher MH, Taylor AH, Faulkner GE. Exercise interventions for smoking cessation. *Cochrane Database Syst Rev* 2014;8:CD002295. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/25170798>.

204. White AR, Rampes H, Liu JP, et al. Acupuncture and related interventions for smoking cessation. *Cochrane Database Syst Rev* 2014;1:CD000009. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/24459016>.

205. Abbot NC, Stead LF, White AR, et al. Hypnotherapy for smoking cessation. *Cochrane Database Syst Rev* 2000:CD001008. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/10796583>.

206. Barnes J, Dong CY, McRobbie H, et al. Hypnotherapy for smoking cessation. *Cochrane Database Syst Rev* 2010:CD001008. Available at: <http://www.ncbi.nlm.nih.gov/pubmed/20927723>.