

# Cigarette Smoking as a Risk Factor for Delirium in Hospitalized and Intensive Care Unit Patients

## A Systematic Review

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### Abstract

**Background:** Active smokers are prevalent in hospitalized and critically ill patients. Cigarette smoking and nicotine withdrawal may increase delirium in these populations. This systematic review aims to determine whether active cigarette smoking increases the risk for delirium in hospitalized and intensive care unit (ICU) patients.

**Methods:** A systematic search of English-, Spanish-, and French-language articles published from 1966 to April 2013 was performed. Studies were included if they measured cigarette smoking as a risk factor and delirium as an outcome in adult hospitalized or ICU patients. Methodologic quality of studies was assessed using both the validated Newcastle Ottawa Scale and an additional evidence-based quality rating scale.

**Results:** A total of 14 cohort studies of surgical and ICU populations were included in the review. No studies in non-ICU inpatients were

identified. The incidence of delirium ranged from 9 to 52%, and the prevalence of active smokers ranged from 9 to 44%. The quality of assessment for active smoking varied widely. None of the studies used biochemical measures to determine cigarette smoke exposure. Of the six studies restricting the smoking group to active smokers only, active smoking was independently associated with delirium in one study, trended toward an association in one study, and showed a dose response in one study. Quantitative summary measures were not calculated due to study heterogeneity and missing data.

**Conclusions:** There is currently insufficient evidence to determine if cigarette smoking is a risk factor for delirium. Future studies should consider using biochemical measures of cigarette smoke exposure to objectively quantify smoking behavior.

**Keywords:** delirium; smoking; risk factors; critical illness; hospitalization

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Delirium is a clinical syndrome characterized by an acute alteration in attention and cognition. Delirium occurs in up to 56 and 80% of hospitalized (1) and ICU patients (2), respectively, and is associated with prolonged hospital stay (3),

subsequent functional and cognitive decline (4, 5), and even death (6, 7). Despite its profound impact on patients, we still have an incomplete understanding of the etiology and risk factors for delirium in hospitalized patients. Although many of

the known predisposing factors are nonmodifiable (e.g., advanced age, dementia, comorbidities), factors that are modifiable in the acute setting are potential targets for delirium prevention (e.g., sedation-induced coma,

immobilization, pain, disorientation, sleep deprivation).

Cigarette smoking may represent a potentially modifiable risk factor for delirium in hospitalized and ICU patients. The primary mechanism through which cigarette smoking is postulated to contribute to delirium is through nicotine withdrawal in the setting of abrupt smoking cessation due to acute illness and hospitalization (8–10). Studies suggest that delirium is caused by imbalances in neurotransmitters due to factors such as systemic inflammation, metabolic derangements, acute stress responses, and exposure to psychoactive medications (11). A leading hypothesized mechanism for delirium is deficiency of acetylcholine in the central nervous system (12). A relative deficiency in acetylcholine also plays a central role in the pathophysiology of nicotine withdrawal. Specifically, the up-regulation and desensitization of nicotinic acetylcholine receptors in the brain in the setting of chronic nicotine exposure and their subsequent unoccupied state during periods of abstinence are believed to be associated with withdrawal symptoms (13). Indeed, nicotine withdrawal and delirium share common features, such as confusion, restlessness, and irritability. Furthermore, the time course of nicotine withdrawal symptoms, which peak in the first week of abstinence and can last up to 2–4 weeks after cessation (14, 15), overlaps with the onset of delirium, which is commonly diagnosed at the time of presentation to the hospital or ICU, or several days postoperatively. If an association between active smoking and delirium is identified, nicotine replacement therapy should be investigated as a potential preventive intervention for hospitalized smokers at risk for delirium. Additional postulated mechanisms for how cigarette smoking increases delirium risk include microvascular changes and increasing atherosclerotic burden. However, these additional effects are likely due to chronic exposure rather than acute smoking cessation; hence, they are less likely to be modifiable in the setting of hospitalization, and are not feasible targets for preventative strategies.

Several studies have tested whether cigarette smoking is a risk factor for delirium. Thus far, these studies have provided conflicting results. Accordingly, we sought to determine whether smokers

who are admitted to the hospital or ICU are at a higher risk for delirium compared with nonsmokers by systematically reviewing the evidence. We hypothesize that active cigarette smoking is an independent risk factor for the development of delirium in these patient populations. Given the high prevalence of active smokers in both hospitalized and ICU populations (40–57%) (16–19), determining whether active smoking contributes to delirium is clinically relevant and important, especially because this represents a potentially modifiable risk factor through the use of nicotine replacement therapy. The results of this study have previously been published in abstract form (20).

## Methods

### Definition of Exposure and Outcome

This systematic review was performed according to *Meta-analysis of Observational Studies in Epidemiology Guidelines* (21). Studies were included if they met the following inclusion criteria: (1) subjects were adult inpatients ( $\geq 18$  yr old); (2) a history of active cigarette smoking was assessed; and (3) data on delirium were provided. The exposure measure of cigarette smoking was defined as biochemical measurement of cigarette smoke biomarkers, or smoking history obtained through self-report, family or surrogate report, or medical chart. The outcome measure of delirium was defined as an acute alteration in attention and cognition, as assessed using validated diagnostic criteria.

### Search Strategy

A PubMed search of English, Spanish, and French studies (1966 to April 2013), using the terms “delirium AND (habits OR smok\* OR smoking cessation OR nicotine OR cigarette\* OR tobacco)” was conducted (see Table E1 in the online supplement). Embase, CINAHL, PsycInfo, Web of Science, and the Centers for Disease Control and Prevention Tobacco Information and Prevention Databases were also searched. The search was supplemented by a manual search of bibliographies of retrieved articles and relevant published reviews. All articles that were deemed potentially relevant were located for full manuscript review.

### Study Selection and Data Abstraction

Studies were excluded if they met one or more of the following criteria: diagnosis of delirium was not based on examination of study subjects; lack of concurrent control group; or study only published in abstract form, case report, or literature review. Two authors (S.J.H. and M.S., or S.J.H. and A.N.L.) independently reviewed titles and abstracts to identify articles for inclusion and exclusion using *a priori* criteria. A log of included and excluded citations, including justification for exclusion, was maintained. In the end, 14 publications were selected for a detailed and independent review by two investigators (S.J.H. and M.S., or S.J.H. and A.N.L.). A structured abstraction form was used to record the author and year of the study, study design, patient characteristics, method of measuring cigarette smoke exposure, prevalence of smoking, delirium assessment method, incidence of delirium, covariates in multivariate model, and univariate and multivariate odds ratios (ORs). Disagreements in the abstracted data were resolved by further review, discussion, and consensus between the two reviewing authors. Attempts were made to contact authors of studies with unreported data ( $n = 8$ ). In the cases where contact was successful, the requested data were not available ( $n = 3$ ). No assumptions were made for missing data. To determine whether meta-analytical pooling would be appropriate, studies were assessed for variability in population type, study design, and outcome reporting.

### Quality Assessment

Two different methods were used to assess methodological quality of the studies. First, two physician reviewers independently assessed the studies using the Newcastle Ottawa Scale (NOS) (22), a validated scoring system that rates the selection, comparability, and assessment of outcome in cohort studies. However, the ability of the NOS to identify flaws in the definition and the assessment of cigarette smoking was limited (Table E2). This limitation could lead to misclassification of active cigarette smoking, and could bias the association between smoking and delirium. Therefore, we devised a scoring system *a priori* to critically concentrate on assessing this potential source of bias and to complement the NOS assessment (Quality

**Table 1.** Quality rating scale for assessment of active cigarette smoking

	Points
Smoking group designation	
Active smoker only	2
Active and former smokers, or not reported	0
Source of smoking assessment	
Biomarker	4
Self-report only	3
Self-, surrogate, or chart report	2
Chart report only	1
Unclear	0

rating scale for assessment of active cigarette smoking; Table 1). In this scale, the highest number of points was given to studies in which (1) the smoking group was restricted to active smokers only, and (2) the source of the smoking status assessment was objectively determined by biomarker measurement (as opposed to smoking history). The justification for the scoring system is as follows. Cigarette smoke biomarkers (e.g., cotinine) are sensitive and specific to smoking, and have been extensively used to establish causal relationships between smoking and disease in outpatient studies (23). Because well validated biomarker cutpoints accurately distinguish active smokers from second-hand smokers, biomarkers can be used to both identify active smokers and quantify the amount of cigarette smoking (24). These biomarkers have been shown to be superior to self-report, surrogate report, and medical records in the setting of inpatient and ICU populations (17, 19). Self-report can be limited by social desirability bias, recall bias, and, in the case of critically ill patients, may not be feasible due to respiratory failure and altered levels of consciousness. Compared with self-report, surrogates tend to underreport overall smoking status, and the accuracy of amount and duration smoked is even more limited (25, 26). Agreement between smoking history obtained from medical records compared with self-report is even poorer, and data on smoking history are often missing. Therefore, smoking history obtained through medical records was given the lowest credit (27, 28). Studies that restricted the smoking group to only active smokers were given credit, whereas studies that included both active and former smokers,

or did not specify the smoking status of the subjects in the smoking group, were given no credit.

**Results**

The comprehensive search yielded 489 unique articles (Figure 1). The 14 articles that were included in the qualitative review were published between 2001 and 2013 for 4,382 patients in 14 different countries (reviewer interrater  $\kappa = 0.92$ ; Table 2) (29–42). All were cohort studies, and only one study was designed to test smoking as the primary risk factor for delirium (34). The 13 remaining studies were designed to assess for multiple risk factors associated with the development of delirium. Study populations in the reviewed literature were composed of three main groups: (1) critically ill patients; (2) cardiovascular surgery patients; and (3) other surgical patients. No studies of adult inpatients were identified. A total of 4 of the 14 studies focused on older adults, a population that is at particularly high risk for developing delirium (11). Quality rating scores using the NOS varied minimally between studies (Table 3).

**Assessment of Active Cigarette Smoking**

The prevalence of active smoking ranged from 9 to 44%, and was not reported in 5 of the 14 studies (Table 2). Six studies restricted the smoking group to active smokers only (29, 31, 32, 36, 39, 41), three studies included both active and former smokers (29, 34, 37, 38), and five did not define the smoking group (30, 33, 35, 40, 42). The source of smoking history also varied widely between studies, ranging from self-report by patients before surgery to chart review only. Four of the studies did not describe how smoking history was obtained (30, 33, 35, 42). None of the studies used cigarette smoke biomarkers to determine smoking status. Overall, five studies had moderate quality assessments for active smoking (score 4–5 out of 6), five studies had low quality assessments (score 2–3), and four studies had poor quality assessments (score 0–1).

**Assessment of Delirium**

The measured incidence of delirium ranged from 9 to 52%. Delirium was diagnosed by Diagnostic and Statistical Manual of

Mental Disorders III or IV criteria, or by validated delirium screening tools. The frequency of delirium assessments ranged from once daily in most studies to several times a day. Three studies did not report the frequency of delirium assessments (31, 35, 41). The experience and training of the assessor performing delirium screening varied widely, from experienced study staff and study physicians to bedside nurses (34).

**Association between Active Smoking and Delirium**

Quantitative summary measures were not calculated, due to study heterogeneity and missing data. Of the six studies that restricted the smoking group to active smokers only (29, 31, 32, 36, 39, 41), a history of active smoking before hospital admission was independently associated with incident delirium in one study on elderly patients who had undergone coronary artery bypass graft surgery (odds ratio [OR], 4.19; 95% confidence interval [CI], 1.35–13.05;  $P = 0.019$ ) (39), and trended toward an association in a study on medical and surgical ICU patients (OR, 2.2; 95% CI, 0.94–4.94) (32). In a third study on patients who had undergone aortic aneurysm repair, the total number of pack-years smoked was independently associated with delirium (OR, 1.05; 95% CI, 1.02–1.08;  $P = 0.001$ ), although active smoking was not (29). Two of the six studies reported a statistically significant increased risk of incident delirium ( $P < 0.05$ ) in active smokers, but that association was lost after controlling for confounders (36, 41). The study with the poorest quality assessment of active smoking did not find an association between active smoking and delirium (31). Although all six studies assessed for advanced age and alcohol as potential risk factors in univariate analyses, the selection of covariates in multivariate analyses varied between studies. Specifically, factors that have been previously demonstrated to have robust associations with a greater risk of delirium, such as age, alcohol use, pre-existing cognitive impairment or dementia, severity of illness, comorbidity burden, and sedation with benzodiazepines, were variably adjusted for (Table 3). Although five out of the six studies adjusted for age, only three adjusted for benzodiazepine use (29, 36, 41), and two adjusted for cognitive impairment or dementia (29, 41). An

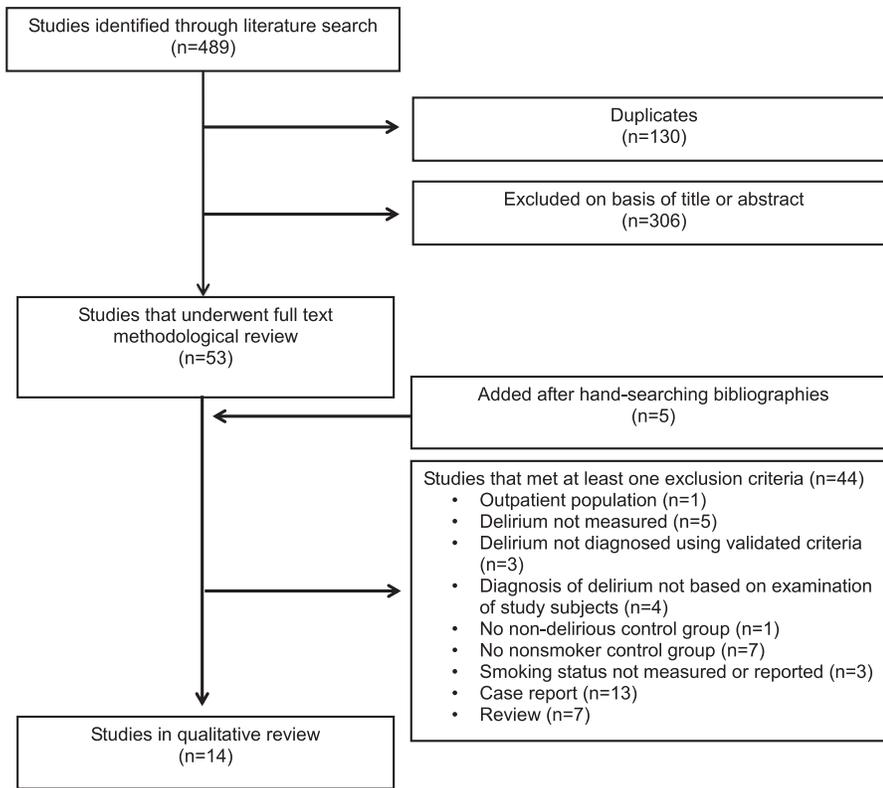


Figure 1. Literature search and selection. Modified by permission from Reference 20.

additional potential confounder for the association between smoking and delirium is prehospitalization depression (43). None of the included studies accounted for depression in their multivariate analyses.

Only 1 of the 14 studies was designed to assess for smoking as the primary risk factor for delirium (34). However, the study was powered for a combined outcome of ICU delirium and agitation, rather than delirium only, and included both active and former smokers in the smoking group. It did not find an association between active/former smoking and delirium in univariate analyses (OR, 1.84; 95% CI, 0.84–3.90), although active/former smoking was independently associated with increased agitation (OR, 3.13; 95% CI, 1.45–6.74). Of note, standardization of delirium assessments was variable in this study.

Of the eight low- to poor-quality assessment studies that either included both active and former smokers in the smoking group (37, 38, 40), or did not specify how cigarette smoke exposure was defined (30, 33–35, 42), smoking was associated with delirium in three studies (30, 35, 38), and trended toward an association in one (OR,

2.78; 95% CI, 0.94–8.24;  $P < 0.1$ ) (40). Only two of these studies included smoking in their multivariate analyses. One reported a trend toward an association between smoking and delirium (OR, 1.6; 95% CI, 1.0–2.6) (38), and one found no association (OR, 2.2; 95% CI, 0.6–8.8) (40).

## Discussion

Although case reports and reviews have suggested that cigarette smoking may increase the risk of delirium in hospitalized and ICU patients, the literature on this association has yielded conflicting results. In the first reported systematic review of the literature to date, we found that there is currently insufficient evidence to determine whether cigarette smoking is a risk factor for delirium in hospitalized and ICU patients.

Delirium is a common and serious form of acute brain injury in hospitalized and ICU patients that carries an enormous societal and financial burden (44–46). In view of the high prevalence of cigarette smoking in these populations, determining whether smoking is a risk factor for

delirium is of extreme importance, as nicotine withdrawal in active smokers may be a potentially modifiable target for both delirium prevention and treatment via nicotine replacement therapy. This question is particularly timely, given the recent debate on the benefits and safety of nicotine replacement therapy in critically ill patients. In fact, a recent prospective cohort study of critically ill smokers suggested that nicotine replacement therapy was associated with increased delirium (47). Currently, no studies support the use of nicotine replacement therapy for delirium prevention or treatment, and, given the potential risks, it should not be routinely used for delirium prevention or treatment until further study.

This systematic review has identified several deficiencies in the existing literature that likely account for the inconclusive nature of the findings. First and foremost, the existing literature was limited by suboptimal assessment of active smoking status. Because risk estimates can be significantly biased with only minor degrees of misclassification (48), an accurate quantitative assessment of active smoking is imperative when investigating the relationship between smoking and delirium. In this systematic review, none of the studies used biomarkers to assess for smoking status, which have been clearly demonstrated to identify more active smokers than self-report in both inpatient and ICU studies (17–19). An alternative measure of cigarette smoking is to determine the degree of nicotine dependence by a validated scale. However, these scales require patient participation, and have not been validated for surrogate use. Therefore, they could not be feasibly administered to delirious patients nor to critically ill patients who are comatose or in respiratory failure, and would not be useful in guiding management. Of note, only one of the reviewed studies assessed nicotine dependence using a validated scale (34). However, it did not investigate the association between nicotine dependence and delirium. In sum, to determine the true impact of acute cessation of active smoking on the development of delirium, future studies should consider using more quantitative and objective measures of active smoking than self-, surrogate-, and medical record report.

Analyzed studies were also limited by incomplete adjustment for potential

Table 2. Summary of included studies

Study No.	First Author, Year, (Ref. No.)	Country, Location	Sample Size	Age (Yr)	Population	Incidence of Delirium n (%)	Delirium Assessment Method	Delirium Assessment Frequency and Duration	Prevalence of Active Smoking (%)
1	Dubois, 2001 (32)	Canada, single center	198	≥18	Medical and surgical ICU, intubated and nonintubated	38 (19)	ICDSC ≥ 4 with psychiatric confirmation	q8 h × 5 d	NR
2	Lucidarme, 2010 (34)	France, multicenter	144	≥18	Medical and surgical ICU, intubated only	40 (28)	ICDSC > 4	Twice daily until ICU discharge	31
3	Ouimet, 2007 (36)	Canada, single center	764	≥18	Medical and surgical ICU, intubated and nonintubated	243 (32)	ICDSC ≥ 4	q1–8 h until ICU discharge	NR
4	Van Rompaey, 2009 (41)	Belgium, multicenter	523	≥18	Medical and surgical ICU, nonintubated only	155 (30)	NEECHAM confusion scale ≤ 19	NR	25
5	Benoit, 2005 (29)	Canada, single center	102	41–88	Elective aortic aneurysm repair surgery	34 (33)	DSM4	Daily × 6 d postop	28
6	Bohner, 2003 (30)	Germany, single center	153	≥18	Elective arterial surgery	60 (39)	DSM4 and DRS ≥ 12	Daily × 7 d postop	43
7	Chang, 2008 (31)	Taiwan, single center	288	≥20	Elective and emergent cardiac surgery	120 (42)	DSM4	NR	NR
8	Koster, 2008 (33)	The Netherlands, single center	112	>45	Elective cardiac surgery	24 (21)	DOS ≥ 2 with DSM4 confirmation	3 times daily × 5 d postop	15
9	Mardani, 2012 (35)	Iran, single center	196	≥48–80	Elective CABG	34 (17)	DSM4 in patients with MMSE ≤ 23	NR	NR
10	Rudolph, 2009 (37)	United States, multicenter	122	≥60	Elective cardiac surgery	63 (52)	Delirium symptom interview and MDAS, or CAM-ICU	Preop; daily beginning 2 d postop	NR
11	Rudolph, 2007 (38)	International, multicenter	1161	≥60	Elective noncardiac surgery	99 (9)	DSM3	Daily × 7 d postop	9
12	Santos, 2004 (39)	Brazil, single center	220	>60	Elective CABG	74 (34)	DSM4	Daily × 5 d postop	15
13	Yoshimura, 2004 (40)	Japan, single center	100	≥18	Elective liver resection for HCC	17 (17)	CAM	Daily × 2 weeks postop	44
14	Zakriya, 2002 (42)	United States, single center	168	50–98	Surgical hip fracture repair	47(28)	CAM	Daily postop until discharge	16

Definition of abbreviations: CABG = coronary artery bypass graft surgery; CAM = Confusion Assessment Method; DOS = Delirium Observation Screening Scale; DRS = Delirium Rating Scale; DSM = Diagnostic and Statistical Manual of Mental Disorders; HCC = hepatocellular carcinoma; ICDSC = Intensive Care Delirium Screening Checklist; ICU = intensive care unit; MDAS = Memorial Delirium Assessment Scale; MMSE = Mini Mental Status Exam; NEECHAM = Neelon and Champagne Confusion Scale; NR = not reported; postop = postoperatively; preop = preoperatively. All studies have prospective cohort design.

Table 3. Results from individual studies arranged by quality of smoking history

First Author, Year (Ref. No.)	Exposed	Unexposed	Smoking Assessment Method	Prevalence of Active Smoking(%)	Univariate OR for Delirium	Multivariate OR for Delirium*	Covariates in Multivariate Model	NOS Score (0-8)	Quality of Smoking Assessment (0-6)
Santos, 2004 (39)	Current smoker	Former + never smoker	Preoperative self-report	15	1.96 (0.93-4.11)	4.19 (1.35-13.05)	1,3,6,8	6	5
Benoit, 2005 (29)	Current smoker	Former + never smoker	Preoperative self-report	28	1.16 (0.46-2.91)	NR†	1,2,3,5,9	8	5
Dubois, 2001 (32)	≥20 cigarettes/d until admission	<20 cigarettes/d until admission	Self-, surrogate or chart report (in order of preference)	NR	2.2 (1.07-4.51)	2.2 (0.94-4.94)	1,6,10,11	5	4
Van Rompaey, 2009 (41)	> 10 cigarettes/d	≤10 cigarettes/d	Self- or surrogate report	25	2.04 (1.05-3.95)	NR	1,2,5,7,9,10,11	6	4
Ouimet, 2007 (36)	Current smoker	Former + never smoker	Self-, surrogate, report or chart report (in order of preference)	NR	NR‡, P = 0.0123	NR	1,2,3,6,7,9,10,11	7	4
Chang, 2008 (31)	≥20 cigarettes/d within 1 mo of surgery	<20 cigarettes/d within 1 mo of surgery	Chart report	NR	NR, P > 0.05	—	—	5	3
Rudolph, 2007 (38)	Current + former smoker	Never smoker	Self-report	9	RR, 1.8 (1.2-2.8)	1.6 (1.0-2.6)	1,3,4,5,6,8	7	3
Rudolph, 2009 (37)	Current + Former smoker: (a) 1-30 pack-years; (b) > 30 pack-years	Never smoker	Self-report	NR	1-30 pack-years RR, 1.1 (0.76-1.6); > 30 pack-years RR, 0.8 (0.5-1.3)	—	—	6	3
Yoshimura, 2004 (40)	"History of smoking"	Quit > 6 mo + never smoker	Self-report	44	2.78 (0.94-8.24)	2.2 (0.6-8.8)	1,3,6	7	3
Lucidarme, 2010 (34)	Current smoker, quit ≤ 6 months	Quit > 6 mo + never smoker	Self- or surrogate report	31	1.84 (0.84-3.90)	Did not perform	—	6	2
Bohner, 2003 (30)	"History of smoking"	NR	NR	43	1.98 (1.03-3.83)	—	—	5	0
Koster, 2008 (33)	"Cigarette use"	NR	NR	15	0.76 (0.20-2.87)	—	—	6	0
Mardani, 2012 (35)	"History of smoking"	NR	NR	NR	8.36 (1.9-37.9)	—	—	5	0
Zakriya, 2002 (42)	"Smoking"	NR	NR	16	1.09 (0.44-2.70)	—	—	6	0

Definitions of abbreviations: NOS = Newcastle Ottawa Scale (see Table E2); NR = not reported; OR = odds ratio; RR=relative risk. Studies that restricted the smoking group to current smokers only are placed above other studies in the table. Covariates included in multivariate model: (1) smoking; (2) alcohol; (3) age; (4) sex; (5) cognitive impairment or dementia; (6) comorbidity; (7) severity of illness; (8) intraoperative/postoperative factors; (9) benzodiazepines; (10) opioids; (11) hospital/intensive care unit factors. Quality of smoking assessment score (6 points maximum): assessment for active smoking: 2 = current smoker only; 0 = current and former smoker, or not reported; source of assessment: 4 = biomarker; 3 = self; 2 = self, surrogate, or chart; 1 = chart only; 0 = not reported. \*Cigarette smoking not included in multivariate model. †No association between active smoking and delirium (OR not reported), but OR for delirium 1.05 (95% CI = 1.02-1.08) per pack-year smoked. ‡OR not reported but study stated active smoking was associated with increased risk for delirium.

confounders of delirium. Factors such as pre-existing cognitive impairment, opioid and benzodiazepine use, and severity of illness have been consistently identified as predisposing and precipitating factors for delirium in both hospitalized and ICU patients (49). These factors were variably measured in the analyzed studies. Depression, an important potential confounder for the relationship between smoking and delirium, was not adjusted for in any of the studies. Because all analyzed studies were observational by design, this limitation could have biased the results.

In studies of critically ill patients, an additional limitation was the lower incidence of delirium (19–32%) compared with previously reported studies (46–80%) (2, 50). This discordance occurred despite the use of validated methods to assess delirium, and may be due to variable delirium definitions (51), variable delirium assessment tools (36), variability in sedation practices, and lower severity of illness of the cohorts (32). Furthermore variability in training for delirium screening (34) can lead to decreased identification of delirious patients, as demonstrated by recent work that showed that detection of delirium by bedside ICU nurses was lower compared with trained researchers (52).

There are some additional limitations to our conclusions. First, none of the studies was powered to investigate the association between active smoking and delirium. Second, nonsignificant ORs were not reported. This not only precludes the generation of a quantitative summary measure, but also biases interpretation of the results. Third, although evidence based, the quality scoring system used for this review has not been validated. However, it has face validity, because biomarker assessment for active smoking has been demonstrated to be a superior measure for cigarette smoke exposure than smoking history in both hospitalized and ICU patients. Fourth, although the intended scope of the review was inpatient and critically ill populations, no inpatient studies met search criteria. Future studies should investigate the association between active smoking and delirium in this important patient population. Fifth, because of individual variation in nicotine withdrawal symptoms, symptom severity and presence of delirium may not be directly associated with intensity of smoking. This question would best be answered with a prospective study. Finally, gray literature was not included in the systematic review, and thus our findings may be limited by publication lag.

These limitations notwithstanding, the strengths of this study are: (1) the comprehensive search of the evidence; (2) the relatively stringent inclusion criteria; and (3) the evidence-based assessment scale to assess the quality of smoking history.

## Conclusions and Implications

In this first systematic review of the literature on active cigarette smoking as a risk factor for delirium in surgical and ICU patients, we found insufficient evidence to determine whether an association exists. Considering the high prevalence of cigarette smokers and delirium in hospitalized and critically ill patients, and the potential preventative and therapeutic implications, a study is needed to carefully and decisively determine whether this association exists. Future studies should be designed to specifically investigate this association, and should use biochemical measures of cigarette smoking to objectively quantify smoking behavior. ■

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