

The major risk factors for delirium in a clinical setting

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Objective: We aimed to determine the major risk factors for the development of delirium in patients at a single general hospital by comparison with a control group.

Subjects and methods: We reviewed the medical records of 260 delirium patients and 77 control patients. We investigated age, sex, and risk factors for delirium in the total delirium group (n=260), the delirium medical subgroup (n=142), and the delirium surgical subgroup (n=118). Logistic regression analysis adjusting for age and sex was performed to identify the odds ratio.

Results: The mean age and the percentage of males were significantly higher in the delirium group compared with the control group (68.9 vs 54.3 years and 70% vs 41.6%, respectively). Risk factors for the delirium group were lower plasma albumin, hypertension, mechanical ventilation, and antipsychotic drug use. Plasma sodium level and hypertension were important risk factors for the delirium medical subgroup. Stroke history, hypertension, ICU care, and medication were important risk factors for the delirium surgical subgroup.

Conclusion: Lower plasma albumin, hypertension, mechanical ventilation, and antipsychotic drug use are important risk factors for delirium.

Keywords: delirium, acute confusional state, psychiatric consultation, risk factor

Introduction

The *Diagnostic and Statistical Manual of Mental Disorders, Fifth Edition (DSM-5)* defines delirium as a disturbance in attention and awareness over a short period of time, representing a change from baseline attention and awareness, with a tendency to fluctuate in severity during the course of a day.¹ Delirium is a very common disorder affecting ~15% of medical elderly inpatients.² In a general population study, the prevalence of delirium was 1%–2% in subjects >65 years old and 10% in subjects >85 years old.³ Patients with delirium have higher mortality and poor long-term prognosis.^{4,5} These results are contrary to the common belief that delirium is a reversible condition and suggest that active intervention should begin at diagnosis due to the adverse effects of delirium on long-term functional outcomes.^{6–8} A recent systematic review of prospective trials suggested that some pharmacological interventions, such as the use of haloperidol, showed success in preventing delirium and in decreasing the duration and/or severity of ongoing delirium, which is associated with better long-term outcomes.⁹

The most important aspect in the treatment and management of delirium is identifying and correcting the underlying causes. The causes of delirium are old age, male sex, underlying cognitive impairment,^{10,11} infection, fever,^{12,13} hypoxia, hypoglycemia, electrolyte imbalance, general medical conditions such as cardiovascular, renal, and hepatic diseases,¹⁴ use of drugs such as benzodiazepine and narcotic analgesics,^{15,16} metabolic disorders such as encephalopathy, and central nervous system diseases such as stroke,

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traumatic brain injury, and epilepsy.^{17,18} In addition, major surgical procedures such as orthopedic and cardiovascular surgeries, conditions associated with major surgeries such as somatic pain and cobalamin deficiency,^{19,20} and treatment environments such as ICUs are known risk factors.²¹

Most previous studies have investigated individual risk factors for delirium separately in their respective clinical situations. Accordingly, their subjects were limited to those with specific diseases or treatments, and their outcomes identified statistically significant risk factors for delirium.^{22–30} When the results of these studies were compared with a meta-analysis of the risk factors for delirium, some risk factors such as age and the use of benzodiazepine were relatively consistent, but others were not.^{31–33} For example, the presence of hypertension or hypoalbuminemia was not associated with delirium in one prospective study, while other evidence suggests that these factors are associated with delirium.¹⁴ Few studies have comprehensively examined the reported risk factors, and no study has examined their relative importance.

In this study, we investigated the risk factors for delirium in patients referred for psychiatric consultation, diagnosis, and management. We aimed to identify the risk factors for inpatients who underwent a major surgery under general anesthesia and patients who did not.

Subjects and methods

Subjects

We selected the delirium group and the control group from among patients referred to psychiatrists from August 2013 to July 2014. The delirium group included patients who were interviewed individually by psychiatrists and diagnosed with delirium according to DSM-5 criteria. We excluded patients whose cognitive impairments or psychiatric symptoms were caused by a psychiatric or neurologic disease, such as hallucination, delusion, or abnormal behavior caused by schizophrenia or schizoaffective disorder, or mood symptoms resulting from major mood episodes. We also excluded patients with Alzheimer's disease, vascular dementia, or dementia with Lewy bodies. Patients with substance use disorder were excluded because they might have had a cognitive impairment indistinguishable from delirium. The control group was selected from among patients referred to psychiatrists but not diagnosed with delirium.

Depending on the nature of the psychomotor disturbance and DSM-5 specifier definition, delirium was classified as follows: 1) hyperactive – increased psychomotor activity that may be accompanied by mood lability, agitation, and/or refusal to cooperate with medical care; 2) hypoactive – decreased psychomotor activity that may be accompanied by

sluggishness, lethargy, or stupor; or 3) mixed – normal psychomotor activity with disturbed attention and awareness, or rapid fluctuation between hyperactive and hypoactive states of delirium.¹ Patients were deemed to have psychotic symptoms when further information from caregivers indicated that there was evidence of hallucination or delusion.

The institutional review board of the Asan Medical Center approved the study, including the data collection and analysis. As this study was performed retrospectively, informed consent was not considered necessary by the institutional review board.

Methods

This study collected data from medical records written by psychiatrists who evaluated the patients. The delirium group was divided into medical and surgical subgroups, each of which was compared with the control group. The delirium medical subgroup included patients who were admitted to the hospital but did not undergo surgery under general anesthesia, while the delirium surgical subgroup included patients who did undergo surgery under general anesthesia. The overall severity of delirium was assessed by the Clinical Global Impression-Severity (CGI-S) scale.

Age, sex, and risk factors for delirium were investigated in the delirium and control groups. Risk factors consistently found in previous studies were comprehensively included and categorized by past medical history, treatment environment, physical illness, and medication history. Past medical history included delirium, dementia, stroke, gait disorder, falling, neurologic diseases, and hypertension. In addition, alcohol use within 1 week before delirium onset was noted, and activities of daily living (ADL) were evaluated on a four-point scale using the caregiver interview conducted by a psychiatrist (1: requires maximum aid for daily living; 2: requires moderate aid; 3: requires minimum aid; 4: does not require aid).

Treatment environments included ICU treatment and the preventive use of chemical or physical restraints before the occurrence of delirium. Chemical restraint was defined as intravenous or intramuscular injection of a benzodiazepine or an antipsychotic drug, and physical restraint was defined as restraint of the upper or lower limbs. Chemical or physical restraint used for patient safety after the occurrence of delirium was not included as a risk factor.

Physical illness included surgery under general anesthesia after admission, use of mechanical ventilation, and infection. Infection was defined by laboratory tests, the identification of pathogens in bacterial culture, or the empirical use of antibiotics. The patient was considered immobile if he or she was unable to ambulate independently due to a physical

impairment such as fracture or a neurological disease, or if absolute bed rest was ordered by the medical staff. Renal function was assessed by evaluating the blood urea nitrogen (BUN) and plasma creatinine (CR) levels, and hepatic function was assessed by estimation of blood levels of aspartate transaminase (AST) and alanine transaminase (ALT), as well as plasma albumin levels. Electrolyte imbalance was assessed by determining sodium and potassium levels.

For medication history, all drugs used within 3 days before the occurrence of delirium were explored. The overall number of medications was counted, and the use of antipsychotic drugs, anticholinergic drugs, antiepileptic drugs, and narcotic analgesics was investigated.

Statistical methods

We compared age, sex, prevalence of psychotic symptoms manifested as hallucinations or delusions, and the frequency of the delirium type between the delirium and control groups. We tried to identify the potential risk factors for delirium in two steps. In Step 1, we used Student's *t*-tests or chi-square tests for demographic and clinical variables. In Step 2, variables with $P < 0.1$ in Step 1 were entered into the next analysis of multiple logistic regression. We identified the odds ratio (OR) for each factor's contribution to the occurrence of delirium using multiple logistic regression analysis, with adjustment for age and sex. Statistical analyses were undertaken using SPSS version 18 (IBM Corporation, Armonk, NY, USA), and a P -value < 0.05 was considered statistically significant.

Results

We included 260 patients in the delirium group and 77 patients in the control group. Among the patients in the delirium group, 142 patients (54.6%) were classified into the delirium medical subgroup and 118 patients (45.4%) into the delirium surgical subgroup. Most of the patients in the control group were diagnosed with major depression, adjustment disorder, sleep disorder, panic disorder, or unspecified anxiety disorder.

The mean age was 68.9 years (standard deviation [SD]: ± 13.0 years) in the delirium group and 54.3 years (SD: ± 16.8 years) in the control group. In addition, 182 patients (70%) in the delirium group and 32 patients (41.6%) in the control group were male. The mean age of the delirium group was significantly higher than that of the control group ($t = -8.05$, $P < 0.001$), and there were more male patients in the delirium group than in the control group (70% vs 41.6%, $\chi^2 = 20.7$, $P < 0.001$). In the delirium group, 167 patients (64.2%) had hyperactive delirium, 18 patients (6.9%) had hypoactive delirium, and 75 patients (28.8%) had mixed delirium. Regarding psychotic symptoms, 91 patients (35%)

experienced hallucination and 52 patients (20%) experienced delusion. The mean severity was a Clinical Global Impression–Severity scale (CGI-S) score of 4.0.

There were no significant differences in terms of age, sex, type of delirium, frequency of psychotic symptoms, or CGI-S score between the two subgroups. There was a significant difference in age between the medical subgroup (69.3 \pm 13.9 years, $t = -7.06$, $P < 0.001$) and the surgical subgroup (68.4 \pm 11.9 years, $t = -6.86$, $P < 0.001$) compared with the control group (54.3 \pm 16.8 years). There were more male patients in the medical subgroup (70.4%, $\chi^2 = 17.37$, $P < 0.001$) and the surgical subgroup (69.5%, $\chi^2 = 14.97$, $P < 0.001$) than in the control group (41.6%) (Table 1).

In Step 1, we identified the potential risk factors by comparison with the control group: 1) By comparing the delirium group with the control group, we could find out that ADL score, the BUN, CR, and plasma albumin levels number of medications, past history of delirium and stroke, ICU care, hypertension, mechanical ventilation, infection, immobility, and use of antipsychotic drugs were the potential risk factors of delirium in the delirium group. 2) By comparing the delirium medical subgroup with the control group, we could find out that ADL score, the BUN, CR, plasma albumin, and plasma sodium levels, past history of delirium and stroke, past history of falls, hypertension, mechanical ventilation, infection, and use of antipsychotic drugs were the potential risk factors of delirium in the delirium medical subgroup. 3) By comparing the delirium surgical subgroup with the control group, we could find out that ADL score, the BUN, CR, and plasma albumin levels, number of medications, past history of delirium and stroke, ICU care, hypertension, mechanical ventilation, immobility, and the use of antipsychotic drugs, anticholinergic drugs, and opioids were the potential risk factors of delirium in the delirium surgical subgroup (Tables 2 and 3).

In Step 2, we identified the risk factors for delirium after multiple logistic regression, after controlling for age and sex (results are presented as OR, 95% confidence interval [CI]): 1) In the delirium group, plasma albumin level (0.55, 0.32–0.94), hypertension (2.86, 1.21–6.72), mechanical ventilation (12.49, 1.13–138.00), and use of antipsychotic drugs (4.07, 1.41–11.72); 2) In the delirium medical subgroup, plasma sodium level (0.90, 0.83–0.98) and hypertension (2.91, 1.12–7.55); and 3) In the delirium surgical subgroup, plasma albumin level (0.37, 0.16–0.87), number of medications (1.09, 1.00–1.18), past history of stroke (1,817.71, 5.37–614,920.5), ICU care (17.95, 1.96–164.55), hypertension (5.90, 1.39–25.09), and the use of antipsychotic drugs (14.57, 2.93–72.37), anticholinergic drugs (14.50, 1.57–134.14), and opioids (6.10, 1.58–23.44) (Table 4).

Table 1 Characteristics of the delirium and control groups

Characteristics	Delirium group			Control group	P-value			
	Total	Medical subgroup	Surgical subgroup		Total vs control	Medical subgroup vs control	Surgical subgroup vs control	Medical subgroup vs surgical subgroup
Number of patients, n (%)	260	142 (54.6%)	118 (45.4%)	77	–	–	–	–
Age, years, mean \pm SD	68.9 \pm 13.0	69.3 \pm 13.9	68.4 \pm 11.9	54.3 \pm 16.8	$t=-8.05$; $P<0.001$	$t=-7.06$; $P<0.001$	$t=-6.86$; $P<0.001$	$t=0.52$; $P=0.603$
Male sex, n (%)	182 (70.0%)	100 (70.4%)	82 (69.5%)	32 (41.6%)	$\chi^2=20.7$; $P<0.001$	$\chi^2=17.37$; $P<0.001$	$\chi^2=14.97$; $P<0.001$	$\chi^2=0.027$; $P=0.87$
Psychotic symptoms								
Hallucination, n (%)	91 (35.0%)	47 (33.1%)	44 (37.6%)	–	–	–	–	$\chi^2=0.57$; $P=0.449$
Delusion, n (%)	52 (20.0%)	31 (21.8%)	21 (18.0%)	–	–	–	–	$\chi^2=0.603$; $P=0.438$
CGI-S score, mean \pm SD	3.99 \pm 0.8	4.01 \pm 0.88	3.96 \pm 0.81	–	–	–	–	$t=0.395$; $P=0.693$
Type of delirium								
Hyperactive, n (%)	167 (64.2%)	90 (63.4%)	77 (65.3%)	–	–	–	–	$\chi^2=1.28$; $P=0.526$
Hypoactive, n (%)	18 (6.9%)	8 (5.6%)	10 (8.5%)	–	–	–	–	
Mixed, n (%)	75 (28.8%)	44 (31.0%)	31 (26.3%)	–	–	–	–	

Abbreviations: CGI-S, Clinical Global Impression – Severity scale; SD, standard deviation.

Discussion

In this study, the mean age of the subjects in the delirium group was 14.6 years, which was greater than that of the control group, and 70% of the delirium group comprised males, whereas only 41.6% of the control group were males. This was concordant with previous reports that identified age and male sex as the major risk factors for the development of delirium.^{10,11,34} Age and sex, which are unmodifiable through medical intervention, are demographic risk factors

for delirium. However, this study was designed to identify the significant risk factors for delirium across a wide range of clinical settings and to evaluate the OR of each risk factor. Therefore, we adjusted for age and sex in the logistic regression in Step 2.

Among the psychomotor disturbance subtypes, hyperactive delirium was the most common (64.2%), while hypoactive delirium was the least common (6.9%). This result contrasts with the general opinion that mixed delirium is the

Table 2 Comparison between delirium group and control group in terms of risk factors for delirium (continuous variables)

Risk factors	Delirium group (n=260), mean \pm SD			Control group (n=77), mean \pm SD	Comparisons between delirium group and control group (P-value)		
	Total	Medical subgroup	Surgical subgroup		Total delirium vs control	Medical subgroup vs control	Surgical subgroup vs delirium
Past medical history							
ADL score	3.2 \pm 0.9	3.03 \pm 1.01	3.4 \pm 0.80	3.7 \pm 0.6	0.0001	<0.0001	0.0308
Physical illness							
BUN (mg/dL)	24.1 \pm 15.0	24.8 \pm 15.0	23.2 \pm 15.0	18.3 \pm 13.8	0.0026	0.0018	0.0226
Creatinine (mg/dL)	1.6 \pm 1.9	1.7 \pm 2.0	1.5 \pm 1.7	1.1 \pm 1.1	0.0105	0.0059	0.0410
AST (IU/L)	46.4 \pm 91.6	42.8 \pm 42.4	50.8 \pm 128.0	46.0 \pm 66.3	0.9729	0.6596	0.7639
ALT (IU/L)	42.8 \pm 104.4	40.3 \pm 53.5	45.9 \pm 143.8	42.6 \pm 67.2	0.9871	0.7762	0.8508
Albumin (g/dL)	2.7 \pm 0.6	2.7 \pm 0.6	2.8 \pm 0.7	3.2 \pm 0.7	<0.0001	<0.0001	0.0001
Sodium (mmol/L)	136.7 \pm 10.0	135.5 \pm 12.5	138.1 \pm 5.4	138.5 \pm 4.2	0.1118	0.041	0.5230
Potassium (mmol/L)	4.1 \pm 0.6	4.1 \pm 0.7	4.1 \pm 0.6	4.2 \pm 0.6	0.2631	0.2779	0.3546
Medication history							
Number of drugs	11.5 \pm 6.3	10.4 \pm 4.7	12.8 \pm 7.6	9.5 \pm 4.9	0.0093	0.1599	0.0008

Abbreviations: ADL, activities of daily living; BUN, blood urea nitrogen; AST, aspartate transaminase; ALT, alanine transaminase; SD, standard deviation.

Table 3 Comparison between delirium group and control group in terms of risk factors for delirium (categorical variables)

Risk factors	Delirium group (n=260), n (%)			Control, n (%)	Comparisons between delirium group and control group (P-value)		
	Total	Medical subgroup	Surgical subgroup		Total delirium vs control	Medical subgroup vs control	Surgical subgroup vs control
Past medical history							
Past delirium history	59 (22.7)	26 (18.3)	33 (28.0)	4 (5.2)	0.001	0.007	<0.001
Dementia history	10 (3.9)	5 (3.5)	5 (4.2)	1 (1.3)	0.269	0.336	0.245
Stroke history	26 (10)	10 (7.04)	16 (13.56)	1 (1.3)	0.013	0.063	0.003
Ambulation difficulty	42 (16.2)	24 (16.9)	18 (15.3)	8 (10.4)	0.211	0.193	0.329
Falling history	18 (6.9)	13 (9.15)	5 (4.24)	2 (2.6)	0.158	0.067	0.547
Past neurologic history	39 (15.0)	23 (16.2)	16 (13.6)	11 (14.3)	0.877	0.709	0.886
Hypertension	124 (47.7)	67 (47.2)	57 (48.3)	12 (15.6)	<0.001	<0.001	<0.001
Alcohol use	9 (3.5)	3 (2.1)	6 (5.1)	3 (3.9)	0.857	0.440	0.699
Treatment environment							
ICU care	66 (25.4)	29 (20.4)	37 (31.4)	9 (11.7)	0.011	0.103	0.002
Chemical restraint	76 (29.2)	38 (26.8)	38 (32.2)	0 (0)	<0.001	<0.001	<0.001
Physical restraint	49 (18.9)	21 (14.8)	28 (23.7)	0 (0)	<0.001	<0.001	<0.001
Physical illness							
Surgery	118 (45.4)	0	118 (100)	13 (16.9)			
Mechanical ventilation	34 (13.1)	20 (14.1)	14 (11.9)	1 (1.3)	0.003	0.002	0.007
Infection	122 (46.9)	83 (58.5)	39 (33.1)	18 (23.4)	<0.001	<0.001	0.147
Immobility	92 (35.4)	48 (33.8)	44 (37.3)	8 (10.4)	<0.001	<0.001	<0.001
Medication history							
Antipsychotic drug use	102 (39.2)	50 (35.2)	52 (44.1)	6 (7.8)	<0.001	<0.001	<0.001
Anticholinergic drug use	26 (10.0)	9 (6.3)	17 (14.4)	4 (5.2)	0.193	0.732	0.043
Anticonvulsant use	24 (9.2)	11 (7.8)	13 (11.0)	5 (6.5)	0.452	0.734	0.286
Opioid use	126 (48.5)	59 (41.6)	67 (56.8)	33 (42.9)	0.387	0.851	0.057

Abbreviation: ICU, intensive care unit.

Table 4 Multiple logistic regression of risk factors for delirium

Risk factors	Total delirium vs control			Medical subgroup vs control			Surgical subgroup vs control		
	OR	95% CI	P-value	OR	95% CI	P-value	OR	95% CI	P-value
Past medical history									
Past delirium history	2.90	0.90–11.97	0.142	3.68	0.62–21.70	0.150	7.47	0.87–63.87	0.066
Past history of stroke	8.98	0.48–168.04	0.142	3.19	0.08–121.62	0.533	1,817.71*	5.37–614,920.5	0.012
Past history of falls	–	–	–	0.98	0.14–6.99	0.981	–	–	–
Treatment environment									
ICU care	1.82	0.56–5.92	0.321	–	–	–	17.95*	1.96–164.55	0.011
Hypertension	2.86*	1.21–6.72	0.016	2.91*	1.12–7.55	0.028	5.90*	1.39–25.09	0.016
Mechanical ventilation	12.49*	1.13–138.00	0.039	10.04	0.97–104.21	0.053	6.80	0.29–157.28	0.232
Infection	2.01	0.87–4.63	0.103	2.49	0.98–6.33	0.055	–	–	–
Immobility	2.61	0.90–7.55	0.077	2.92	0.88–9.71	0.080	2.12	0.40–11.15	0.375
Physical illness									
ADL score	0.86	0.52–1.42	0.554	0.67	0.37–1.18	0.165	0.79	0.39–1.61	0.518
BUN	1.00	0.97–1.03	0.997	1.01	0.98–1.04	0.561	1.00	0.94–1.05	0.855
CR	1.15	0.813–1.64	0.422	1.13	0.80–1.61	0.479	0.92	0.53–1.61	0.782
Albumin	0.55*	0.32–0.94	0.028	0.63	0.33–1.21	0.165	0.37*	0.16–0.87	0.022
Sodium	–	–	–	0.90*	0.83–0.98	0.010	–	–	–
Medication history									
Number of medications	1.06	0.99–1.15	0.1	–	–	–	1.09*	1.00–1.18	0.050
Antipsychotic drug use	4.07*	1.41–11.72	0.009	2.38	0.73–7.81	0.151	14.57*	2.93–72.37	0.001
Anticholinergic drug use	–	–	–	–	–	–	14.50*	1.57–134.14	0.018
Opioid use	–	–	–	–	–	–	6.10*	1.58–23.44	0.009

Note: *P<0.05.

Abbreviations: ADL, activities of daily living; BUN, blood urea nitrogen; CI, confidence interval; CR, creatinine; ICU, intensive care unit; OR, odds ratio.

most common delirium subtype.³⁵ This discrepancy may be due to the fact that hypoactive delirium is difficult for medical staff to identify and is easily missed. The natural course of delirium fluctuates, and assessment is often made when the symptoms have become aggravated, leading to a determination of hyperactive delirium. According to a recent study,³⁶ one of the major reasons that nurses fail to detect delirium is that the natural course of delirium changes. Because we included patients who had been referred to psychiatrists, there was a greater chance of including patients whose behavioral problems were relatively severe and easily noticeable.

In order to identify the risk factors in two major clinical situations, we categorized the delirium group into medical and surgical subgroups. Lower albumin level, hypertension, mechanical ventilation, and use of antipsychotic drugs were identified as risk factors in the delirium group. Among these factors, lower albumin level, hypertension, and use of antipsychotic drugs are modifiable through proper medical care, and it might be important to correct these risk factors aggressively to prevent and/or manage delirium. A recent meta-analysis determined that the risk of delirium can be lowered in surgical inpatients by the preventive use of antipsychotic drugs.³⁷ However, the use of antipsychotic drugs did not have a significant effect on the duration of delirium, the severity of symptoms, or the length of hospital stay.³⁷ In contrast, another study reported that the use of antipsychotic drugs in ICU patients without a specific diagnosis might increase the duration of hospital stay and mortality.³⁸ Clearly, the effect of antipsychotic drugs on delirium requires further research.

When we identified the risk factors for delirium in the two subgroups, there were some differences. While the risk factors in the medical subgroup were lower plasma sodium level and hypertension, the risk factors in the surgical subgroup were lower plasma albumin level, past history of stroke, ICU care, hypertension, number of medications, and the use of antipsychotic drugs, anticholinergic drugs, and opioids. The fact that the two subgroups showed different results suggests that different risk factors should be considered in each clinical situation. For nonsurgical patients, it might be important to control electrolyte imbalance. For surgical patients, surgery is a known risk factor for delirium, and delirium is believed to be associated with preoperative and postoperative stroke in patients with risk factors for cardiovascular diseases.^{39–42} Furthermore, postoperative surgical patients are often treated in the ICU, which itself is a risk factor for delirium. Thus, stroke prevention should be a priority in surgical management.

This study had several limitations of note. First, we could not clarify the temporal relation between risk factors and the

occurrence of delirium because of the cross-sectional study design. However, this limitation was minimized by the fact that we collected the risk factor information prior to the onset of delirium. Second, the etiology of delirium is multifactorial and has not been completely clarified. In addition to the risk factors included in this study, there might be unidentified or inconsistently reported risk factors that were not included. Because we could not process the information about medication use appropriately, the effect of medication history on the development of delirium might have been underestimated. Third, the control group and the delirium group were not precisely matched. There were differences in physical illness and major procedures such as surgery under anesthesia. It was difficult to enroll matched controls who had no psychiatric problems and, accordingly, to obtain information from psychiatric assessments for them. To reduce the effect of the control group differences, however, we excluded certain risk factors. For example, we excluded the use of benzodiazepines in the analysis as we were certain that the control group had a higher rate of use. Nevertheless, the results should be cautiously interpreted. Fourth, because the number of patients involved in our study is limited, it is difficult to extrapolate the results to the general population. Moreover, some risk factors (mechanical ventilation and past history of stroke) showed wide CI of the ORs. It may be due to the small number of patients who had risk factors. The results with wide CI should be cautiously interpreted because these results may lose their statistical significance in other studies.

We attempted to overcome the limitations of previous studies by including various risk factors of delirium. We were able to identify significant risk factors under general clinical situations, ie, the study was not limited to one specific clinical setting. Furthermore, we investigated both the delirium group and the control group for the same risk factors. The use of the control group for the analysis is one strength of our study.

Disclosure

The authors report no conflicts of interest in this work.

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