

Relationship between cobalamin deficiency and delirium in elderly patients undergoing cardiac surgery

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Background: Delirium is common after cardiac surgery and is independently associated with increased morbidity, mortality, prolonged hospital stays, and higher costs. Cobalamin (vitamin B₁₂) deficiency is a common cause of neuropsychiatric symptoms and affects up to 40% of elderly people. The relationship between cobalamin deficiency and the occurrence of delirium after cardiac surgery has not been examined in previous studies. We examined the relationship between cobalamin deficiency and delirium in elderly patients undergoing coronary artery bypass grafting (CABG) surgery.

Material and methods: A total of 100 patients with cobalamin deficiency undergoing CABG were enrolled in this retrospective study. Control group comprised 100 patients without cobalamin deficiency undergoing CABG. Patients aged 65 years or over were included. Diagnosis of delirium was made using Intensive Care Delirium Screening Checklist. Delirium severity was measured using the Delirium Rating Scale-revised-98.

Results: Patients with cobalamin deficiency had a significantly higher incidence of delirium (42% vs 26%; $P=0.017$) and higher delirium severity scores (16.5 ± 2.9 vs 15.03 ± 2.48 ; $P=0.034$) than patients without cobalamin deficiency. Cobalamin levels were significantly lower in patients with delirium than patients without delirium ($P=0.004$). Delirium severity score showed a moderate correlation with cobalamin levels ($\rho=-0.27$; $P=0.024$). Logistic regression analysis demonstrated that cobalamin deficiency was independently associated with postoperative delirium (OR 1.93, 95% CI 1.03–3.6, $P=0.038$).

Conclusion: The results of our study suggest that cobalamin deficiency may be associated with increased risk of delirium in patients undergoing CABG. In addition, we found that preoperative cobalamin levels were associated with the severity of delirium. This report highlights the importance of investigation for cobalamin deficiency in patients undergoing cardiac surgery, especially in the elderly.

Keywords: cobalamin deficiency, coronary artery bypass grafting, delirium, delirium severity score, elderly

Background

Delirium is an acute confusional state characterized by disturbance of consciousness with inattention accompanied by a change in cognition or perceptual disturbance that develops over a short period of time (hours to days) and fluctuates over time.¹ Delirium is common in patients undergoing cardiac surgery. Its incidence after cardiac surgery has been reported as 8%–52% and is independently associated with increased morbidity, mortality, prolonged hospital stays, and higher costs.^{2,3} Therefore, recognizing the factors associated with delirium is crucial to prevent and decrease the negative consequences related to delirium.

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Delirium usually has a multifactorial etiology. Previous studies have shown that advanced age, preoperative cognitive impairment, depression, benzodiazepine use, cerebrovascular and peripheral vascular disease, preoperative cerebral oxygen saturation, smoking, atrial fibrillation, renal dysfunction, heart failure, low intraoperative perfusion pressure and cardiogenic shock are associated with increased risk of postoperative delirium.^{4,5}

Cobalamin (vitamin B₁₂) deficiency is characterized by hematologic, psychiatric, and neurologic disturbances and affects up to 40% of elderly people.⁶ The reported prevalence of deficient and marginal values is much higher in developing countries.⁷ Cobalamin deficiency is a common cause of neuropsychiatric symptoms and has been linked to psychiatric disorders such as depression, dementia, catatonia, delirium, schizophrenia, and psychosis.^{8–10} Even occult cobalamin deficiency has been associated with neuropsychiatric disorders.¹¹ Neuropsychiatric symptoms usually precede hematologic signs and are often the presenting manifestation of vitamin cobalamin deficiency.¹¹ Mental or psychological changes may precede hematological signs by months or years.^{11,12}

The relationship between cobalamin deficiency and the occurrence of delirium after cardiac surgery has not been examined in previous studies. The aim of this study was to examine the relationship between cobalamin deficiency and postoperative delirium in elderly patients undergoing coronary artery bypass grafting (CABG) surgery.

Material and methods

Study population

The present study was approved by the Diyarbakir Gazi Yasargil Education and Research Hospital ethics committee (Diyarbakir, Turkey) and complies with the requirements of the Declaration of Helsinki. We retrospectively reviewed the clinical records of patients who underwent CABG with cardiopulmonary bypass (CPB) in our hospital between January 2011 and November 2014 for coronary artery disease. Inclusion criteria were as follows: 1) cobalamin deficiency; 2) measurement of serum B12 levels performed within one month prior to the surgery; 3) age ≥ 65 years; 4) elective, first time CABG with CPB; 5) intensive care unit (ICU) length of stay < 24 hours.

The exclusion criteria were as follows: 1) known psychiatric and/or neurologic diseases before surgery; 2) alcohol dependence; 3) chronic kidney and hepatic diseases; 4) chronic heart failure; 5) reduced myocardial function (ejection fraction $< 45\%$); 6) severe systemic illness; 7) malignant tumors; 8) acute or chronic infections; 9) carotid artery stenosis; 10) folate deficiency; 11) visual and hearing impairment;

12) emergency procedures; 13) redo-CABG; 14) off-pump CABG; 15) preoperative myocardial infarction (MI) within a week; 16) ICU length of stay > 24 hours; 17) bleeding recorded 12 hours after operation $> 1,000$ cc; 18) postoperative MI; 19) perioperative hemodynamic instability; 20) presence of aortic plaque; and 21) perioperative hypoxia and hypercapnia.

Between January 2011 and November 2014, 437 patients underwent CABG in our clinic. A total of 159 patients met inclusion criteria. Of the 159 patients, seven were excluded due to incomplete data, three due to known neurologic disease, one due to alcohol dependence, two due to chronic kidney disease, one due to hepatic disease, four due to chronic heart failure and reduced myocardial function (ejection fraction $< 45\%$), one due to severe systemic illness, one due to malignant tumor, two due to infections, five due to carotid artery stenosis, five due to folate deficiency, one due to visual and hearing impairment, two due to emergency procedures, one due to redo-CABG, two due to off-pump CABG, two due to preoperative MI within a week, five due to ICU length of stay > 24 hours, five due to bleeding $> 1,000$ cc recorded 12 hours after operation, one due to postoperative MI, four due to perioperative hemodynamic instability, one due to presence of aortic plaque, and three due to perioperative hypoxia and hypercapnia.

After applying inclusion and exclusion criteria, 100 patients with cobalamin deficiency were included in the study. Control group comprised 100 patients without cobalamin deficiency. Diagnosis of vitamin B₁₂ deficiency was based on low serum vitamin B₁₂ levels (< 191 ng/L), along with megaloblastic anemia. The lowest hematocrit value on bypass was defined as the nadir hematocrit value throughout the CPB. Type 2 diabetes mellitus (DM) was defined as having a fasting plasma glucose levels ≥ 126 mg/dL or treatment with insulin or an hypoglycemic agent. Hypertension (HT) was defined as systolic blood pressure > 140 mmHg, diastolic blood pressure > 90 mmHg, or use of an antihypertensive medication. Hyperlipidemia (HL) was defined as total serum cholesterol (TC) ≥ 200 mg/dL and/or low-density lipoprotein (LDL) cholesterol ≥ 130 mg/dL and/or triglycerides (TG) ≥ 200 mg/dL and/or history of current use of lipid-lowering regiment.

Anesthesia

The following premedication was administered: 10 mg diazepam, per os, was given to all patients on the night prior to operation, and 0.1 mg/kg morphine HCl was administered intramuscularly half an hour before the operation. Anesthesia was induced with intravenous fentanyl 5–10 μ g/kg, midazolam 0.1 mg/kg, rocuronium bromide 0.6–0.8 mg/kg

and was maintained with fentanyl 1–2 µg/kg/hour, rocuronium bromide 0.3 mg/kg/hour, supported with inhalation of 1%–2% sevoflurane.

Management of CPB

All procedures were performed with median sternotomy. Standard nonpulsatile CPB with a roller pump and a membrane oxygenator was used. The extracorporeal system was primed with ringer lactate. CPB was established between the ascending aorta and right atrium using dual-stage venous cannula. Prior to the cannulation, 300–400 U/kg heparin sulfate was administered to every patient in order to maintain ACT values above 480 seconds. During CPB, nonpulsatile pumpflow was kept at 2.2–2.5 L/m²/minute. Mean arterial pressure during CPB was stabilized between 50 and 70 mmHg. Cold blood cardioplegia was used in all patients. Core temperature was cooled to 28°C. Alpha-stat strategy was used for pH management, and all patients were kept at normocapnic levels (PaCO₂ =35–45 mmHg). Concentrated erythrocyte suspensions were added to pump prime volume, if required to keep the hematocrit level above 20% during CPB. Proximal anastomoses were completed using a partial occlusion clamp. Postoperative analgesia was achieved by administration of paracetamol or tramadol.

Delirium assessment

Diagnosis of delirium was made using Intensive Care Delirium Screening Checklist (ICDSC).¹³ The evaluation is based on eight fields (one point each): altered level of consciousness, inattention, disorientation, hallucination-delusion-psychosis, inappropriate speech or mood, psychomotor agitation/retardation, sleep–wake cycle disturbance, and fluctuating course of aforementioned items. Patients were considered to have delirium if the ICDSC score was equal to or greater than four.

Delirium severity was measured using the Delirium Rating Scale-revised-98 (DRS-R-98).¹⁴ The DRS-R-98 is a 16-item, clinician-rated scale, with anchored item descriptions corresponding to both symptoms and temporal aspects of delirium. The severity scale has 13 items, each rated from 0 to 3, with a maximum score of 39 points, and where higher scores indicate greater severity of delirium. Three additional items (rated from 0 to either 2 or 3) capture temporal course and attribution to an underlying etiology, and, when added to the sum of the 13 symptom-items, produce the DRS-R-98 total score, which ranges from 0 to 46. The original validation study suggested cutoff scores for differential diagnosis of around 17–18 points on the total scale and around 15 for

the severity scale. However, milder forms of delirium can occur with scores ≥ 11 .¹⁴

In our ICU, patients are assessed for delirium by using the Turkish versions of ICDSC and DRS R-98 as a standard of care. In recent studies both ICDSC and DRS-R98 were found to be a valid and reliable tool in determining delirium in Turkish population.^{15,16}

Neuropsychological assessment by DRS R-98 and ICDSC criteria for delirium was made by experienced bedside nurses twice per day, in the morning and in the afternoon, until discharge from the hospital.

Data collection and laboratory tests

Demographic, clinical, and laboratory parameters were collected from the patients' charts. All blood samples were collected before CABG. Serum cobalamin levels were determined by chemiluminescent enzyme immunoassay in an automated analyzer (Roche Diagnostic Modular Analytics E170, Roche Ltd., Basel, Switzerland). The normal range for cobalamin in our laboratory is 191–663 ng/L.

Statistical analysis

Statistical analysis was conducted using SPSS for Windows version 17 (SPSS Inc Chicago, IL, USA). All variables were investigated using visual (histograms, probability plots) and analytic methods (Kolmogorov–Smirnov test) to determine whether or not they are normally distributed. Continuous variables were reported as means and standard deviation for normally distributed variables, and as medians and interquartile range (IQR) for the non-normally distributed variables. Categorical variables were presented using numbers and percentages.

Patients were divided into two subgroups according to the presence or absence of cobalamin deficiency. Comparison between two groups was performed using the chi-square test for qualitative variables, independent *t*-test for normally distributed continuous variables and Mann–Whitney *U*-test for non-normally distributed continuous variables. Patients were further categorized into two subgroups according to the presence or absence of delirium. A similar analysis to that above was made for delirium groups. The strength of linear relations between DRS and clinical variables was assessed by the Pearson or Spearman correlation coefficient as appropriate. Logistic regression analysis was used to evaluate the associations between delirium and cobalamin deficiency. Presence of delirium was used as the dependent variable. Potential risk factors for delirium, and potential predictor variables that had a *P*-value <0.25 in the univariate analysis were identified and included as covariates in the multivariate model. The multivariate model was adjusted

for the following variables: age, sex, DM, COPD, CPB time, and number of transfusions. *P*-values <0.05 were considered to indicate statistical significance.

Results

Relationship between cobalamin deficiency and delirium

Clinical and biochemical characteristics of the patients with cobalamin deficiency (*n*=100; 66 males; mean age 70.4±3.8) and patients without cobalamin deficiency (*n*=100; 62 males; mean age 70.9±4.1) are presented in Table 1.

Patients with cobalamin deficiency had a significantly higher incidence of delirium than those without cobalamin deficiency (42% vs 26%; *P*=0.017). In addition, patients with cobalamin deficiency had significantly higher delirium severity scores than patients without cobalamin deficiency (16.5±2.9 vs 15.03±2.48; *P*=0.034). There were no statistically significant differences in other clinical characteristics between the two groups.

Relationship between delirium and clinical variables

Clinical and biochemical characteristics of the patients with delirium (*n*=68; 41 males; mean age 69.9±4.4) and patients

without delirium (*n*=132; 87 males; mean age 71±3.66) are presented in Table 2. Cobalamin levels were significantly lower in patients with delirium than patients without delirium (*P*=0.004). There were no statistically significant differences in other clinical characteristics between the two groups.

Multivariate analysis

The correlation between delirium rating score and evaluated parameters is presented in Table 3. Delirium score showed a moderate correlation with cobalamin levels ($\rho=-0.27$; *P*=0.024). After adjusting for age, sex, DM, COPD, CPB time, and number of transfusions, logistic regression analysis revealed that cobalamin deficiency was independently associated with postoperative delirium (OR 1.93, 95% CI 1.03–3.6, *P*=0.038).

Discussion

In the present study, we found that, cobalamin deficiency was associated with increased risk of postoperative delirium and higher delirium severity scores in patients undergoing cardiac surgery. Moreover, preoperative cobalamin levels were significantly lower in patients with delirium compared to patients without delirium. As far as we know, this is the first report of

Table 1 Differences in clinical and biochemical characteristics of the patients, according to presence or absence of cobalamin deficiency

	Patients with Cbl deficiency (n=100)	Patients without Cbl deficiency (n=100)	P-value
Age, years, mean ± SD	70.4±3.8	70.9±4.1	0.35
Sex, males, n	66	62	0.33
HT, n	83	76	0.22
DM, n	29	41	0.07
HL, n	57	64	0.31
Smoking, n	63	58	0.47
PVD, n	4	9	0.15
COPD, n	28	41	0.053
EUROSCORE II, mean ± SD	0.98±0.26	1.06±0.3	0.07
Preoperative HCT	34.16±2.8	34.2±2.6	0.32
Anastomoses, median (IQR)	4 (3–4)	3 (2.25–4)	0.36
CPB time, median (IQR)	103.5 (80.25–103.75)	100 (85.25–126.75)	0.93
X clamp time, median (IQR)	65 (51–79.5)	60.5 (48.25–75)	0.26
Nadir HCT on CPB, median (IQR)	20 (17.1–24)	20.2 (18–22.75)	0.9
Operation time, mean ± SD	204.7±29.6	206±31.07	0.75
Ventilator time, mean ± SD	8.3±2.4	8.5±2.5	0.85
RBC transfusion, mean ± SD	2.2±1.3	2.25±0.98	0.91
FFP transfusion, mean ± SD	0.74±0.7	0.74±0.5	0.72
Bleeding recorded 12 hours after operation, median (IQR)	350 (250–450)	300 (200–500)	0.55
Delirium, n	42	26	0.017*
DRS-R-98 severity score, mean ± SD	16.5±2.9	15.03±2.48	0.034*
Preoperative Cbl levels, ng/L, median (IQR)	160 (134.2–176)	352 (286.5–459)	<0.001*

Notes: **P*-values <0.05 were considered to indicate statistical significance.

Abbreviations: Cbl, cobalamin deficiency; HT, hypertension; DM, diabetes mellitus; HL, hyperlipidemia; PVD, peripheral vascular disease; COPD, chronic obstructive pulmonary disease; CPB, cardiopulmonary bypass; X clamp, cross clamp; HCT, hematocrit value; RBC, red blood cell; FFP, fresh frozen plasma; DRS-R-98, Delirium Rating Scale-Revised-98; SD, standard deviation; IQR, interquartile range.

Table 2 Differences in clinical and biochemical characteristics of the patients, according to presence or absence of delirium

	Patients without delirium (n=132)	Patients with delirium (n=68)	P-value
Age, years, mean $\pm$ SD	71 $\pm$ 3.66	69.9 $\pm$ 4.4	0.052
Sex, males, n	87 (65.9)	41 (60.3)	0.43
HT, n	106 (80.3)	53 (77.9)	0.69
DM, n	45 (34.1)	25 (36.8)	0.7
HL, n	76 (57.6)	45 (66.2)	0.23
Smoking, n	83 (62.9)	38 (55.9)	0.33
PVD, n	9 (6.8)	4 (5.9)	0.8
COPD, n	51 (38.6)	18 (26.5)	0.08
EUROSCORE II, mean $\pm$ SD	1.02 $\pm$ 0.27	1.03 $\pm$ 0.31	0.9
Preoperative HCT	34.14 $\pm$ 2.7	34.3 $\pm$ 2.8	0.22
Anastomoses, median (IQR)	3 (3–4)	4 (3–4)	0.75
CPB time, median (IQR)	101.5 (82–124)	107 (77.2–125)	0.97
X clamp time, median (IQR)	62 (51–78)	65 (47.2–77.2)	0.78
Nadir HCT on CPB, median (IQR)	20 (18–23)	21 (18–24)	0.09
Operation time, mean $\pm$ SD	204.7 $\pm$ 30.1	206.8 $\pm$ 30.7	0.64
Ventilator time, mean $\pm$ SD	8.3 $\pm$ 2.5	8.66 $\pm$ 2.4	0.29
RBC transfusions, mean $\pm$ SD	2.3 $\pm$ 1.1	2.04 $\pm$ 1.22	0.13
FFP transfusion, mean $\pm$ SD	0.77 $\pm$ 0.6	0.68 $\pm$ 0.67	0.3
Bleeding recorded 12 hours after operation	350 (250–500)	300 (200–400)	0.08
Cbl deficiency, n	58 (43.9)	42 (61.8)	0.017*
Preoperative Cbl levels, ng/L, median (IQR)	254 (165–388)	184.5 (142.5–288.2)	0.004*

Notes: *P-values <0.05 were considered to indicate statistical significance.

Abbreviations: HT, hypertension; DM, diabetes mellitus; HL, hyperlipidemia; PVD, peripheral vascular disease; COPD, chronic obstructive pulmonary disease; CPB, cardiopulmonary bypass; X clamp, cross clamp; HCT, hematocrit value; RBC, red blood cell; FFP, fresh frozen plasma; Cbl, cobalamin; SD, standard deviation; IQR, interquartile range.

an association between cobalamin deficiency and postoperative delirium in elderly patients undergoing CABG surgery.

The underlying cause of delirium is often multifactorial, and the pathogenesis of delirium is not entirely understood. Alterations in the levels of neurotransmitters and the neurotransmitter imbalances between the monoaminergic (dopamine, serotonin, norepinephrine) and cholinergic neurotransmitter systems have been proposed to be a leading

pathobiological mechanism for delirium.^{17,18} Monoamine neurotransmitter system, including the neurotransmitters dopamine, serotonin, and norepinephrine, has a balancing role for the cholinergic activity. Thus, the development of delirium involves interaction between these two neurotransmitter systems, and any process interfering with neurotransmitter function or with the supply or use of substrates can cause delirium.¹⁷ Other possible mechanisms are neural injury, inflammation, and stress response.¹⁷ Variable degrees of neurologic injury, ranging from relatively minor neurocognitive deficits to more severe cerebral infarction may occur following cardiac surgery.¹⁹ Cerebral hypoxia and hypoperfusion and embolization have been implicated in CPB-related cerebral injury.¹⁹ Inflammation and increase in TNF- α , interleukin-10, and soluble TNF-receptor I levels after CPB were also demonstrated to be associated with brain injury in patients undergoing cardiac surgery.²⁰

Although the exact mechanism of how vitamin B₁₂ deficiency causes neuropsychiatric disorder is unclear, several mechanisms detailed can contribute to delirium in cobalamin-deficient patients undergoing CABG: 1) impaired monoamine neurotransmitter metabolism, 2) derangements in DNA synthesis, 3) vasculotoxic effects associated with elevated Hcy and demyelination associated with elevated methylmalonic acid levels, 4) diminished protection against inflammatory oxidative stress

Table 3 Correlation between the delirium severity score and clinical variables

	ρ	r	P
Age	–	–0.04	0.7
EUROSCORE II	–	0.23	0.06
Cbl level	–0.27	–	0.024*
Number of anastomosis	–0.19	–	0.11
CPB time	–0.05	–	0.69
X clamp time	–0.09	–	0.44
Nadir HCT on CPB	–0.1	–	0.4
Operation time	–	–0.13	0.29
Ventilator time	–	0.18	0.13
RBC transfusion	–	0.07	0.57
FFP transfusion	–	0.13	0.27
Bleeding recorded 12 hours after operation	–0.004	–	0.94

Notes: *P-values <0.05 were considered to indicate statistical significance.

Abbreviations: Cbl, cobalamin; CPB, cardiopulmonary bypass; X clamp, cross clamp; HCT, hematocrit; RBC, red blood cell; FFP, fresh frozen plasma.

in the perioperative period, and 5) mild cognitive impairment (MCI) and resultant increase in plasma IL-2 and cortisol levels. Cobalamin is critical to central nerve system functioning and is required for the proper biosynthesis of the monoamine neurotransmitters serotonin, epinephrine, and dopamine.²¹ Thus, cobalamin deficiency can lead to decreased production of monoamine neurotransmitters.²¹ A potential mechanism for delirium in patients with cobalamin deficiency may be neurotransmitter imbalances between monoaminergic and cholinergic neurotransmission. Cobalamin deficiency leads to elevated homocysteine and methylmalonic acid levels.²¹ Methylmalonic acid is a myelin destabiliser and can affect normal myelin formation.²² Homocysteine has been shown to be neurotoxic, leading to DNA damage and apoptosis.²³ Previous studies suggested that elevated homocysteine may induce oxidative stress through activation of glutamate receptors and consequent reactive species generation, or by autooxidation to homocysteine and other disulfides releasing O₂ and H₂O₂.^{24,25} These findings suggest that, neurotoxicity of homocysteine and methylmalonic acid, which is further augmented by CPB, may provide another mechanism for delirium in patients with cobalamin deficiency. Recent observations suggest that cobalamins exhibit a marked antioxidant activity and provide significant cellular protection against oxidative stress, including those of the inflammatory response.²⁶ Cobalamin modulates the immune response and influences cytokine and growth factor production.²⁷ Thus, it was suggested that cobalamins could be used to treat a number of pathological conditions in which inflammatory oxidative stress is a clinically important component.^{26,28} Diminished protection against inflammatory oxidative stress in the perioperative period may be another possible mechanism responsible for delirium in patients with cobalamin deficiency. Vitamin B₁₂ deficiency has been linked to MCI in previous studies. MCI has been associated with risk for delirium after cardiac surgery. Patients with MCI were found to have higher cortisol and IL-2 concentrations when compared to non-MCI subjects. Thus, it was suggested that cortisol and inflammatory alterations may underlie the susceptibility of these patients for delirium.^{29,30}

Vitamin B₁₂ deficiency prevalence in the United States has been reported to range from 3% to 6%.³¹ The reported prevalence of deficient and marginal values is much higher in developing countries. It was shown that 40% of individuals in Latin America had deficient or marginal levels of serum vitamin B₁₂.³² Even higher Vitamin B₁₂ deficiency prevalence rates, up to 80%, have been reported in African and Asian countries.^{33–36} In a previous study which was conducted in our country, the prevalence of vitamin B₁₂ deficiency has been reported to range from 50% to 73.5%.⁷ The prevalence

of Vitamin B₁₂ deficiency in our study population was found to be 36.3%.

Prompt treatment of B₁₂-deficient patients is essential to avoid irreversible neurological and cognitive impairment. Improvement of delirium, depression, and psychotic symptoms after B₁₂ replacement therapy has been reported in previous reports in patients with B₁₂ deficiency.^{37–39} The management of vitamin B₁₂ deficiency varies between countries and between individual practices. Vitamin B₁₂ treatment should start preoperatively and continue postoperatively. Options available for treating a clinical vitamin B₁₂ deficiency include high dose oral and parenteral (intramuscular or subcutaneous) preparations. Classical treatment for vitamin B₁₂ deficiency consists of parenteral administration.⁶ To date, no studies have examined whether B₁₂ replacement therapy reduces the delirium incidence in patients with B₁₂ deficiency undergoing cardiac surgery. Further studies are needed to examine whether vitamin B₁₂ replacement therapy reduces the incidence of delirium in patients with vitamin B₁₂ deficiency after cardiac surgery. Nevertheless, the Food and Nutrition Board recommends that even healthy older adults should take a daily vitamin B₁₂ supplement.⁴⁰

This study has several limitations. The major limitation of our study is its retrospective design. Second, postoperative cobalamin levels were not measured. Some of the potential risk factors such as renal deficiency, cognitive dysfunction, and brain chronic ischemic changes were not included in the analysis. This is another limitation of our study. Finally, this study was conducted in patients undergoing CABG surgery. Whether patients with cobalamin deficiency undergoing other types of cardiac surgery are at increased risk for delirium warrants further investigation.

Conclusion

The results of our study suggest that cobalamin deficiency may be associated with increased risk of delirium in patients undergoing CABG. In addition, we found that preoperative cobalamin level was associated with the severity of delirium. This report highlights the importance of investigation for cobalamin deficiency in patients undergoing cardiac surgery, especially in the elderly.

Disclosure

The authors report no conflicts of interest in this work.

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