REVIEW

Incidence of Postoperative Cognitive Dysfunction Following Inhalational vs Total Intravenous General Anesthesia: A Systematic Review and Meta-Analysis

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Abstract: Postoperative cognitive dysfunction (POCD) has been increasingly recognized as a contributor to postoperative complications. A consensus-working group recommended that POCD should be distinguished between delayed cognitive recovery, ie, evaluations up to 30 days postoperative, and neurocognitive disorder, ie, assessments performed between 30 days and 12 months after surgery. Additionally, the choice of the anesthetic, either inhalational or total intravenous anesthesia (TIVA) and its effect on the incidence of POCD, has become a focus of research. Our primary objective was to search the literature and conduct a meta-analysis to verify whether the choice of general anesthesia may impact the incidence of POCD in the first 30 days postoperatively. As a secondary objective, a systematic review of the literature was conducted to estimate the effects of the anesthetic on POCD between 30 days and 12 months postoperative. For the primary objective, an initial review of 1913 articles yielded ten studies with a total of 3390 individuals. For the secondary objective, four studies with a total of 480 patients were selected. In the first 30 days postoperative, the odds-ratio for POCD in TIVA group was 0.46 (95% CI = 0.26–0.81; p = 0.01), compared to the inhalational group. TIVA was associated with a lower incidence of POCD in the first 30 days postoperatively. Regarding the secondary objective, due to the small number of selected articles and its high heterogeneity, a metanalysis was not conducted. Given the heterogeneity of criteria for POCD, future prospective studies with more robust designs should be performed to fully address this question.

Keywords: postoperative cognitive dysfunction, POCD, total intravenous anesthesia, TIVA, inhalational anesthesia, postoperative complications, psychometric tests

Introduction

Postoperative cognitive dysfunction (POCD) is a common condition after surgery and anesthesia.^{1,2} Recent studies showed an incidence of POCD between 10% and 18%.^{3–7} The International Study of Post-Operative Cognitive Dysfunction (ISPOCD-1) has estimated the incidence of POCD after non-cardiac surgery is as high as 9.9% at three months.⁸

Regarding the choice of the type of general anesthesia, previous studies have identified a possible role of propofol in attenuating the inflammatory cascade.^{9,10} Moreover, an increase in various cytokines, including IL-6, TNF- α , IL-8, and IL-10, have been found to be associated with the presence of POCD.^{11,12} Consequentially, TIVA may be hypothesized as being protective against POCD.

A consensus-working group published recommendations from a panel of specialists suggesting that cognitive assessments on POCD should be distinguished into delayed cognitive recovery (DCR), ie, evaluations up to 30 days postoperative, and postoperative neurocognitive disorder (pNCD), ie, assessments performed between 30 days and 12 months after surgery. The consensus-working group stressed that cognitive decline after the first 30 days postoperatively might potentially be linked to long-term consequences and should, therefore, also be a topic for research.¹³

The primary objective of this study was to conduct a systematic review of the literature and a meta-analysis on the clinical impact of the choice of general anesthesia on the incidence of POCD-DCR, either inhalational or total intravenous anesthesia (TIVA) in the first 30 days, excluding assessments on the same day of surgery. As a secondary goal, we conducted a systematic review of the literature to study the impact of the choice of anesthetic on the incidence of POCD-pNCD between 30 days and 12 months postoperatively.

Methods

Search Strategy

The 2020 updated Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines were followed when performing and reporting this study.¹⁴ A health sciences librarian (BH) conducted an initial literature search on March 6, 2020, and an updated search on May 24, 2021. The following databases were queried: Ovid MEDLINE(R); Embase.com; Web of Science, Google Scholar. Conference abstracts/papers were excluded in Embase. No other limits were applied.

All retrieved records were organized using the citation management software Endnote version 20 (Clarivate, London, U.K.). For removal of duplicates Covidence (Melbourne, Australia), a systematic review citation reviewing and screening software, was used.

The search strategy was designed to capture the association between post-operative cognitive dysfunction (POCD) with surgical anesthetics, specifically propofol and inhalational agents. The full search strategy is presented in Figure 1.

Searches were supplemented by hand searching and retrieval of any additional articles meeting eligibility criteria that were cited in our reference lists.

The full protocol for this systematic review and meta-analysis is registered and approved at the PROSPERO database under the registration number CRD42021239283.

Study Selection

Only randomized controlled trials (RCT) comparing the impact on POCD between TIVA and inhalational anesthesia were selected. All papers including cardiac, carotid, or neurosurgical procedures, and non-adult patients were excluded. Studies that only assessed cognitive function on the same day of surgery were also excluded. If the title and/or abstract suggested that a paper matched the inclusion and exclusion criteria, the full article was screened and assessed for eligibility.

Primary Objective

For our primary objective, we considered POCD-DCR assessed in the first 30 days postoperatively. When assessments were performed multiple times in the postoperative period, we selected the first measurement after surgery, excluding assessments on the same day of surgery.

Secondary Objective

For our secondary aim, we focused only on papers evaluating POCD-pNCD between 30 days and 12 months postoperatively.

Methodological Quality and Risk of Bias Analysis

The methodological quality of the included studies for both objectives was assessed using the Cochrane Collaboration's tool for assessing risk of bias in randomized trials,¹⁵ which accounts for six potential risks of bias: selection, performance, detection, attrition, reporting, and other sources of bias. Ultimately, each domain was assessed as low, high, or unclear.

Database(s	s) [,] Ovid	MEDLINE	(R) ALL	1946 to May	/ 24 2021

#	Searches	Results
1	(((Postoperative Cognit* or post operative Cognit* or post surg* cognit*) adj3 (Complic* or Dysfunction* or Decline or impair* or disorder*)) or POCD).tw,kf. or Postoperative Cognitive Complications/ or ((exp Postoperative Period/ or exp Postoperative Complications/) and exp Cognition Disorders/)	3081
2	(an?est* or propofol or Disoprofol or Diprivan or Disoprivan or Fresofol or Ivofol or Recofol or Aquafol or anepol or anesia or cryotol or diprofol or disoprivan or fresofol or gobbifol or hiremon or plofed or pofol or profast or propocam or propolipid or propoven or provive or rapinovet or rapiva or ripol or safol or spifol or spiva or unifol or sevoflurane or Sevorane or Ultane or sevocalm or sevoflo or sevofrane or sevohale or sevotec or sojourn or ultane).tw,kf. or exp Anesthesia/ or exp Propofol/ or exp Sevoflurane/	532390
3	1 and 2	1181
4	remove duplicates from 3	1178

#	Query	Results
ŧ5	#3 NOT #4	1464
4	#1 AND #2 AND ([conference abstract]/lim OR [conference paper]/lim OR [conference review]/lim)	335
#3	#1 AND #2	1799
#2	an?est*:ti,ab OR propofol:ti,ab OR disoprofol:ti,ab OR diprivan:ti,ab OR ivofol:ti,ab OR recofol:ti,ab OR aquafol:ti,ab OR anepol:ti,ab OR anesia:ti,ab OR cryotol:ti,ab OR diprofol:ti,ab OR disoprivan:ti,ab OR fresofol:ti,ab OR gobbifol:ti,ab OR hiremon:ti,ab OR plofed:ti,ab OR profol:ti,ab OR profast:ti,ab OR propocam:ti,ab OR propolipid:ti,ab OR propoven:ti,ab OR provive:ti,ab OR rapinove:ti,ab OR rapiva:ti,ab OR ripofit:,ab OR sofol:ti,ab OR spiva:ti,ab OR rapiva:ti,ab OR propoven:ti,ab OR sevorane:ti,ab OR sevoralm:ti,ab OR sevoflo:ti,ab OR sevoflurane:ti,ab OR sevorane:ti,ab OR sevote:ti,ab OR sevoflo:ti,ab OR ultane:ti,ab OR sevorane:ti,ab OR sevote:ti,ab OR sofol:ti,ab OR ultane:ti,ab OR anesthesia/exp OR 'propofol/exp OR 'sevoflurane'/exp	557240
1	((('postoperative cognit*' OR 'post operative cognit*' OR 'post surg* cognit*') NEAR/3 (complic* OR dysfunction* OR decline OR impair* OR disorder*)):ti,ab) OR pocd:ti,ab OR 'postoperative cognitive dysfunction'/exp OR (('postoperative complication'/exp OR 'postoperative period'/exp) AND ('cognitive defect'/de OR 'mild cognitive impairment'/de))	6456

Web of	Web of Science (Indexes=SCI-EXPANDED, SSCI Timespan=All years) UPDATE: 5/24/21								
#	Searches	Results							
# 3	#1 AND #2	449							
# 2	TS=(an?est* or propofol or Disoprofol or Diprivan or Disoprivan or Fresofol or Ivofol or Recofol or Aquafol or anepol or anesia or cryotol or diprofol or disoprivan or fresofol or gobbifol or hiremon or plofed or pofol or profast or propocam or propolipid or propoven or provive or rapinovet or rapiva or ripol or safol or spifol or spiva or unifol or sevoflurane or Sevorane or Ultane or sevocalm or sevoflo or sevofrane or sevohale or sevotec or sojourn or ultane)	184,966							
# 1	TS=(("Postoperative Cognit*" or "post operative Cognit*" or "post surg* cognit*") near/3 (Complic* or Dysfunction* or Decline or impair* or disorder*) or POCD)	2,002							

Google Scholar Date: 5/24/21	postoperative cognitive dysfunction anesthesia propofol sevoflurane

Figure I The complete search strategy.

Two investigators (DN and YAW) independently selected the studies, extracted the relevant information from the included trials, and assessed the risk of bias. In the case of disagreement, a third investigator (AO) resolved the conflict. Studies judged with an unknown or high risk of bias were excluded.

Outcomes

The outcome for the primary objective was the incidence of POCD-DCR, as described by the authors of the primary studies, in patients exposed to either TIVA or inhalational anesthesia in the first 30 days postoperatively. For the primary objective, we estimated the odds ratio of POCD between the two groups. For the secondary objective, the outcome was POCD – pNCD, also as defined the authors of the primary studies.

Data Synthesis and Statistical Analysis

All analyses were performed using Stata version 15.1 (StataCorp LLC, College Station, Texas, USA). The percentage of the total variability in the set of effect sizes due to true heterogeneity was tested with the I^2 statistic. Random Effects Mantel-Haenszel model was used to estimate adjusted odds ratio and 95% confidence intervals for the pooled data for the primary objective.

Assessment of the Quality of Evidence

The quality of evidence was evaluated using the GRADE system, which accounts for four different levels of evidence, ie, high, moderate, low, and very low.¹⁶

Results

The flowchart for data extraction is shown in Figure 2. We identified 3381 total citations. After removal of duplicates, 1913 total unique citations were selected. After screening for eligibility criteria based on the title and/or the abstract, 17 potentially eligible articles were retrieved in full text, with 2 additional studies being further included after hand searching on our reference lists. Ten published studies were then selected for the primary objective. Four studies were selected for the secondary objective.

Primary Objective

Overall, ten RCTs compared TIVA vs inhalational anesthesia reporting the incidence of POCD-DCR in both groups (Table 1). The mean sample size among those studies was 339 subjects, considering both groups. In total, 1660 participants were assigned to the TIVA group and 1730 to inhalational anesthesia. The range of age among all the ten studies varied from 18 to 86 years, with median age of 70 years. In the TIVA group, the median age was 70.58 years (18–86) whereas in the inhalational group, the median age was 69.43 years (24–85). The pooled incidence of POCD in the TIVA group was 11.4%, while in the Inhalational group was 27.7%.

Nine out of the ten studies seemed to favor TIVA. Moreover, three of those nine studies reached statistical significance.^{7,17,18} None of the included studies favored inhalational anesthesia. The pooled OR significantly favored the use of TIVA 0.46 (95% CI = 0.26–0.81; p = 0.01) (Figure 3). Additionally, since one of the included studies contributed with nearly half of the total amount of subjects included in the pooled analysis,¹⁷ we conducted a separate analysis, excluding this study, to evaluate the overall effect measure this paper could potentially have on the pooled odds ratio. Our results indicated that we still found a significant association favoring TIVA (0.67; 95% CI = 0.51–0.87; p < 0.01), even after excluding this paper (Figure 4).

The Random Effects Mantel-Haenszel model was used to estimate adjusted OR and 95% confidence intervals for the pooled data based on the value of the I^2 statistic, which was judged as high. We assumed a cut-off value of 75% for I^2 statistic to choose between models.

Two studies used the Mini-Mental state examination (MMSE) as the only tool for evaluation.^{18,19} Specific tests used by different authors in their respective studies are summarized in Table 1. When more than one assessment of cognitive performance was conducted in the postoperative period, we chose the measurement closest to the day of surgery. Among studies with multiple testing in the first 30 postoperative days,^{6,17,18} results were similar in all assessments, apart from one study,¹⁷ in which cognitive decline was observed in inhalational group, compared to propofol, only in postoperative days one, two and three, but not in day ten. Consequently, we used data from 1st day after surgery from four studies,^{4,18,19} day 2 from three studies,^{6,7,20} day 7 from four studies^{3,5,21} and day 10 in a single study.²²

The definition used to diagnose POCD-DCR varied a lot among those studies, ranging from a statistical difference in the means between pre- and postoperative values in MMSE, up to more sophisticated concepts, such as the Z-score or more than one SD in at least two different tests evaluating different cognitive domains, between pre- and postoperative values. Four studies used health controls not submitted to any surgery or anesthesia in the comparison.^{3,5,19,21} All included studies used monitoring of the level of consciousness, except for one.¹⁹

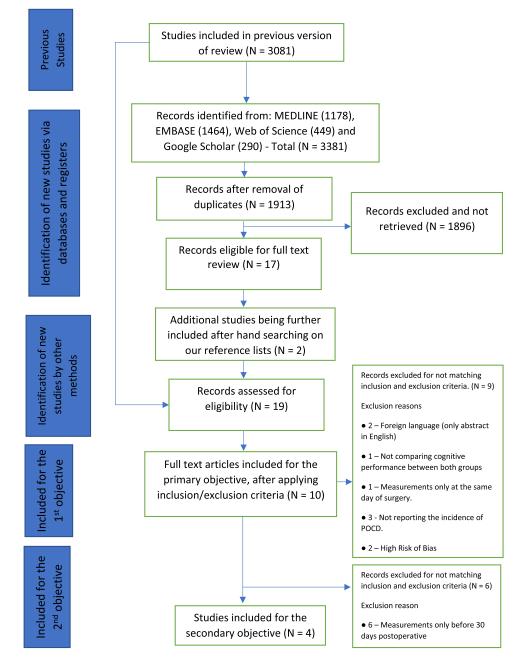


Figure 2 PRISMA[®] flow chart showing article selection process.

Notes: Adapted from: Page MJ, Moher D, Bossuyt PM, et al. PRISMA 2020 explanation and elaboration: updated guidance and exemplars for reporting systematic reviews. *BMJ*. 2021;372. doi:https://doi.org/10.1136/bmj.n160.¹⁴ Copyright © 2021, BMJ Publishing Group Ltd. Creative Commons CC BY 4.0 license (<u>https://creativecommons.org/</u> licenses/by/4.0/legalcode).

In all included studies, the type of opioids and regimen of administration were similar between groups. Most studies used sevoflurane as inhalational agent. Two studies used isoflurane,^{17,18} and one used desflurane.²⁰ In most studies, usual clinical doses of inhalational agents were always used and titrated to maintain an adequate level of consciousness based on the BIS monitor. In particular, the one study that admitted in the inhalational group doses of Isoflurane ranging two to three percent, the BIS monitor was also used to titrate the adequate level of consciousness.¹⁷

The overall risk of bias in the included studies for the meta-analysis (Figure 5) was considered low in all the ten included studies. As mentioned earlier, studies assessed with an unknown or high risk of bias were excluded. Two studies were excluded during the extraction phase based on these criteria.^{3,23} We have also estimated the impact of the use of

Author	Year	Study Design	Type of Surgery	Age	Number of Patients	Tests Used	Tests Altered	Control Group Beyond Pre-Op Testing	Measure of Significance for Diagnosing POCD	BIS Used in Both Groups	Moment of Cognitive Assessment
Rohan et al ²⁰	2005	RCT	Cystoscopy or hysteroscopy	65–86	Total = 30 (Propofol = 15 vs Sevo = 15)	Stroop test and the modified word-recall test	Both	Yes	Z-score analysis	No	l day post-op
Cai et al ¹⁷	2011	RCT	Esophageal cancer, gastric cancer, renal carcinoma, and fracture	>60	Total = 2000 (Propofol = 1000 vs Inhalational = 1000)	MMSE	MMSE	No	MMSE < 25	Yes	I, 3 and 10 days post-op
Tang et al ²²	2014	RCT	Thoracic surgery	>60	Total = 210 (Propofol = 101 vs Inhalational = 99)	TMT (trail-making test); AVLT (auditory verbal learning test); DSF (digit span forward); DSB (digit span backward); DSST (digit symbol substitution test); SCWT (Stroop colour and word test)	Not clear	Yes	Cognitive decline	Yes	7 days post- op
Egawa et at ⁶	2016	RCT	Lung surgery	63–73	Total = 144 (Propofol = 72 vs Inhalational = 72)	MMSE; trail making test A and B; digit span forward and backward; grooved pegboard test dominant hand and non-dominant hand	Not clear	No	Decline ≥ 20% in preop test score in at least two different tests.	Yes	2 and 5 days, and 3 months
Micha et al ⁷	2016	RCT	Tumor resection (non-neural), longer than 2 hours	60–74	Total = 73 (Propofol = 36 vs Inhalational = 37)	Rey auditory verbal learning trail making (A and B), Stroop neuropsychological screening, COWA, three word-three shapes Babcock story recall clock test Beck depression inventory (BDI), instrumental activities daily living (IADLS)	COWA; Stroop neuropsychological screening; clock test; three word-three shapes; Babcock story recall; instrumental activities daily living (IADLS); trail making-B	No	Statistically difference in means in any of the tests	Yes	2 days and 9 months

Table I Characteristics of Included Articles in the Review and Meta-Analysis

Geng et al ¹⁸	2017	RCT	Laparoscopic cholecystectomy	>65	Total = 150 (Propofol = 50 vs Inhalational = 100)	(MMSE); a vision test; the digit symbol substitution test the cumulative test; digit span: forward and backward; the trail making test part A; the Rey auditory verbal learning test (RAVLT): and the grooved pegboard test (dominant and non dominant hands)	Not clear	No	Decline ≥ 20% in preop test score in at least two different tests.	Yes	I and 3 days post-op
Tanaka et al ²¹	2017	RCT	Total knee arthroplasty	>65	Total = 90 (Propofol = 45 vs Inhalational = 45)	Digit symbol substitution test (DDST) [23], the mini mental status exam (MMSE) [24], the trail making test, the digit span subtest (DST) [25] of the Wechsler adult intelligence scale (revised), and the geriatric depression scale (GDS)	All tests	No	Decline ≥ 20% in preop test score in any test.	Yes	2 days post- op
Kletecka et al ⁴	2018	RCT	Single space lumbar disc hernia resection	18–65	Total = 43 (Propofol = 20 vs Inhalational = 23)	Digital span test (forward and backward), letter number sequence test, verbal fluency, trial making test (A and B), Stroop test	Semantic verbal fluency (executive function); letter number sequence test (speed and visual space working memory)	Not clear	I SD in at least three tests	Yes	6 and 42 days
Zhang et al ¹⁹	2018	RCT	Major cancer surgery (> 2H)	65–90	Total = 379 (Propofol = 189 vs Inhalational = 190)	Mental control, visual retention, paired associate verbal learning, digit span (forward and backward), digit symbol, trail making (part A) and grooved pegboard (favored and non-favored hand)	Not reported	Yes	Z-score analysis	Yes	7 days post- op.
Guo et al ⁵	2019	RCT	Tumor resection (Both abdominal and thoracic)	>than 65	Total = 220 (Propofol = 110 vs Inhalational = 110)	Verbal learning test (learning trial and delay); concept shifting task (part C); Stroop color word test (part 3); letter digit coding	All tests	Yes	Z-score altered in any two tests	Yes	7 days and 3 months

Notes: RCT (randomized controlled trial); Z-score analysis (Z = Z = [X-X reference]/standard deviation (SD); where X is the difference between baseline and postoperative cognitive test scores in the trial group, X reference is the difference between baseline and final time point in the control group, and SD is the change in score for the control group. When Z-score is ≥ 1.96 it's considered positive.

Abbreviations: MMSE, mini-mental state examination; CERAD, consortium to establish a registry for Alzheimer's disease; AVLT, auditory-verbal learning (AVLT); TMT A and B, trail making test A and B; DSST, digit symbol substitution test; COWAT, controlled oral word association test; GPB d and nd, grooved pegboard test dominant and non-dominant hand.

	TI	VA	Inhala	ational	Odds Ratio	Weig
Study	Yes	No	Yes	No	with 95% Cl	(%
Egawa	16	56	24	48	0.57 [0.27, 1.20]	11.5
Guo L	30	80	32	78		12.5
Kletecka	4	16	4	19	1.19 [0.26, 5.52]	6.9
Micha	1	35	10	27	0.08 [0.01, 0.64]	4.73
Geng	2	48	25	75	0.13 [0.03, 0.55]	7.18
Rohan	7	8	7	8	1.00 [0.24, 4.20]	7.44
Tanaka	19	20	26	14	0.51 [0.21, 1.26]	10.53
Tang	30	71	33	66		12.44
Cai Y	44	956	275	725	0.12 [0.09, 0.17]	13.78
Zhang Y	28	189	44	190		12.9
Overall	181	1479	480	1250	0.46 [0.26, 0.81]	
Heteroger	neity:		² :	= 81.33%		
Test of θ =	= 0: z =	= -2.71	l, p =	0.01		
Random-effects REML model					1/64 1/16 1/4 1 4 Favours TIVA Favours Inhalational	

Figure 3 Forest plot presented OR, for each of the ten studies included for primary objective, and pooled data (TIVA vs Inhalational).

	TIVA		TIVA Inhalational		ational	Odds Ratio	Weight
Study	Yes	No	Yes	No	with 95% Cl	(%)	
Egawa	16	56	24	48	0.57 [0.27, 1.20]	12.92	
Guo L	30	80	32	78		20.53	
Kletecka	4	16	4	19	— 1.19 [0.26, 5.52]	3.00	
Micha	1	35	10	27	0.08 [0.01, 0.64]	1.58	
Geng	2	48	25	75	0.13 [0.03, 0.55]	3.21	
Rohan	7	8	7	8	—— 1.00 [0.24, 4.20]	3.45	
Tanaka	19	20	26	14	0.51 [0.21, 1.26]	8.68	
Tang	30	71	33	66		19.87	
Zhang Y	28	189	44	190	0.64 [0.38, 1.07]	26.75	
Overall	137	523	205	525	• 0.67 [0.51, 0.87]		
Heteroger	Heterogeneity: $I^2 = 0.00\%$		0.00%				
Test of θ = 0: z = -2.97, p < 0.01					1/64 1/16 1/4 1 4		
Random-effects REML model					Favours TIVA Favours Inhalational		

Figure 4 Forest plot presented OR, for each of the nine studies included for primary objective, excluding "Cai et al", and pooled data (TIVA vs Inhalational).

TIVA and POCD including those two studies, and still found an association between TIVA and a lower incidence of POCD, with OR = 0.60 (95% CI = 0.40-0.91; p = 0.02).

The quality of the evidence with found based on the GRADE system,¹⁶ estimating the association between the use of TIVA and a lower incidence of POCD, was judged as moderate (Figure 6).

Secondary Objective

Characteristics of eligible studies for the secondary objective are shown in Table 1. From the four included articles (4–7), a total of 480 patients were included, 238 in the TIVA group, with a median age of 68.15 years (20–85), and 242 in the inhalational group, with a median age of 67.83 years (24–81). The range of age of the participants enrolled in all four studies varied from 18 to 85 years.



Figure 5 Cochrane collaboration's tool for assessing risk of bias for the included studies.

Quality of evidence and definitions - GRADE System

High quality — Further research is very unlikely to change our confidence in the estimate of effect Moderate quality — Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate Low quality — Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate Very low quality — Any estimate of effect is very uncertain

Figure 6 Application of the GRADE system to our findings. We believe our evidence should be classified as moderate.

As previously mentioned, the type of opioids used, and the regimen of administration were similar in both groups for all included studies. All included studies for the secondary objective used sevoflurane as inhalational agent.

The overall risk of bias for the included studies (Figure 5) indicated that the risk was low in all the four included studies. One study was excluded based on the risk of bias, judged as unknown, during the extraction phase.³

In the study by Kletecka et al,⁴ postoperative measurements were conducted at 42 days postoperatively, but authors only considered a diagnosis of POCD-pNCD if three of the following tests were altered: the Digital Span Test (Forward and Backward), the Letter Number Sequence Test, the Verbal Fluency, the Trail Making Test (TMT), A and B, and the Stroop Test. Egawa et al⁶ conducted another study using a definition of POCD-pNCD when there was a difference in

means of test scores of at least 20% between postoperative and preoperative scores. Postoperative measurements were performed three months after surgery. The tests used were the TMT (A and B), the Digit Span Forward and Backward, the Grooved Pegboard Test, as well as the MMSE. An additional study by Guo et al⁵ used a difference of at least one SD preoperatively and three months postoperatively in two of the following tests: the Verbal Learning Test (Learning Trial and Delay), the Concept Shifting Task (part C), the Stroop Color Word test (Part 3), and the Letter Digit Coding. The author did not mention which specific tests were altered in the postoperative period. Finally, in the study conducted by Micha et al,⁷ the authors diagnosed POCD based on a significant statistical difference between means of tests performed at 9 months postoperatively, compared to the preoperative results. Among the tests used, the ones reported as altered were: The Controlled Oral Word Association Test (COWAt), the Stroop Neuropsychological Screening, the Clock Test, the Three Word-Three Shapes, the Babcock Story Recall, the Instrumental Activities Daily Living (IADLS) and the Trail Making-B.

Discussion

The results from our systematic review and meta-analysis suggested that the incidence of POCD-DCR following the use of TIVA may be lower compared to inhalational anesthesia in the first 30 postoperative days. Even though we have succeeded in including a high total number of subjects in our review and meta-analysis, the heterogeneity in definitions of POCD-DCR, different psychometric tests used and its cuff-off values, among other factors, limit the reach of our conclusions. This is reflected in our heterogeneity analysis ($I^2 = 85\%$). Our study suggests that the concept of POCD should be redefined into a more objective definition. Moreover, it would be of interest to evaluate a more basic cognitive domain, such as attention, in all awake and alert individuals, since it is well known from the literature that attention plays a pivotal role to the functions of all other cognitive domains. It is reasonable to assume that specific cognitive deficits, such as memory, executive function, among others, may reflect a subjacent attention impairment. Future research should focus on objective attention measurements prior to other specific cognitive domains. This would allow a reduction in heterogeneity on POCD research.

Our results agree with a previous Cochrane systematic review on the same topic published in 2018,²⁴ before the consensus stressed the importance of subclassifying POCD definitions according to time in the postoperative period.¹³ This explains why these authors only analyzed the period up to 30 days postoperative, since the late postoperative period, ie, between 30 days and 12 months postoperative, only came into focus for researchers after the 2018 consensus. Additionally, this is the reason why we included 10, instead of only 7 studies in the previous mentioned review paper. Moreover, we decided to conduct a systematic review in the period between 30 days and 12 months postoperative. Just like in our paper, those authors decided to define the outcome measure, ie, POCD, based on author's definition in each individual included study, as a binary variable (yes or no), to proceed with the meta-analysis.

The potential benefits of propofol and TIVA in POCD might be mediated through its positive effects in diminishing the inflammatory cascade. Evidence has shown that propofol has anti-inflammatory properties compared to inhalation agents²⁵ as in vivo study results have shown lower levels of circulating cytokines and other mediators of inflammation in animals injected with propofol.¹¹ Inflammation has been associated with POCD in many different studies. An increase in various cytokines, including IL-6, TNF- α , IL-8, and IL-10, have been correlated with postoperative cognitive impairment.²⁶ Recently, a meta-analysis was conducted assessing the association between various inflammatory biomarkers and POCD and concluded that higher postoperative C-reactive protein (n = 11 studies) and IL-6 (n = 17 studies) were associated with POCD.²⁷ It is true that the possible role of the anesthetics in the inflammatory cascade is still yet to be clarified, with some evidence favoring the use of inhalational agents, such as sevoflurane, specifically in ischemia-reperfusion cell models.²⁸ However, data favouring inhalational anesthetics in ischemia-reperfusion models were produced in the scenario of cardiopulmonary bypass surgeries, in which the etiopathogenesis of POCD is far more complex and diverse.²⁹

The population enrolled in our review has a high median age, consequentially to the fact that most of the research on POCD involves elder individuals. Only one of the included studies admitted patients younger than 60 years old,⁴ and we did not have enough information to perform a subgroup analysis stratified by age. Even though all included studies relied on comparisons of the results of psychometric tests postoperative with the preoperative evaluation, only five studies used

healthy controls not submitted to surgery or anesthesia in its study design,^{3,5,19,21,22} to make sure the cognitive decline was not consequence of the advanced age itself.

One limitation that must be stressed refers to the fact that propofol was used in both groups in all included studies, at least as a single bolus agent at the induction phase of anesthesia. It is uncertain if a single dose of propofol might exert any potential beneficial effects on POCD, consequentially to its potential effects at the inflammatory cascade, even considering that in TIVA group propofol is used in a continuous infusion through all the duration of the procedure. Maybe future studies in POCD should consider using another induction agent in the inhalational group, at the study design phase.

Regarding our secondary aim, we decided to proceed only with a systematic review of the literature, considering the few studies included and, consequently, the small number of subjects, given that the recommendations for testing between 30 days and 12 months postoperatively are relatively recent. Further studies, considering this testing period, are necessary in the future.

Most of these psychometric tests aim at a specific domain of cognitive function (Supplementary Table 1). Many of these tests have been validated in different clinical scenarios, including the postoperative period.^{30–37} It seems reasonable to hypothesize that different psychometric tests, targeting different cognitive domains, might differ in their ability to diagnose POCD. In addition, little attention has been spent on which specific tests and cognitive domains would be most likely altered in the postoperative period. So far, we have scarce evidence of which cognitive domains are more susceptible to POCD, with few data pointing towards the attention domain and executive function as potentially more affected in the postoperative period.³⁸ As mentioned earlier, the attention domain plays a pivotal role in cognition since its proper function is essential to the functioning of all other domains. However, all the studies included in this review established the diagnosis of POCD accepting any altered domain as equally valid. This should, as well, be an important topic for future research.

Only two authors in our review reported which specific tests showed a significant difference in their postoperative assessment. One study reported that tests most frequently altered were the Semantic Verbal Fluency and the Letter Number Sequence Test, which measures the executive function and speed and visual space working memory cognitive domains.⁴ Another author reported that the COWAT, the Stroop Neuropsychological Screening, the Clock Test, the Three Word-Three Shapes, the Babcock Story Recall, the Instrumental Activities Daily Living (IADLS), and the TMT-B as the tests showed a difference in their postoperative assessment.⁷

Additionally, the application of psychometric tests for diagnosis of POCD that relies on a cut-off, such as one SD from the mean, or similar, could be insensitive for detecting minor but significant changes in cognitive status in the postoperative period. We hypothesize that the use of a test more focused on the attention domain, a pre-requisite for the proper function of all the other cognitive domains, could potentially be more sensitive in detecting subtle changes in the cognitive function perioperatively. This should be an additional relevant topic for future research.

It should be emphasized that nine out of the ten studies that we included in our present review titrated the level of anesthesia in both groups with the use of EEG-derived monitors, such as the BIS, all studies targeting a value between 40 and 60. The use of these devices might potentially lead to improved titration of anesthesia.³ Therefore, our results cannot be explained as a consequence of monitoring the level of consciousness on a particular group.

We considered it was appropriate to judge the quality of the evidence we found based on the GRADE system.¹⁶ Systematic reviews of RCTs are usually classified as a high level of evidence. However, the assessment of POCD is still a matter of controversy due to the difficult to select adequate instruments to evaluate basic cognitive domains, such as the attention domain. Consequently, we acknowledge that the quality of evidence produced in our study is most likely moderate, and the strength of recommendations based on our findings should be cautious. We believe that the evidence we found in this study should lead to future research on the association between TIVA and a potential lower incidence of POCD, with more objective definition and assessment of the outcome.

In conclusion, TIVA might be associated with a lower incidence of POCD, compared with inhalational anesthesia, at least in the first 30 postoperative days. However, future studies investigating POCD should focus on assessments of attention because the proper functioning of other cognitive subdomains (eg, memory, executive functions, etc.) relies on attention integrity. This could also potentially reduce heterogeneity on POCD research.

Disclosure

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