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Determinants of post-operative cognitive decline in elderly people

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Highlight:

- **Drop of MoCA after surgery is not associated with the type of anesthesia**
- **Pre-operative treatment and well-being are associated to MoCA points loss after surgery**
- **Per-operative complications and ketamine administration affect post-operative MoCA**

Abstract:

Background: Surgery and anesthesia can result in temporary or permanent deterioration of the cognitive functions, for which causes remain unclear.

Objectives: In this pilot study, we analyzed the determinants of cognitive decline following a non-emergency elective prosthesis implantation surgery for hip or knee.

Design: Prospective single-center study investigating psychomotor response time and changes in MoCA scores between the day before (D-1) and 2 days after (D+2) following surgery in Lariboisière Hospital (Paris, France).

Participants: 60 patients (71.9±7.1-year-old, 72% women) were included.

Measurements: Collected data consisted in sociodemographic data, treatment, comorbidities and type of anesthesia (local, general or both). Furthermore, we evaluated pain and well-being before as well as after the surgery using point scales.

Results: Post-operative (D+2) MoCA scores were significantly lower than pre-operative ones (D-1) with a difference of median[IQR]=2[4]pts ($p<0.001$), we found no significant difference between locoregional and general anesthesia. Pre-operative benzodiazepine or anticholinergic treatment were also associated to a drop in MoCA scores ($p=0.006$). Finally, the use of ketamine during anesthesia ($p=0.043$) and the well-being ($p=0.006$) evaluated before intervention, were both linked to a reduced cognitive impact.

Conclusion: In this pilot study, we observed a post-operative short-term cognitive decline following a lower limb surgery. We also identified pre and perioperative independent factors linked to cognitive decline following surgery. In a next stage, a larger cohort should be used to confirm the impact of these factors on cognitive decline.

Keyword: Cognitive Decline, lower limb surgery, benzodiazepine, ketamine, well-being

Abbreviations: AC: Anticholinergic, BDZ: Benzodiazepine, MoCA: Montreal Cognitive Assessment, POCD: Post-Operative Cognitive Decline.

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INTRODUCTION

Post-Operative Cognitive Decline (POCD) is a major cause of mortality and morbidity costing over \$150 billion dollars yearly in health care expenses in the United States (Braunwald, et al. 2001). POCD includes post-operative delirium, NeuroCognitive Disorder (NCD) and delayed NeuroCognitive Recovery (Mahanna-Gabrielli, et al. 2019). NCD definition from the DSM-V consists in a significant cognitive decline from a previous level of performance, diagnosed at least 30 days after the surgery and assessed by standardized neuropsychological testing. Delayed NeuroCognitive Recovery are tested using the same criteria as for NCD except that the diagnosis window must be less than 30 days after the intervention (Mahanna-Gabrielli, et al. 2019).

POCD can lead to a loss of autonomy in the elderly (>60 year-old (yo)) (Fritz, et al. 2016, Wildes, et al. 2019). However, clear negative consequences of anesthesia on POCD are still debatable (Rasmussen, et al. 1999, Dokkedal, et al. 2016). Some papers speculate that POCD in the elderly is attributable to per-operative use of benzodiazepines (BDZ) and anticholinergic (AC) drugs (Pandharipande, et al. 2006, Pratico, et al. 2005), while others suggest that POCD would result from a pre-existing cerebral fragility (Wildes, et al. 2019, Dokkedal, et al. 2016). Taking into account the ageing of the population linked to a growing demand for surgery, the identification of factors increasing the risk of POCD or more severe cognitive dysfunction is needed.

Diagnosing NCD and delayed NeuroCognitive Recovery requires preoperative cognitive status along with additional stages into the routine care. Indeed, a minimal setup depends upon establishing a baseline of cognitive functions prior the surgery, then on the administration of at least a second neuropsychometric test in the post-operative period (Tsai, Sands and Leung 2010). Tools for quick assessment of POCD are for example the Confusion Assessment Method (CAM), Mini Mental State (MMS) or the Montreal Cognitive Assessment (MoCA) (Fritz, et al. 2016, Nasreddine, et al. 2005).

In particular, the MoCA test evaluates several cognitive functions which impairment is a common mechanism shared by most of POCD, including memory, concentration and verbal abstraction among others. Identifying

POCD risk factors (drugs, comorbidity, surgical or anesthesia setups) might help targeting patients that should be followed-up, leading to a personalized care. This prospective pilot study aims to investigate the cognitive decline following a non-emergency elective hip or knee replacement, and to identify independent risk factors associated with it. Additionally, we search for factors linked with persistent cognitive decline evaluated six weeks after the surgery.

METHODS

Study population

Between February and September 2017, patients from the orthopedic department of Lariboisière Hospital, programmed for a non-emergency hip or knee replacement and who provided consent, were included in this monocentric, prospective study based on the daily clinical practice. We did not include non-French-speaking patients and those refusing to participate to the study. Based on the routine clinical practice, all patients underwent a standardized clinical examination, including medical history and physical examination, laboratory tests were performed in all subjects including chemistry panel and complete blood count. Patients underwent either a general anesthesia (GA), a locoregional anesthesia (LRA) or LRA+ propofol (LRA+P).

Cognitive assessment

To evaluate the cognitive diminution, we administered the MoCA one day before, then two days and six weeks after the surgery. The same practitioner administered MoCA tests. Additionally, the same patient never performed twice the same version of the test.

Covariates

To identify determinants of post-operative decline, we collected during the pre-, per-, and postoperative periods the following covariates:

- Preoperative period: Age; Tobacco; Alcohol; Diabetes; Cholesterol; Hypertension (HT); Thyroids; Feeling of defective memory; Previous Neoplasia; Previous GA; Previous LRA; Cognitive Complaints; Anti-hypertensive treatment; Anti-diabetic treatment (per os); AC drugs; Pre-op. Analgic; Pre-op. NSAID; BDZ; Antidepressants; MoCA (D-1) score; MoCA (D-1) evaluation duration (min); Self-evaluation of pain (D-1) based on a point scale ranging from 0 (no pain) to 10 (extreme pain); Well-being score evaluated using a custom point scale ranging from 0 (no happiness) to 10 (extreme happiness); Instrument Activity Daily Living (iADL) based on the Lawton scale.
- Peroperative period (surgery and anesthesia): Type of anesthesia as described above; Ketamine; Sufentanyl; Corticoids; Droperidol, Ephedrine; Atropine; Tranexamic acid; Clonidine chlorhydrate; Surgery duration; Occurrence of complications (see Appendix).
- Post-operative period: Analgic; NSAID; Nefopam chlorhydrate; Pregabalin; AC; BDZ; Peri-operative complications; MoCA (D+2, W+6) score; MoCA (D+2, W+6) evaluation duration (min); Well-being scale; Self-evaluation of pain (D+2, W+6); Well-being (D+2) score.

Statistical analysis

Numerical variables were expressed by the median and interquartile range or mean and standard deviation, categorical variables were expressed as the count and percentage. For all statistical test, we chose $\alpha=0.05$ as significance level. Patient characteristic data, duration of surgery, and MoCA scores were compared using chi2, Student's t-test or Wilcoxon test as appropriate. When t-test was used, the distribution normality was assessed with the two-tailed Shapiro-Wilk and Lilliefors tests. Sensitivity of variable of interest were computed from confusion matrices.

Comparison of MoCA (D-1) scores distribution and medians between GA, LRA, and LRA+P categories were analyzed with Kolmogorov-Smirnov and Mann-Whitney tests, both two-tailed.

For binary analysis, we associated for each patient the dummy variable 1 for the loss of at least one MoCA point between (D-1) and (D+2), 0 otherwise. We compared this response variable to binary variable and non-binary variables. To identify factors associated to a post-operative decline, we first proceeded with a univariate analysis (Chi2 for dummy variables and Logistic Regression (LR) for non-binary variable). We computed associated factors unless there were less than 5 occurrences in the entire population. Time duration of MoCA administration were compared using a two tailed paired t-test, after log-transform and normality evaluation (Shapiro-Wilk).

To investigate the impact of AC and BDZ treatment, we constructed two dummies variables for 'AC' AND 'BDZ' and for 'AC' OR 'BDZ' treatments. We searched for an association between these variables and cognitive decline ($\Delta\text{MoCA} \geq 1$) between D-1 and D+2.

We used a multivariate class weight corrected logistic regression model to produce a risk model for loss of at least one MoCA point between D-1 and D+2. Variables with a p-value ≤ 0.05 in the univariate analysis were introduced in the multivariate model. In the case of incomplete data, patients were excluded from the multivariate analysis. We performed statistical analysis using R-studio software.

RESULTS

Sixty patients (71.9 ± 7.1 yr., 72% women) above 60 year-old were prospectively included in this study (Fig. 1, Tab. 1). Among them, 27 (resp. 33) underwent a non-emergency elective prosthesis hip (resp. knee) implantation surgery. The distributions of age, time of surgery, MoCA, pain and well-being scores were not different between patient from the hip and the knee groups (Kolmogorov-Smirnov test, insignificant difference, data not shown). Therefore, for the statistical analysis we decided to consider patients that had a hip and knee surgery as a single study group.

Patients underwent GA, (n=27) or LRA (n=14) or LRA+ P (n = 13). There was no significant difference, neither in mean nor in variance, on the Δ MoCA between D-1 and D+2 between the 3 groups of anesthesia (Fig. 2). We investigated if MoCA scores obtained one day before (D-1), two days after (D+2) and six weeks (W+6) following the surgery were significantly different. MoCA score drop was 2[4] points (median[IQR], $p < 0.001$) between (D-1) and (D+2) (Fig. 3A) including 8, 14 and 19 patients who lost 1, 2 or 3 or more points respectively. We found no significant difference between D-1 and W+6 (MoCA median[IQR]: 24[3.5] versus 24 [4], p -value= 0.831), although among the 31 patients evaluated at 6 weeks, 32% did not recovered their baseline level score. We observed a significant decrease in the evaluation duration of the MOCA between (D-1) and (D+2) ($p < 0.001$) and between (D+2) and (W+6) ($p < 0.001$). The surgery duration did not affect the cognitive functions (median[IQR] = 70[40] min, ranging: 30 to 200min).

In the univariate analysis, we found a positive significant relationship between Δ MoCA ≥ 1 and BDZ treatment (p -value = 0.047, sensitivity = 93.8%), AC drugs (p -value = 0.023, sensitivity = 100%), 'BDZ or AC' (p -value = 0.006, sensitivity = 93.8%), perioperative complications (p -value = 0.035, sensitivity = 100%) and a negative significant result with the use of ketamine agent (p -value = 0.043, sensitivity = 62.5%). Results are summarized in Tab. 1. Furthermore, among non-binary variable, only well-being (p -value = 0.006, sensitivity = 94.6%) was significantly associated with cognitive decline. Using a multivariate analysis, we found that 'well-being', 'Ketamine', 'AC' or 'BDZ', were independent risk factors associated with cognitive decline between D-1 and D+2 (see Tab. 2A).

Similarly, we found that, persistent cognitive decline was associated with BDZ (p -value = < 0.001) and perioperative complications (p -value = 0.013). Additionally, we found that post-operative confusion was also associated with a W+6 cognitive decline (p -value = 0.002). Results are shown in Tab. 2B.

DISCUSSION

In this study, MoCA scores were significantly impacted by the surgery, with a loss of 2.25 points in average. We described that POCD were independently associated with pre-operative well-being, per-operative use of ketamine, peri-operative complications and pre-existing treatment based on either BDZ or AC drugs. In particular, well-being and ketamine had a protective effect, while complications and BDZ/AC treatments were linked to cognitive decay. Here, the type of anesthesia was not linked to cognitive decline and, persistent cognitive decline was linked to pre-existing BDZ medication and per-operative complications. Finally, MoCA evaluation duration was unchanged according the kind of anesthesia and the occurrence of POCD, however, the duration at W+6 was significantly shorter than D-1 and D+2 reflecting the training effect (Fig. 3B). These results suggest insight to improve perioperative cognitive protection and in particular to identify upstream of the intervention patients that could be affected by surgery or anesthesia.

Ketamine is a non-barbiturate anesthetic inhibiting acetylcholine that could lead to an increased number of delirium (Sleigh, et al. 2014). Per-operative use of ketamine is still debated, with on one side studies defending its neuroprotective and anti-inflammatory actions (Dale, et al. 2012), and on the other side studies arguing for no beneficial effects (Avidan, et al. 2017). Interestingly ketamine was shown to speed up recovery to consciousness after GA, yet possible impact on POCD was unclear (Hambrecht-Wiedbusch, Li and Mashour 2017). In the present study, per-operative ketamine administration was associated to fewer POCD, this support previous findings suggesting that administration of ketamine during a surgery reduces occurrences of POCD.

Chaiwat et al 2019 (Chaiwat, et al. 2019) reported more POCD among patients sedated with propofol, which was also suspected to decrease mean arterial pressure with a risk of brain hypoxia leading to cognitive decline (Wild, et al. 2008). However, ketamine could preserve the hemodynamic stability during propofol-based anesthesia. In our study, we compared three groups with and without propofol. We did not find any impact of the kind of anesthesia on the cognitive decline suggesting that propofol did not impact cognition in this cohort.

Furthermore, patients who received ketamine also received propofol supporting the hypothesis of a benefic effect of the ketamine during propofol infusion. However, even if several studies are in favor of ketamine protective effect, further prospective randomized controlled studies would be needed to confirm this hypothesis.

BDZ and AC drugs have long been suspected to be involved in drug-induced cognitive decline (Starr and Whalley 1994, Bartus, et al. 1982). On one hand, BDZ act as positive allosteric modulators of GABA-A (γ -aminobutyric acid) receptors, which results in an overall increased cortical inhibition, the latter possibly responsible for mental deterioration (Griffin III, et al. 2013). On the other hand, delirium pathway arises from a dopamine excess and acetylcholine decrease, that could be due or accentuate directly by AC drugs (Maldonado 2013). Furthermore, consequences of BDZ or AC treatment on cognition have been extensively described for speed-up age-related cognitive decay and Alzheimer's disease, not for POCD (Gage, et al. 2014, Grande, et al. 2018). In the present study, using a homogenous prospective non-emergent population, we observed detrimental effects of BDZ and AC drugs on cognitive functions. This finding supports previous results and hypothesis, however larger prospective and multicentric studies will be needed to confirm these results, allowing recommendations regarding the use of BDZ and AC drugs in non-emergent surgery. Clarifying a possible mechanisms underpinning BDZ/AC deleterious effects on sedated patients will benefit the prevention of temporary delayed NeuroCognitive Recovery as well as permanent POCD (NCD). We already know that the GABA-A agonist role of propofol is potentiated by BDZ, that could in turn promote POCD (Orser and Miller 2001). However, among our 54 patients, 15 had no propofol, yet for this subpopulation the cognitive decline was statistically undistinguishable from the 39 other patients. A regular intake of BDZ was also associated to a reduced cognitive reserve (Gage, et al. 2014), the latter linked to more POCD (Feinkohl, Winterer and Pischon 2017). Nevertheless, we found no correlation between MoCA (D-1) scores or sociocultural levels, and intake of BDZ/AC drugs.

Per-operative complications were associated with Δ MoCA both two days and six weeks after the surgery. These findings are consistent with previous works (Greene, et al. 2009), and even more with recent works about complications following total hip arthroplasties (Aziz, et al. 2018). This suggests that an extended period under anesthesia does not affect post-operative cognition while a physiological response to a surgical complication does. Surgery-related inflammatory response has been associated to POCD (Alam, et al. 2018). Although we did not collect inflammatory-related biomarkers, we can note that among patient with peri-operative complications, only two had NSAIDs and three had ketamine.

Finally, depressed mood, in particular among the elderly, was associated with cognitive impairment (Brommelhoff, et al. 2009). Several studies outlined the link between POCD and elderly mood deterioration, assessed using Geriatric Depression Scale for patients receiving a cardiac as well as a non-cardiac surgery (Rudolph and Marcantonio 2011, Greene, et al. 2009). In our study, we found no link between cognitive decline and patient treated for depression, which agrees with these studies. Interestingly, we found that well being measured before and after the surgery were not linked to pain scores, in fact pre-operative well-being was here an independent factor associated with POCD, which was not the case for pain. These results might be explained by the fact that pain was carefully managed after the surgery (av. score \approx 4), which does mitigate POCD (Inouye, et al. 2015). In the end, it emerges from this study that well-being measured before the surgery is a better indicator of cognitive decline than post-operative pain, which suggests that the patient's state before surgery affects post-operative cognitive trajectory.

Limitations and perspectives

This study has some limitations. The patients included were very homogeneous but the inclusion criteria, the prospective methodology during a limited period led to a small number of patients. Larger multicentric studies

would be needed to confirm or not these results and to propose recommendation regarding non-emergency arthroplasty.

In some group the number of patients was too low to provide useful data and/or multivariate analysis. However, the prospective methodology and our statistical approach allowed us to underline interesting insights about POCD. This article is the first result of a pilot study suggesting that cognitive decline occurring after a surgery could be anticipated based on pre-operative factors, yet only partially. We are performing a larger study in order to identify patients who are at risk of both delayed neurocognitive recovery and permanent cognitive decline.

CONCLUSION

To conclude, we reported a significant cognitive decline two days following a non-emergency lower limb surgery. BDZ or AC drugs were associated with a poor cognitive trajectory, both in short-term, but also in long-term in the case of BDZ drugs. At the same time, use of per-operative ketamine appeared to have a cognitive protective effect. Pre-operative well-being and perioperative complications were the two non-drug factors associated with positive and negative outcome respectively. Findings in the present study suggest that patient's medical prescriptions and the surgery/anesthesia smooth progress can both independently affect POCD. Nevertheless, further research, based on a larger cohort, should investigate separately factors discussed above, in particular drugs for which neurophysiological mechanisms remain unclear and the role in cognitive decline/protection is still a divisive topic.

STATEMENT OF INFORMED CONSENT

We obtained an authorization by the French data privacy administrative body 'Commission Nationale de l'Informatique et des Libertés' (CNIL), under the reference number z2c2680420w. In agreement with the ethics

committee for this non-interventional study an oral agreement was obtained from each patient during their first medical visit.

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CONFLICTS OF INTEREST

Pr. C. PAQUET is member of the International and National Advisory Boards of Lilly, ROCHE, Biogen. She is consultant of Fujiribio, ALZOHIS, NEUROIMMUNE and GILEAD and is involved as investigator in several clinical trials for Roche, Eisai, Lilly, Biogen, Astra-Zeneca, Lundbeck, Neuroimmune.

Dr. J. DUMURGIER is investigator in several passive anti-amyloid immunotherapies and other clinical trials for Roche, Eisai, Lilly, Biogen, Astra-Zeneca, Lundbeck. Similarly, Dr. C. HOURREGUE is investigator for Roche, Eisai and Biogen.

Pr. R. NIZARD, Dr. C. RABUEL, Dr. L. COBLENTZ BAUMANN, Dr. C. LOYER, Dr. J. CARTAILLER and Dr. E. VANDERLYNDEN declare that they have no conflict of interest.

AUTHORS' CONTRIBUTIONS

Claire Paquet designed the study and analyzed the data and prepared the manuscript. Remy Nizard, Christophe Rabuel and Camille Loyer included the patients. Jerome Cartailier, Julien Dumurgier and Claire Paquet have performed statistical analysis and results interpretation. All authors read and validated the manuscript

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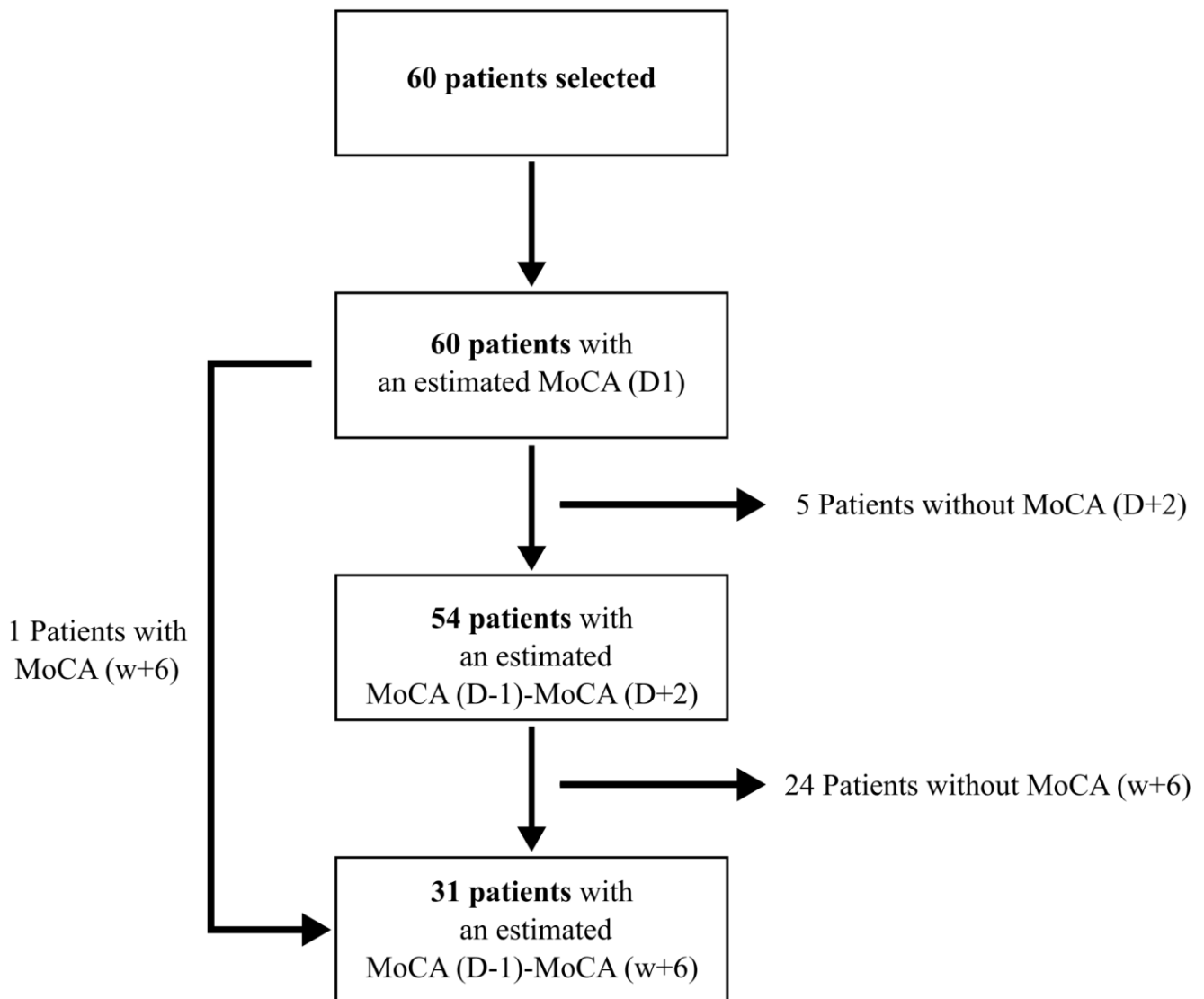


Figure 1: Flowchart

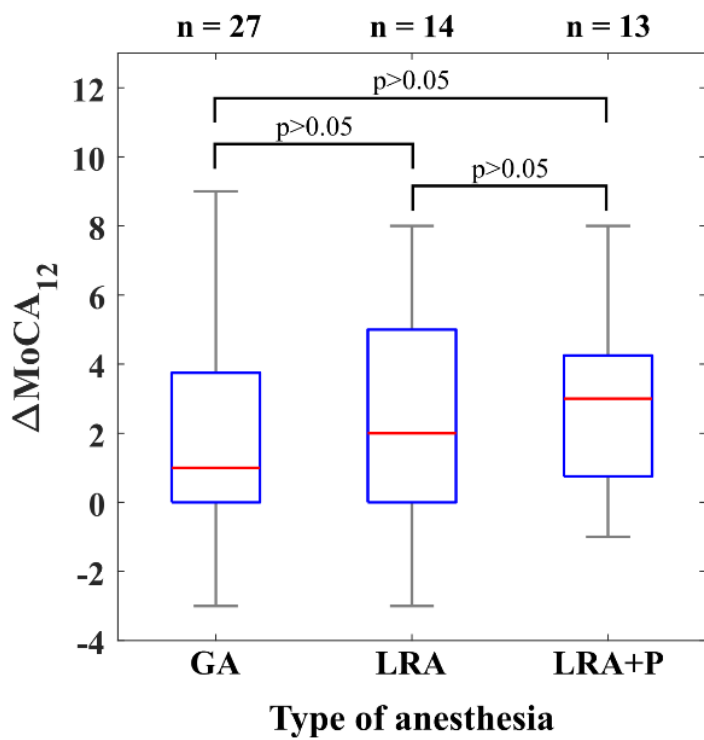


Figure 2: Distribution of ΔMoCA between D-1 and D+2 for the three types of anesthesia (GA, LRA, LRA+Propofol) showing no significant difference in (Wilcoxon test).

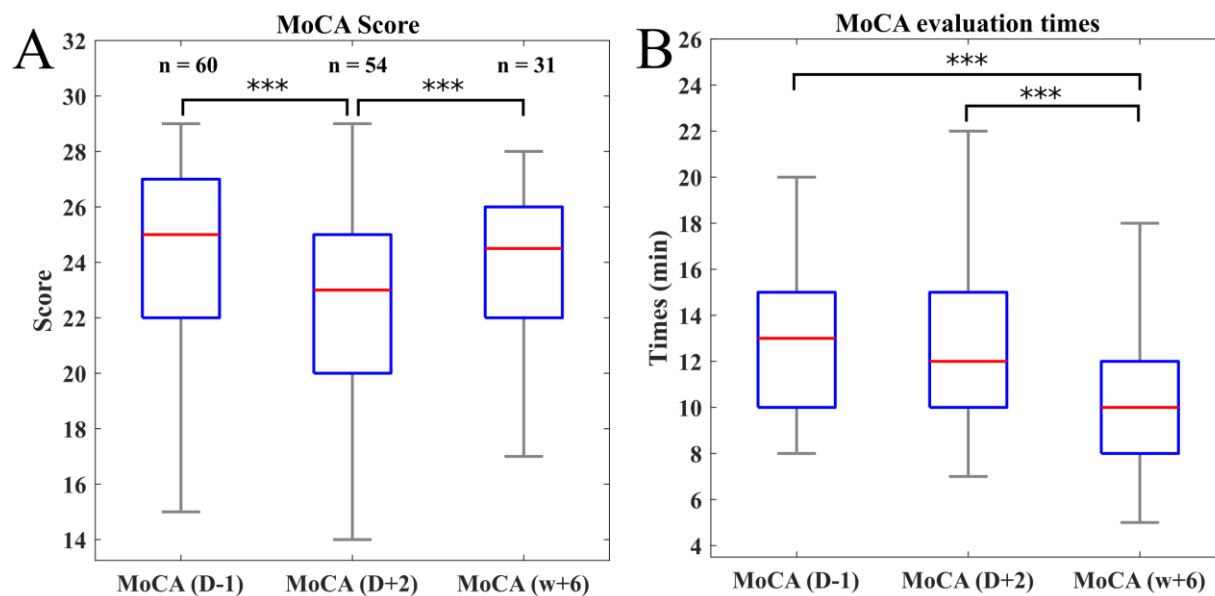


Figure 3: Distribution of MoCA score (**A**) and evaluation time (**B**) for three different periods: one day before (D-1), two days (D+2) and six weeks (w+6) following the surgery and anesthesia.

Table 1: Comparison of population characteristic data between the patient with $\Delta MoCA \geq 1$ pt and patients with no point loss. Binary data are number and percentage, while non binary data are shown in median and interquartile range (IQR). NSAIDs ; Nonsteroidal anti-inflammatory drugs, † includes the following complications: failed LRA – modification of the surgery – Hemorrhage – incisors defect (see Supplementary).

VARIABLES	All (n=54)		Unaffected MoCA group (n=16)		Altered MoCA group (n=38)		p-value
	n or median	(%) or [IQR]	n or median	(%) or [IQR]	n or median	(%) or [IQR]	
PRE-OPERATIVE							
Age (yrs.)	70.5	[7.5]	69.5	[5.5]	72	[7.8]	0.259
Sex (female)	39	72.2%	9	56.3%	30	79.0%	0.089
Sociocultural level	3	[2]	3	[2]	3.5	[2]	0.571
Instrumental Activities of Daily Living	4	[0]	4	[0]	4	[0]	0.731
Smoker	8	14.8%	2	12.5%	6	15.8%	0.756
Alcohol (0/1)	6	11.1%	3	18.8%	3	7.9%	0.246
Hypertension (0/1)	35	64.8%	12	[75]	23	60.5%	0.309
Cholesterol	24	44.4%	8	[50]	16	42.1%	0.594
Diabetes (0/1)	11	20.4%	3	18.8%	8	21.1%	0.848
Thyroid	7	13.0%	1	6.3%	6	15.8%	0.341
Cognitive trouble in family (0/1)	6	11.1%	1	6.3%	5	13.2%	0.461
Nb. of drug	4.5	[5]	3	[3.8]	5	[4]	0.204
Anti-hypertensive treatment (0/1)	1	[1]	1	[0.5]	1	[1]	0.163
Anti-diabetic treatments (0/1)	10	18.5%	3	18.8%	7	18.4%	0.977
Preexisting NSAIDs treatment (0/1)	16	29.6%	3	18.8%	13	34.2%	0.256
Preexisting antalgic treatment (0/1)	42	77.8%	11	68.8%	31	81.6%	0.3
Preexisting antidepressant treatment (0/1)	7	13.0%	1	6.3%	6	15.8%	0.341
Preexisting benzodiazepine treatment (0/1)	13	24.1%	1	6.3%	12	31.6%	0.047
Preexisting anticholinergic treatment (0/1)	10	18.5%	0	0.0%	10	26.3%	0.023
Preexisting anticholinergic OR benzodiazepine treatment (0/1)	18	33.3%	1	6.3%	17	44.7%	0.006

MoCA score (D-1)	25	[4.8]	24	[3.2]	25.5	[4]	0.035
Time to perform MoCA (min)	13	[5]	14	[3]	12	[5]	0.683
Memory complaint (0/1)	16	29.6%	3	18.8%	13	34.2%	0.256
Pain scale (D-1)	0	[5]	0	[0.2]	0	[5]	0.084
Well-being (D-1)	8	[2]	8	[1]	7.5	[2.5]	0.006
Nb. Previous general anesthesia	2	[3]	1.5	[2.2]	2	[2]	0.092
Nb. Previous loco-regional anesthesia	0	[0]	0	[1.2]	0	[0]	0.259
PER-OPERATIVE							
General anesthesia (0/1)	27	50.0%	8	50.0%	19	50.0%	0.999
Loco-regional anesthesia (0/1)	14	25.9%	4	25.0%	10	26.32%	0.712
Loco-regional + propofol (0/1)	13	24.1%	4	25.0%	9	23.68%	0.690
Surgery duration (min)	70	[40]	70	[35]	75	[40]	0.812
Propofol	39	72.2%	11	68.8%	28	73.7%	0.712
Sufentanyl	30	55.6%	9	56.3%	21	55.3%	0.947
Atracurium	24	44.4%	7	43.3%	17	44.7%	0.947
Ropivacaine	25	46.3%	9	56.3%	16	42.1%	0.341
Marcaïne/bupivacaine	21	38.9%	5	31.3%	16	42.1%	0.455
Ketamine	11	20.4%	6	37.5%	5	13.2%	0.043
Ephedrine	25	46.3%	6	37.5%	19	50.0%	0.4
Atropine	8	14.8%	3	18.75%	5	13.2%	0.597
Clonidine chlorhydrate	6	11.1%	2	12.5%	4	10.5%	0.833
Droperidol	30	55.6%	9	56.3%	21	55.3%	0.947
Tranexamic Acid	29	53.7%	6	37.5%	23	60.5%	0.121
Per-operative complications†	8	14.8%	0	0.0%	8	21.1%	0.047
POST-OPERATIVE							
Post-operative Confusion	7	13.0%	0	0.0%	7	18.4%	0.066
Pain scale (D+2)	4.2	[4]	2.8	[5.2]	4.8	[4]	0.244
Post-operative well being	8	[3]	9	[2.5]	8	[2]	0.3
MoCA at D+2	23	[5]	24.5	[4.2]	22	[5]	0.014
Time to perform MoCA (min)	12	[5]	11	[4.2]	12	[5]	0.294

Table 2: Factors associated with cognitive decline at D+2 and w+6.

2A Sensitivity of risk factors of cognitive decline at D+2 computed with their significance computed from multivariate logistic regression for Ketamine, Well-being and 'AC or BDZ'

VARIABLE	SENSITIVITY	P-VALUE
Per-op. complication	100%	-
Well-being = 0	94.6%	0.012

2B Patient characteristic associated to cognitive decline at six weeks (w+6) using a chi2 analysis

VARIABLE	$\chi^2(1)$	P-VALUE
BDZ	15.62	<0.001
Post-op. Confusion	9.64	0.002
Per-op. complication	6.22	0.013

AC OR BDZ	93.8%	<0.001
Ketamine	62.5%	0.026

Appendix:

Statistic/ ΔMoCA₁₂	AG	ALR	ALR + P	All
n	27	14	13	54
Minimum	-3	-3	-1	-3
Maximum	9	8	8	9
Mean	1.89	2.36	2.92	2.25
Standard deviation (n-1)	3.18	3.30	2.57	3.05

Table A1: MoCA D-1/D+2 drop statistics for the three types of anesthesia

Type of peroperative complications encountered	n
LRA installation failed	2
A bilateral hip surgery became a hip and knee surgery	1
Hemorrhage	3

Incisors broken during oro-tracheal intubation	1
Signaled but not detailed	3

Table A2: List of peroperative complications