Association between preoperative autonomic nervous system function and post-induction hypotension in elderly patients: a protocol for a cohort study

Quexuan Cui, Lu Che, Han Zang, Jiawen Yu, Li Xu, Yuguang Huang

ABSTRACT

Introduction Post-induction hypotension (PIH), which is prevalent among elderly patients, is associated with adverse perioperative outcomes. As a critical part of blood pressure regulation, baroreflex control is believed to be closely related to intraoperative blood pressure fluctuations. Spontaneous baroreflex sensitivity and heart rate variability measurement can aid evaluation of patients’ autonomic function. This study aims to determine the association between preoperative decreased baroreflex function and PIH in elderly patients.

Methods and analysis This prospective cohort study will enrol patients who are 65 years old and above, scheduled for elective non-cardiac surgery under general anaesthesia, and American Society of Anesthesiologists physical status I–III (n=180). Baseline assessment will include routine preoperative evaluations as well as symptoms and anamneses associated with baroreflex failure. Preoperative autonomic function monitoring will be performed through 20 min of continuous beat-to-beat heart rate and blood pressure monitoring using LiDCO rapid (Masimo Corporation, USA). The primary outcome will be PIH. Detailed use of anaesthetic agents during induction and maintenance will be documented for adjustment in multivariable analyses.

Ethics and dissemination The Research Ethics Committee of Peking Union Medical College Hospital approved the study protocol (I-22PJ008). We aim to publish and disseminate our findings in peer-reviewed journals.

Trial registration number NCT05425147.

INTRODUCTION

Intraoperative hypotension (IOH) is a common event during general anaesthesia, occurring in approximately 40% of patients. Low arterial pressure is associated with perioperative mortality and ischaemia of vital organs, including the brain, heart and kidneys in a ‘dose-dependent’ manner. Advanced age is one of the risk factors for IOH. As an increasing number of elderly people receive surgery under general anaesthesia, IOH in elderly patients becomes an urgent issue.

Hypotension can happen throughout perioperative period but is most prevalent during post-induction phase, which refers to the time frame between anaesthesia induction and skin incision. During this phase, arterial hypotension is associated with anaesthesia induction, baseline haemodynamic status and other patient-related factors rather than surgical factors, such as bleeding and surgical interventions. It is widely recognised that the risk of post-induction hypotension (PIH) increases with age.

Apart from senior age, risk factors for PIH includes the American Society of Anesthesiologists (ASA) Physical Status classification ≥3 and low preoperative mean arterial pressure (MAP), which are all non-modifiable in nature. Preoperative hypovolaemia, a potentially modifiable factor, has also been widely studied in PIH-related research. Preoperative echocardiographic assessment and preoperative fluid replenishment strategies have been put forward to achieve euvoalma.
It is widely recognised that ageing is closely related to the role of baroreflex function in PIH is still unclear. However, there are few studies investigating the relationship between baroreflex function and PIH; therefore, it is widely recognised that ageing is closely related to decreased baroreflex function. Since PIH is prevalent in elderly patients and is associated with adverse postoperative outcomes, we postulate that measurement of BRS in preoperative period may serve as a surrogate marker of individual baseline baroreflex control, and provide information on haemodynamic management among elderly patients undergoing general anaesthesia.

In addition to BRS, heart rate variability (HRV) can also be used to assess ANS. Several studies have revealed the relationship of low HRV and IOH, but the preoperative fluid status assessment was missing in these studies. Time, frequency and non-linear analysis measures were all considered to be related to IOH. We hypothesise that decreased baroreflex function in elderly patients plays a significant role in PIH. This study will enable us to explore whether perioperative short-time BRS and HRV assessment can help to identify elderly patients at high risk of PIH. We will also perform frailty assessment in elderly patients to explore the role of frailty in the association of autonomic regulation and PIH. Overall, this cohort study will provide the relationship between perioperative assessment of autonomic function and outcomes.

**METHODS**

**Study design and setting**
Our study is a single-centre prospective cohort study to investigate the potential impacts of preoperative autonomic function on haemodynamic stabilisation after induction. This study will be conducted at Peking Union Medical College Hospital (PUMCH), Beijing, China. This study has been approved by the Research Ethics Committee of PUMCH (reference number: I-22PJ008) and has been registered in ClinicalTrials.gov (registration number: NCT05425147).

**Participants**
Enrolled patients will be at least 65 years old, scheduled for major elective non-cardiac surgery under general anaesthesia (estimated operation time ≥2 hours) and ASA physical status I–III. The airway management tool during operation will be endotracheal tubes. Exclusion criteria will be severe vascular disease, secondary hypertension, Parkinson’s disease, or other conditions that cause tremors, persistent atrial fibrillation, inability to measure upper extremity blood pressure and mental disorders.

**Protocol**
Our study includes baseline assessment, autonomic function measurement, intraoperative monitoring and postoperative follow-up (Figure 1). All collaborators will be trained on autonomic function measurement. All data will be recorded in the Clinical Report Form (see online supplemental material 1) and kept securely.

**Baseline assessment**
In addition to routine preoperative clinical and laboratory evaluations, we also specifically assess patients for...
autonomic function. As is shown in table 1, the symptoms and anamneses associated with baroreflex failure will be taken.\textsuperscript{25, 26} Medications that may affect autonomic regulation will also be inquired. We will also perform frailty phenotype assessment using FRAIL Scale and Fried phenotype.\textsuperscript{27, 28}

**Table 1** Medical history, symptoms and medications related to autonomic function

<table>
<thead>
<tr>
<th>Medical history</th>
<th>Symptoms</th>
<th>Medications</th>
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<tr>
<td>Afferent baroreflex failure</td>
<td>Blood pressure</td>
<td>Anticholinergics</td>
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<td>Radiation therapy for head and neck cancer</td>
<td>Orthostatic hypotension or syncope</td>
<td>Sympathomimetics</td>
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<td>Resection of neck tumours (bilateral)</td>
<td>Hypertensive crisis</td>
<td>Parasympathomimetics</td>
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<td>Carotid endarterectomy</td>
<td>Hypotensive episode</td>
<td>Mineralocorticoids</td>
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<td>Central baroreflex failure</td>
<td>Heart rate</td>
<td>Adrenergic antagonists</td>
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<td>Multiple system atrophy</td>
<td>Tachycardia at rest, during exercise or orthostasis</td>
<td>Diuretics</td>
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<tr>
<td>Efferent baroreflex failure</td>
<td>Vasomotion</td>
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<tr>
<td>Diabetic neuropathy</td>
<td>Heat/cold intolerance</td>
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<tr>
<td>Parkinson’s disease</td>
<td>Skin colour change</td>
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*Patients with Parkinson’s disease are usually excluded in our study because of tremor.

Autonomic function monitoring

Autonomic function monitoring will be performed approximately 40 min before the start of anaesthesia on the day of surgery (figure 2). Patients will be monitored for 20 min using LiDCO rapid (Masimo Corporation, USA) to obtain continuous heart rate and beat-to-beat blood pressure measurement. The measurement will be performed in the pre-anaesthesia waiting room with an ambient temperature of 25°C and constant lighting status. All included patients will have been fasted for at least 8 hours before the test. Patients will be in supine position and undergo the following instructions.\textsuperscript{14, 29}

1. Baseline monitoring for 2 min.
2. Passive leg raising will be performed to predict central hypovolaemia. LiDCO rapid will be used to measure stroke volume (SV) non-invasively. If ΔSV ≥12%, the patient will be suspected for hypovolaemia and excluded.\textsuperscript{30} Fasting time and preoperative intravenous fluid volume will also be recorded.
3. Spontaneous baroreflex function will be monitored for 15 min during which time patients will be instructed to relax and breathe spontaneously.
4. Enhanced baroreflex function will be measured for 1 min during which time patients will be instructed to do pursed-lips breathing at a frequency of 0.1 Hz, to induce periodical fluctuations of heart rate and blood pressure.\textsuperscript{31}

Matlab programs will be used to calculate BRS and HRV measures from the continuous heartbeat and blood pressure recordings. BRS is expressed as milliseconds change in RR interval per mm Hg in response to changes in systolic blood pressure (SBP). The sequence method will be used for assessment of baroreflex function, which identifies SBP and RR interval have both progressively increased over three or more consecutive beats.\textsuperscript{19, 32} The threshold values for SBP and RR interval changes will be set at 1 mm Hg and 6 ms, respectively. BRS will be obtained by calculating the slope of the regression line relating changes in SBP to changes in RR interval. Baroreflex Effectiveness Index (BEI) will be defined as the percentage of progressive increase/decrease swings in SBP that are effectively regulated by heart rate, that is, RR interval changes.\textsuperscript{33} We define the increase in RR interval with rising SBP as BRS+/BEI+, and vice versa as BRS−/BEI−.

The RR interval data collected during BRS monitoring will also be used for HRV measurement. We will perform time and frequency domain analysis according to the standards of HRV published by the American Heart Association. Time domain analysis, which reflects overall variability of RR intervals, includes indicators such as SDNN (SD of the normal-to-normal intervals),

![Figure 2](image-url) **Figure 2** Timeline of measures. BRS, baroreflex sensitivity; HRV, heart rate variability; POD, postoperative day.
vasopressors will not be used.

Intraoperative measures and anaesthetic agents use
All enrolled patients will receive standard monitoring with a torso-positioned limb leads ECG, pulse oximetry, a peripheral intravenous line and an automated non-invasive blood pressure (NIBP) according to ASA Standards for Basic Anaesthetic Monitoring. NIBP will be measured every 2 min during post-induction period and every 3 min during intraoperative period. Bispectral index monitoring (BIS; Medtronic, Dublin, Ireland) will be used to maintain the proper depth of anaesthesia, with a BIS target range of 40–60.

Intravenous anaesthesia induction will be used for all patients. For pragmatic reasons, anaesthesia agents used (propofol or etomidate) in the induction and maintenance phase are not strictly regulated and will be chosen at the discretion of the anaesthesiologist in charge who is independent from the study. However, during induction, anaesthetic and other accompanying medications will be given according to the following institutional protocol: fentanyl (1–3µg/kg), propofol (1.5–2.5mg/kg) or etomidate (0.1–0.3mg/kg) and rocuronium (0.5–0.9mg/kg). Total intravenous anaesthesia will be induced with fentanyl (1–3µg/kg), propofol (1.5–2.5mg/kg) or etomidate (0.1–0.3mg/kg) and rocuronium (0.5–0.9mg/kg). We will record anaesthetic agents used during anaesthesia and correct for confounding factors through multivariable analysis. Vasopressors will be allowed during post-induction and intraoperative periods. Patients will receive boluses of ephedrine 6mg or phenylephrine 100µg whenever patients meet the standards of hypotension. Pre-emptive vasopressors will not be used.

Postoperative follow-up
We will conduct daily follow-up of patients until hospital discharge and a telephone follow-up at 30 days after surgery. Postoperative recovery, complications and routine laboratory test results will be recorded.

Study outcomes
Primary outcomes
The primary outcome in this study is PIH, occurring within 20 min after induction or before incision. Hypotension is defined as SBP <90 mm Hg, MAP <65 mm Hg or a decrease of more than 30% of baseline.

Secondary outcomes
The secondary outcomes include:
1. Early intraoperative hypotension, occurring within 30 min of surgery.
2. Mortality within 30 days after surgery.
3. Postoperative complications in accordance with the European Perioperative Clinical Outcome definitions. Postoperative complications will be classified according to the Clavien-Dindo system.

Sample size consideration
To investigate the association between BRS, HRV and outcomes, we used the HRV results in the literature to estimate the sample size, considering few previous studies have explored the correlation between BRS and PIH. Our preliminary study suggested that around 35% of elderly patients in our centre developed PIH. Based on the study by Padley and Ben-Menachem and considering the need for a multivariable analysis according to the induction strategy in our study, to achieve a 95% power and a significance level of 5%, 180 patients will be required (60 with PIH and 120 with stable blood pressure). A drop-out rate of 10% is assumed in sample size calculation. The statistical power of various HRV metrics is shown in table 2.

Previous studies have shown that elderly women had lower BRS compared with elderly men. To reduce the impact of differences in sex distribution on measures and outcomes, we included patients in a specific sex ratio. From 2014 to 2022, 41.1% of the elderly patients who received elective non-cardiac surgery under general anaesthesia in our centre were female. We will include patients in a ratio of 6:4 male to female (108 male and 72 female).

Statistical analysis
Statistical analysis will be performed using the SPSS V.22.0 for Windows (IBM). Quantitative variables will be reported as the mean±SD or median (25th–75th IQR). BRS and

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<th>Table 2</th>
<th>Power calculation according to a previous study on HRV</th>
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<td>Time domain analysis</td>
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<td>Mean RR (ms)</td>
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<td>SDNN (ms)</td>
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<td>Frequency domain analysis</td>
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<td>HF (ms²)</td>
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<td>Total power (ms²)</td>
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<td>LF/HF ratio</td>
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<td>HF, high frequency; HRV, heart rate variability; LF, low frequency; RMSSD, square root of mean squared differences of normal-to-normal intervals; SDNN, SD of the normal-to-normal intervals.</td>
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HRV measures will be compared in elderly patients with and without PIH. Categorical variables will be compared using χ² or Fisher’s exact test. Differences in mean or median values between PIH and stable groups were compared using Student’s t-test or the non-parametric Mann-Whitney U test to explore the association between PIH and preoperative ANS function. Logistic regression will be used to estimate the relationship between frailty, BRS and PIH. Since different anaesthetic agents (mainly propofol vs etomidate) used for induction might bring in confounding factors, we will use logistic regression to adjust for confounding. We will consider p<0.05 as statistical significance.

Ethics and dissemination
The Research Ethics Committee of PUMCH approved the study and the reference number was I-22P[J]008 (approved on 10 June 2022). The investigator will comprehensively explain the study to patients before recruitment. A written informed consent form will be obtained from all patients. Patients will be informed that they may decline to participate or withdraw from the study at any time. The study was registered with the National Institutes of Health, US National Library of Medicine at ClinicalTrials.gov (NCT05425147, registered on 15 June 2022).

We aim to publish and disseminate our findings in peer-reviewed journals. Our findings will also be presented at Chinese and international conferences.

Patient and public involvement
Patients or the public were not involved in the design, or conduct, or reporting, or dissemination plans of our research.

DISCUSSION
This study aims to find out the association between decreased baroreflex function and PIH or other adverse events in elderly patients. Through preoperative non-invasive BRS and HRV measurement, we can obtain information on the patient’s preoperative baroreflex function and sympathetic/parasympathetic tone from real-world evidence.

Considering the impact of PIH on prognosis, the mechanism of PIH has attracted attention of investigators. Previous studies have confirmed that absolute volume status could not explain the complete picture of PIH. Blood pressure rhythms, which are regulated by autonomic nerves, are thought to be a potentially critical link in PIH. However, to our knowledge, the study on the association between PIH and baseline BRS is still missing. Accordingly, performing preoperative BRS assessment is an effective way to identify this association. Furthermore, through passive leg raising, we can exclude the effect of hypovolaemia on PIH. Like many studies focused on PIH, anaesthetic agents used during induction cannot be strictly standardised due to clinically practical reasons. To reduce the bias caused by them, we will attempt to correct for confounding factors by statistical methods. In conclusion, despite the limitations, this study might help with early identification and precaution of PIH mediated by impaired autonomic function in elderly patients.

Contributors Concept, study design and first draft of the manuscript—QC, LC and LX. Manuscript review and data collection—HZ, JY, LX and YH. Editing and critical review—LC and LX. All (GC, LC, HZ, JY, LX, YH) authors have given final approval for manuscript submission.

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Competing interests None declared.

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