Effects of levosimendan on occurrence of cerebral vasospasm after aneurysmal subarachnoid hemorrhage: a case–control study

Antoine Trinh-Duc1,2†, Marc-Antoine Labeyrie3,4†, Anaïs Caillard1,2, Wagih Ben Hassen5, Alexandre Mebazaa1,2 and Benjamin Glenn Chousterman1,2*

Keywords: Subarachnoid hemorrhage, Cerebral vasospasm, Delayed cerebral ischemia, Levosimendan

Dear Editor,

Delayed cerebral ischemia (DCI) is a common complication following aneurysmal subarachnoid hemorrhage (aSAH) contributing to poor prognosis [1]. Macrovascular dysfunction related to cerebral vasospasm (CVS) remains the main pathophysiological hypothesis and therapeutic target of DCI [2]. No preventive treatment for CVS has yet proven to be effective [3].

Levosimendan, a non-adrenergic calcium-sensitizing inotrope, has been used to treat neurogenic stress-induced cardiomyopathy (NSIC) related to aSAH. Only few cases of patients treated with levosimendan for NSIC related to aSAH have been reported and have suggested that levosimendan could be associated with hemodynamic and neurological improvement [4]. The aim of this study was to retrospectively evaluate the efficacy of levosimendan as a therapy targeting occurrence of CVS in a cohort of patients admitted in ICU for aSAH.

We retrospectively reviewed the medical records of patients admitted to the Lariboisière Hospital Surgical Intensive Care Unit (ICU) (Paris, France) for an aSAH and treated with levosimendan during the first 48 h after ICU admission in order to treat neurogenic stress-induced cardiomyopathy (NSIC) secondary to aSAH (“Levo”) defined as an elevation of circulating troponin. We matched with a 1:2 ratio all Levo patients to patients that were not treated with levosimendan. Matching was performed on age, World Federation of Neurosurgical Societies (WFNS) grade, Fischer grade, need for external ventricular derivation (EVD) and severity assessed by Simplified Acute Physiology Score II (SAPS II). We excluded patients who died or had treatment withholding or withdrawal within the first four days following the bleeding. Levosimendan was administered intravenously as a continuous infusion over 24 h at a 0.1 μg/kg/min dose.

The primary endpoint of the study was the rate of occurrence of CVS defined as vessel narrowing higher than 50% requiring angioplasty and/or associated with DCI. DCI was defined according to Vergouwen et al. [5] Secondary outcomes included occurrence of DCI, 3-month modified Rankin scale (mRS) and variation of the cerebral arteries diameters between days 5–7 and day 1.

Between January 2018 and May 2020, a total of 652 patients were admitted for aSAH in our institution, and 18 Levo patients and 36 controls could be included in the study. The characteristics and outcomes of the 54 included patients are presented in Table 1.
Considering the primary outcome of the study, the rate of CVS was lower in patients treated with levosimendan compared to the control group (5/18 (28%) vs 24/36 (67%), respectively, \( p = 0.009 \)).

There was no brain infarction related to CVS in the Levo group, but the difference with the control did not reach statistical significance (0/18 (0%) vs 7/36 (19%) for the Levo vs control groups, respectively, \( p = 0.08 \)).

There was no association between levosimendan administration and mRS at 3 months (OR 1.56 [0.43; 5.92]; \( p = 0.56 \)) neither for mortality at 3 months (OR 0.18 [0.00; 1.52]; \( p = 0.14 \)).

Regarding evolution of vessels diameters, we observed a slight increase in the median cerebral arteries diameters between D0 and D5–7 (0.03 mm (IQR [−0.01; 0.06])) in the Levo group, while cerebral arteries showed signs of vasospasm with a decreasing diameter in the control group (−0.47 mm (IQR [−0.51; −0.24])) (\( p = 0.002 \); Fig. 1). Characteristics of the patients included in this analysis did not differ.

We found an association between the administration of levosimendan and a reduced incidence of CVS. Our study presents several limitations including its retrospective and non-randomized design that may induce several biases. Cardiac impairment was more frequent in the Levo group, since NSIC is associated with occurrence of CVS [6]; this argues in favor of a positive impact of levosimendan.

Table 1 Characteristics, management and outcome for levosimendan cases and non-levosimendan controls

| Characteristics                                    | Cases (“Levo”)a (\( n = 18 \)) | Controlsb (\( n = 36 \)) | \( P \)-value | Odds-Ratio |
|----------------------------------------------------|---------------------------------|--------------------------|---------------|------------|
| **Patients characteristics**                       |                                 |                          |               |            |
| Age—median (IQR)—years                             | 61 (50–70)                      | 61.5 (52–68)             | 0.63          |            |
| Female—No (%)                                      | 15 (83.3)                       | 30 (83.3)                | 1.00          |            |
| Arterial hypertension—No (%)                       | 6 (33.3)                        | 17 (47.2)                | 0.39          |            |
| Smoking—No (%)                                     | 10 (55.6)                       | 10 (27.8)                | 0.07          |            |
| Diabetes—No (%)                                    | 0 (0)                           | 1 (2.8)                  | 1.00          |            |
| Stroke history—No (%)                              | 2 (11.1)                        | 4 (11.1)                 | 1.00          |            |
| SAPS 2—median (IQR)                                | 60 (43; 65)                     | 49 (40; 57)              | 0.15          |            |
| SOFA—median (IQR)                                  | 8 (7; 11)                       | 6 (2; 9)                 | 0.03          |            |
| **Characteristic of SAH**                          |                                 |                          |               |            |
| WFNS grade—median (IQR)                            | 5 (4; 5)                        | 4 (4; 5)                 | 0.43          |            |
| Fischer grade—median (IQR)                         | 4 (4; 4)                        | 4 (4; 4)                 | 0.48          |            |
| Coiling—No (%)                                     | 16 (88.9)                       | 32 (88.9)                | 1.00          |            |
| External ventricular drainage—No (%)               | 7 (38.9)                        | 23 (63.9)                | 0.09          |            |
| Decompressive craniectomy—No (%)                   | 2 (11.1)                        | 3 (8.3)                  | 1.00          |            |
| **Cardiac impairment**                             |                                 |                          |               |            |
| Cardiac troponin peak—median (IQR)—ng/L           | 3999 (2210; 7382)               | 821 (40; 4532)           | 0.007         |            |
| **BNP peak—median (IQR)—ng/L**                     | 1122 (762; 1718)                | 138 (65; 937)            | 0.001         |            |
| **Outcome**                                        |                                 |                          |               |            |
| Cerebral vasospasm—No (%)                          | 5 (27.8)                        | 24 (66.7)                | 0.009         | 0.199 [0.0443; 0.766] |
| Delayed cerebral ischemia—No (%)                   | 0 (0)                           | 7 (19.4)                 | 0.08          | NA         |
| MRS at 3 months \( \leq 3 \)—No (%)               | 11 (61.1)                       | 18 (50)                  | 0.57          | 1.558 [0.4331; 5.922] |
| Death at 3 months—No (%)                           | 1 (5.6)                         | 9 (25)                   | 0.14          | 0.181 [0.004; 1.516] |

Characteristics of patients are compared with Mann–Whitney U test for quantitative variables and Fisher’s exact test for categorical variables

BNP, B-type natriuretic peptide; mRS, modified Rankin score; NA, not available; SAPS2, Simplified Acute Physiology Score 2; SOFA, Sepsis-related Organ Failure Assessment; WFNS, World Federation of Neurosurgical Societies

* No missing data

b Missing data: arterial hypertension status for one patient, smoking status for one patient, diabetes status for one patient, stroke history for one patient, SOFA for six patients, type of aneurysmal securization for one patient, cardiac troponin peak for four patients, BNP peak for ten patients, use of norepinephrine for two patients
Trinh-Duc et al. Critical Care (2021) 25:396

Our results suggest that levosimendan could represent a therapeutic candidate for early prevention of cerebral vasospasm secondary to aSAH.

Acknowledgements
The authors are in debt to Prof. Etienne Gayat for his help regarding data analysis and manuscript drafting.

Authors’ contributions
ATD and MAL collected the data, analyzed the data and drafted the manuscript. AC and GBH collected the data, analyzed the data and made critical revision to the manuscript. AM analyzed the data and made critical revision to the manuscript. BGC analyzed the data, supervised the work and drafted the manuscript. All authors read and approved the final manuscript.

Funding
This study was not funded.

Availability of data and materials
Data are available upon request to corresponding author.

Declarations

Ethics approval and consent to participate
This research is a single-centre retrospective case–control study of patients admitted to the Lariboisière Hospital Surgical Intensive Care Unit (ICU) (Paris, France) for an aSAH. The study was approved by an ethics committee (Comité d'éthique de la Recherche en Anesthésie-Réanimation (CERAR) no. IRB 00010254-2018-062 and Commission Nationale Informatique et Liberté (CNIL) (no. 2173692)).

Consent for publication
All authors approved the content of the manuscript and its publication.

Competing interests
BGC was a member of an advisory board for Roche Diagnostic; AM received speaker’s honoraria from Abbott, Novartis, Orion, Roche and Servier and fees as a member of the advisory board and/or steering committee from Cardiorentis, Adrenomed, MyCartis, Neurontonik and Sphingotec. EG received research grant from Sphingotec and consultancy fees from Magnisense and Roche Diagnostics. All other authors have nothing to disclose.

Author details
1 Department of Anesthesiology and Critical Care, DMU Parabol, FHU PROMICE, APHPNord, Lariboisière Hospital, 2 rue Ambroise Paré, 75010 Paris, France. 2 INSERM U942 MASCOT, Université de Paris, Paris, France. 3 Department of Interventional Neuroradiology, Hospital Lariboisière, Paris, France. 4 EA 7334 REMES, Université de Paris, Paris, France. 5 UMR 1266, Department of Neuroradiology, GHU Paris, Université de Paris, INSERM, Paris, France.

Received: 27 September 2021 Accepted: 11 November 2021
Published online: 16 November 2021

References
1. Macdonald RL. Delayed neurological deterioration after subarachnoid haemorrhage. Nat Rev Neurol. 2014;10(1):44–58.
2. Brami J, Chousterman B, Boulouis G, Dorze ML, Majlath M, Saint-Maurice J-P, et al. Delayed Cerebral Infarction is Systematically Associated with a Cerebral Vasospasm of Large Intracranial Arteries. Neurosurgery. 2019 Sep 9;nyz340.
3. Li K, Barras CD, Chandra RV, Kok HK, Maingard JT, Carter NS, et al. A review of the management of cerebral vasospasm after aneurysmal subarachnoid hemorrhage. World Neurosurg. 2019;126:513–27.
4. Busani S, Rinaldi L, Severino C, Cobelli M, Pasetto A, Girardis M. Levosimendan in cardiac failure after subarachnoid hemorrhage. J Trauma Inj Infect Crit Care. 2010;68(5):E108–10.
5. Vergouwen MDI, Vermeulen M, van Gijn J, Rinkel GJE, Wijdicks EF, Muizelaar JP, et al. Definition of delayed cerebral ischemia after aneurysmal subarachnoid hemorrhage as an outcome event in clinical trials and observational studies: proposal of a multidisciplinary research group. Stroke. 2010;41(10):2391–5.
6. Zhang L, Zhang B, Qi S. Impact of echocardiographic wall motion abnormality and cardiac biomarker elevation on outcome after subarachnoid hemorrhage: a meta-analysis. Neurosurg Rev. 2020;43(1):59–68.

Publisher's Note
Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.