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Physiological effects of invasive ventilation with neurally adjusted ventilatory assist (NAVA) in a crossover study

Jean-Michel Liet1*, François Barrière1, Bénédicte Gaillard-Le Roux1, Pierre Bourgoin1, Arnaud Legrand1,2 and Nicolas Joram1

Abstract

Background: Neurally Adjusted Ventilatory Assist (NAVA) is a mode of assisted mechanical ventilation that delivers inspiratory pressure proportionally to the electrical activity of the diaphragm. To date, no pediatric study has focused on the effects of NAVA on hemodynamic parameters. This physiologic study with a randomized cross-over design compared hemodynamic parameters when NAVA or conventional ventilation (CV) was applied.

Methods: After a baseline period, infants received NAVA and CV in a randomized order during two consecutive 30-min periods. During the last 10 min of each period, respiratory and hemodynamic parameters were collected. No changes in PEEP, FiO2, sedation or inotropic doses were allowed during these two periods. The challenge was to keep minute volumes constant, with no changes in blood CO2 levels and in pH that may affect the results.

Results: Six infants who had undergone cardiac surgery (mean age 7.8 ± 4.1 months) were studied after parental consent. Four of them had low central venous oxygen saturation (ScvO2 < 65 %). The ventilatory settings resulted in similar minute volumes (1.7 ± 0.4 vs. 1.6 ± 0.6 ml/kg, P = 0.67) and in similar tidal volumes respectively with NAVA and with CV. There were no statistically significant differences on blood pH levels between the two modes of ventilation (7.32 ± 0.02 vs. 7.32 ± 0.04, P = 0.34). Ventilation with NAVA delivered lower peak inspiratory pressures than with CV: -32.7 % (95 % CI: -48.2 to –17.1 %, P = 0.04). With regard to hemodynamics, systolic arterial pressures were higher using NAVA: +8.4 % (95 % CI: +3.3 to +13.6 %, P = 0.03). There were no statistically significant differences on cardiac index between the two modes of ventilation. However, all children with a low baseline ScvO2 (<65 %) tended to increase their cardiac index with NAVA compared to CV: 2.03 ± 0.30 vs. 1.91 ± 0.39 L/min/m2 (median ± interquartile, P = 0.07).

Conclusions: This pilot study raises the hypothesis that NAVA could have beneficial effects on hemodynamics in children when compared to a conventional ventilatory mode that delivered identical PEEP and similar minute volumes.

Trial registration: ClinicalTrials.gov Identifier: NCT01490710. Date of registration: December 7, 2011.

Keywords: Mechanical ventilation, Cardiac index, Doppler ultrasonography, Neurally adjusted ventilatory assist, Interactive ventilatory support, Electrical activity of diaphragm, Intensive care, Children, Infants, Cardiac surgery

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Background
To better match the level of ventilator assistance to the patient’s needs, manufacturers have developed new modes that deliver a level of assistance proportional either to the patient’s inspiratory muscle effort, with proportional assist ventilation (PAV); or to the diaphragmatic electrical activity (EAdi), with neurally adjusted ventilatory assist (NAVA) [1]. For now, only NAVA is usable for infants. With this mode of mechanical ventilation, the collected electrical signal allows synchronization of ventilation to spontaneous breathing efforts of the child, as well as permitting pressure assistance proportional to the electrical signal. NAVA provides both fine synchronization of respiratory support and pressure assistance varying with the needs of the child. To our knowledge, no data have been published on the impact of NAVA on the hemodynamics in children mechanically ventilated.

The impact of mechanical ventilation on the circulatory system remains a concern in some infants following surgery for repair of congenital cardiac defects [2]. After the completion of the operation, when there is consensus regarding a good surgical result, many children nowadays are rapidly weaned and successfully separated from mechanical ventilation shortly after the procedure. However, duration of mechanical ventilation is inversely proportional to the patient’s age and directly related to the duration of cardiopulmonary bypass and complexity of the surgical procedure [3].

This physiologic study with a randomized cross-over design compared hemodynamic parameters when neurally adjusted ventilatory assist (NAVA) or conventional ventilation (CV) was applied.

Methods
This study was performed in the 12-bed PICU of the university hospital center of Nantes (France), between June 2012 and March 2013. Last year, 712 children were admitted in this unit. Among them, 182 underwent a cardiac surgery with cardiopulmonary bypass (55 % were extubated within 8 h). This trial was registered (ClinicalTrials.gov Identifier: NCT01490710). Research has been performed in accordance with the Declaration of Helsinki and has been approved by an appropriate ethics committee (Comité de Protection des Personnes Ouest III de POITIERS, Reference comité: Protocole n°11.07.20). The parents of all the children gave their written consent.

Study design: prospective randomized cross-over study
After a baseline period, children received conventional ventilation (CV) and NAVA during two consecutive periods of 30 min, in random order. This randomization was performed by contacting a server to define the order in which the two modes of ventilation were administered. There was no washout time between these two periods. During the last 10 min of each period at a steady state, hemodynamic parameters, respiratory parameters, blood gas and cardiac index were collected. No changes in PEEP, FiO₂, sedation or inotropic doses were allowed during these two periods. The study protocol is depicted in Fig. 1.

Participants
Children hospitalized for postoperative care after cardiac surgery were considered for inclusion. Inclusion criteria were: admission weight >5 kg; invasive ventilatory support ongoing; NAVA ventilation available; sedation status allowing the NAVA ventilation functioning. Non inclusion criteria were: parental refusal; oesophageal disease prohibiting gastric tube; brain damage conflicting with spontaneous ventilation; clinical instability requiring treatment and/or management inconsistent with research.

Intervention
Children were ventilated with a Servo-i ventilator (Maquet Critical Care, Solna, Sweden). During a baseline period, conventional ventilation was applied and adjusted according to blood gases. After the baseline period in which NAVA probe and Doppler monitoring were introduced,
all children received conventional ventilation (CV) and NAVA during two consecutive periods of 30 min, in random order. The ventilation settings were supposed to provide the same minute volumes.

**Conventional ventilation (CV)**
With the goal of providing the lowest peak pressures, we used the Volume Control Intermittent Mandatory Ventilation (VC-IMV). Ventilatory parameters were as follows: tidal volume (between 5 and 8 ml/kg), respiratory rate (between 20 and 30 cycles/min). Pressure Support was set at 12 cmH2O above the PEEP. A flow pneumatic trigger was used to synchronize the support to spontaneous breaths. The flow pattern during the volume control ventilation was constant flow. The inspiratory time was initially set at 0.5 s and adjusted according to the inspiratory flow waveforms if needed. The respiratory rate of VC-IMV was set at a high level (between 20 and 30 cycles/min), not allowing many spontaneous cycles. The CV settings were adjusted during the baseline period with no changes allowed during the study period.

**Neurally Adjusted Ventilatory Assist (NAVA)**
During NAVA, the ventilator delivers a pressure proportional to the electrical activity of the diaphragm (EAdi). The settings in NAVA mode included PEEP, FiO2 and level of NAVA assistance. The NAVA level was set to 1 cmH2O/μV. We had previously observed that children ventilated with NAVA usually adjust their EAdi to maintain a normal blood pH by preserving physiologic tidal volumes and minute volumes. This is no longer true when children are highly sedated. Backup ventilation was Pressure Control set at 12 cmH2O above PEEP in case of failure EAdi signal detection. During NAVA, the positive pressure is triggered when the deflection in the EAdi curve exceeds 0.5 μV, and is cycled-off when the EAdi drops to 70 % of its peak value.

**Outcomes measurements**

**Respiratory measurements**
Peak inspiratory pressures (PIP), PEEP, FiO2. Tidal volume (VT), Minute volume, EADI max. Respiratory rate, were recorded for all infants during the last 10 min of each period. Measurements of each parameter were repeated at steady state a total of six times to guarantee reliability and reproducibility. We collected the most representative values rather than gathering data at constant interval because of the variability of the respiratory parameters in NAVA. Hemodynamic and respiratory parameters were collected always in the same time. Respiratory system compliance (Crs) according to the ventilation mode was estimated by the ratio VT/(PIP-PEEP) in ml/kg.cmH2O.

**Biological measurements**
Cardiac index (CI, L/min.m2) was assessed by transesophageal Doppler ultrasonography (CardioQP, Gamida, France) [4]. Other hemodynamic parameters included heart rate, arterial pressures (by means of an arterial catheter placed before the surgery), and central venous pressure (by means of a central venous catheter inserted into the right internal jugular vein). Cerebral Near Infrared Spectroscopy (NIRS) was measured with a non invasive optical monitor of regional cerebral oxygen saturation (Invos Oximeter, Covidien, Boulder, CO). The pulse oximeter perfusion index was provided by the monitors (Intellivue MP70 monitor, Philips Medical Systems) [5]. Hemodynamic parameters were collected always in the same time than respiratory parameters. Left cardiac work index [6] was calculated as: LCWI (kg.min/m2) = CI × MAP × 0.0136, with CI for cardiac index and MAP for mean arterial pressure.

**Results**
The study was concluded after one year. Technical problems were greater than expected: a measure of cardiac output sufficiently accurate, and especially stable in time, was very difficult to reach in children who already had a stomach tube for the NAVA ventilation. We asked the research office of our institution for continuing recruiting more patients. In view of the technical difficulties we had, this request was not accepted.

Nine children were randomized. The quality of the surgical repair was always checked by a cardiologist prior to inclusion in the study. Analysis was performed on only six of them. We were unable to start NAVA on one child due to a lack of EAdi signal capture. Thereafter, when the NAVA signal was not stable enough, children were not included. In two other children we were unable to record a Doppler signal likely due to interference between the
CardioQP probe and the EAdi probe both intraoesophageal placed. Patient characteristics are provided in Table 1. The mean (± SD) age was 7.8 ± 4.1 months, and weight was 6.7 ± 1.2 kg. Four children had low central venous saturation (ScvO2 < 65 %) at the baseline period: the children 1, 2, 3, 6 in the Table 1.

Respiratory parameters, hemodynamic parameters, and blood gases are provided in Table 2. The ventilatory settings resulted in similar minute volumes and in similar tidal volumes in NAVA compared to CV. These similar minute volumes in each periods of the study were supposed when the study was designed. Nevertheless, significant lower peak inspiratory pressures (PIP) were observed during NAVA compared to CV. Apparent respiratory system compliance improved with NAVA in all children. We observed higher systolic arterial pressures during NAVA compared to CV. Pulse oximeter perfusion index were also higher during NAVA.

Cardiac index during NAVA compared to CV did not statistically differ. However, all four children with ScvO2 < 65 % tended to have higher cardiac index after 30 min of ventilation with NAVA compared to CV: 2.03 ± 0.30 vs. 1.91 ± 0.39 L/min.m² (median ± interquartile, P = 0.07). Individual values of cardiac index and ScvO2 of children with low baseline ScvO2 are shown in Fig. 2. Upper values of pulse oximeter perfusion index were also observed. Turning now to left cardiac work index, the increase is +12.4 % (95 % CI: +3.8 % to +20.9 %, P = 0.07).

Discussion

NAVA can be used in infants receiving postoperative mechanical ventilation after congenital heart surgery, as previously reported [7–10]. Infants included in this study all weighed less than 10 kg and had been operated less than 24 h before. To deliver a same tidal volume, NAVA required lower inspiratory pressures than conventional ventilation and had some beneficial effects on hemodynamics. Several data validate that the children were not over assisted when ventilated in CV: the delivered volumes, EAdi max, PCO2 and pH are similar in CV compared to NAVA. We suggest that lower EAdi peaks observed in CV are related to the normalization of the blood pH by CV ventilation. If children were over-assisted in CV, EAdi would have been missing, and minute volume would have been high, with a blood alkalosis.

In a large randomized controlled trial where NAVA was used as a primary mode of ventilation, lower peak inspiratory pressures were found in the NAVA group [11]. It looks as if gas volume delivery, and probably alveolar expansion, required less pressure. This could be enhanced by an improvement in patient–ventilator interactions with a greater respiratory variability as reported in previous studies [12–16].

The effectiveness of this mode of mechanical ventilation was associated with some beneficial effects on the hemodynamics. Systolic arterial pressures and pulse oximeter perfusion index were significantly higher in NAVA. This pulse oximeter perfusion index provides a monitoring of illness severity in neonates [5]. In fact, we observed that the children with a low cardiac index had also a low perfusion index. This index was significantly higher when the children were ventilated with NAVA compared to CV.

The overall values of the cardiac index and of the central venous oxygen saturation during NAVA compared

| Child | Gender | Age (mo) | Weight (kg) | Cardiopathy | Cardiopulmonary bypass duration (min) | Time between PICU entry and study (hours) | Baseline cardiac index (L/min.m²) | Milrinone (mcg/kg/min) | Adrenaline (mcg/kg/min) | ScvO2 (%) | Perfusion index (%) | Random order |
|-------|--------|----------|-------------|-------------|--------------------------------------|-----------------------------------------|-------------------------------|-----------------------|-----------------------|-----------|-------------------|--------------|
| 1     | M      | 3        | 5.7         | AVSD        | 96                                   | 20                                      | 2.00                          | 0.9                   | 0                     | 64        | 0.31              | CV then NAVA |
| 2     | M      | 13       | 8.9         | VSD with pulmonary stenosis tetralogy of Fallot | 83                                   | 7                                       | 2.85                          | 0.5                   | 0                     | 63        | 3.65              | NAVA then CV |
| 3     | M      | 4        | 7.0         | atrial septal defect coarctation of the aorta | 130                                  | 20                                      | 1.93                          | 0.3                   | 0                     | 52        | 0.60              | CV then NAVA |
| 4     | F      | 10       | 6.0         | pulmonary atresia with VSD | 48                                   | 4                                       | 2.60                          | 0.5                   | 0                     | 82        | 3.62              | CV then NAVA |
| 5     | F      | 6        | 6.0         |             | 0                                    | 5                                       | 2.75                          | 0                     | 0                     | 68        | 1.90              | NAVA then CV |
| 6     | F      | 11       | 7.0         |             | 130                                  | 7                                       | 1.85                          | 0.5                   | 0.05                  | 52        | 0.87              | NAVA then CV |

All children were receiving morphine 0.5 mg per kilo per day. All, except the 2 first, were receiving midazolam 40 micrograms per kilo per hour

AVSD atrioventricular septal defect, VSD ventricular septal defect, NAVA neurally adjusted ventilatory assist, CV conventional ventilation, ScvO2 central venous oxygen saturation
Table 2 Respiratory parameters, hemodynamics and biological data

| Respiratory parameters       | CV       | NAVA     | NAVA versus CV | P       |
|-----------------------------|----------|----------|----------------|---------|
| Respiratory rate (/min)     | 30 (7)   | 35 (13)  | +26.4 (-3.6, +56.4) | 0.17    |
| VT (ml/kg)                  | 6.6 (0.7)| 6.9 (0.3)| +5.3 (-5.5, +16.1) | 0.46    |
| Minute volume (L/min)       | 1.7 (0.4)| 1.6 (0.6)| +5.4 (-4.3, +15.1) | 0.67    |
| PIP (cm H₂O)                | 21 (6)   | 11 (4)   | -32.7 (-48.2, -17.1) | 0.04    |
| PEEP (cm H₂O)              | 4 (2)    | 4 (2)    | -                | (a)     |
| EAdi max (microVolt)        | 3.9 (3.8)| 6.3 (1.4)| +61.2 (-4.6, +126.9) | 0.34    |
| SpO₂ (%)                    | 98 (3)   | 96 (3)   | -1.2 (-2.8, +0.5)  | 0.14    |
| FiO₂                        | 30 (4)   | 30 (4)   | -                | (a)     |
| Crs (ml/kg.cm H₂O)          | 0.37 (0.19) | 0.87 (0.32) | +98.4 (+43.8, +153.0) | 0.04    |

Hemodynamic parameters

| Parameter                        | CV       | NAVA     | NAVA versus CV | P       |
|----------------------------------|----------|----------|----------------|---------|
| Heart rate                       | 156 (15) | 156 (22) | +2.1 (-0.7, +4.5) | 0.17    |
| Systolic arterial pressure (mmHg)| 93 (6)   | 99 (13)  | +8.4 (+3.3, +13.6) | 0.03    |
| Diastolic arterial pressure (mmHg)| 54 (12) | 57 (6)   | +3.6 (-3.0, +10.1) | 0.46    |
| Central venous pressure (mmHg)   | 11 (5)   | 10 (5)   | +3.9 (-5.3, +13.1) | 0.92    |
| Cerebral NIRS (%)                | 62 (5)   | 61 (3)   | +1.6 (-2.6, +5.8)  | 0.34    |
| Cardiac index (L/min.m²)         | 2.33 (0.84) | 2.26 (0.70) | +1.4 (-3.4, +6.2) | 0.17    |
| Pulse oximeter perfusion index (%)| 1.50 (2.45) | 1.78 (2.29) | +18.8 (+3.0, +34.7) | 0.04    |

Venous blood gases

| Parameter | Value (interquartile) | Value (interquartile) | Difference (95% CI) | P |
|-----------|-----------------------|-----------------------|---------------------|---|
| pH        | 7.32 (0.04)           | 7.32 (0.02)           | -0.1 (-0.4, +0.2)   | 0.34 |
| PCO₂ (mm Hg) | 47.3 (5.1)            | 45.5 (8.1)            | +0.6 (-5.1, +6.3)   | 0.50 |
| ScvO₂ (%) | 60.1 (20.9)           | 58.4 (15.4)           | +3.3 (-5.7, +12.2)  | 0.60 |

Measurements in NAVA and CV are reported as median (interquartile). Variations between NAVA versus CV are reported as mean percentage (95% CI). Statistical analyses between NAVA versus CV were performed by a Wilcoxon test for paired samples.

(a) No changes in PEEP, FiO₂, neither in sedation or inotropic doses were allowed during these study periods.

NAVA = Neurally adjusted Ventilatory Assist, CV = Conventional Ventilation, VT = tidal volume, PIP = peak inspiratory pressure, Crs = respiratory system compliance, EAdi = electrical activity of the diaphragm, ScvO₂ = central venous oxygen saturation.

Fig. 2 Cardiac index and ScvO₂ of the low baseline ScvO₂ infants, according to the ventilation mode. White bars are cardiac index when CV is applied; back bars are cardiac index when NAVA is applied. The triangles above the bars indicate the values of the corresponding ScvO₂ (%). ScvO₂ = central venous oxygen saturation, NAVA = Neurally adjusted Ventilatory Assist, CV = Conventional Ventilation.
to CV were not significantly higher, but there are possibly no reasons to observe an increase in these two parameters when tissue perfusion is already effective. And actually, the four children with a low ScvO₂ trend to have a higher when tissue perfusion is already effective. And actually, no reasons to observe an increase in these two parameters for critically ill children [21, 22]. This bed-side device for cardiac output measurement is minimally invasive, and provides better results in monitoring the slight variations rather than in measuring absolute values. As we were performing a cross-over study, we were interested primarily by changes. Finally, only comparison with CV-SIMV with relatively high preset respiratory rate preventing many spontaneous cycles was made, not with other ventilation strategies.

Conclusion

In conclusion, this cross-over study provides new data on the NAVA ventilation. This pilot study raises the hypothesis that a ventilatory assistance with NAVA could provide improvements in hemodynamics when compared to a conventional ventilatory mode that delivered identical PEEP and similar minute volumes. Thus, because minute volumes were not different between the two modes of ventilation, hemodynamic effects are linked to NAVA mode itself and NOT to incorrect settings in conventional ventilation. Further studies with larger population are needed to confirm these promising results.

Abbreviations

AVSD: Atrioventricular septal defect; CI: Cardiac index; Crs: Respiratory system compliance; CV: Conventional ventilation; EAdi: Diaphragmatic electrical activity; LCWI: Left cardiac work index; NAVA: Neurally adjusted ventilatory assist; NIRS: Cerebral near infrared spectroscopy; PAV: Proportional assist ventilation; PIP: Peak inspiratory pressures; ScvO₂: Central venous oxygen saturation; VC-IMV: Volume control intermittent mandatory ventilation; VSD: Ventricular septal defect; VT: Tidal volume

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Availability of data and materials

Data supporting our findings can be found in additional supporting file (excel xls spreadsheet).

Authors’ contributions

AI was responsible for the data collection, ethical approval and data cleaning. JML, NJ, PB and BGL jointly were responsible for conceptualizing the study, the data analysis, drafting of the paper and approval of the final manuscript. JML is the corresponding author. All authors read and approved the final manuscript.

Competing interests

The authors declare that they have no competing interests.

Consent for publication

The parents of all the children gave their written consent to publish.

Ethics approval and consent to participate

Research has been performed in accordance with the Declaration of Helsinki and has been approved by an appropriate ethics committee (Comité de Protection des Personnes Ouest III de POITIERS, Référence comité: Protocole n° 11.07.20) approved the protocol. The parents of all the children gave their written consent to participate.
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