

Factors affecting the use of neurally adjusted ventilatory assist in the adult critical care unit: a clinician survey

Daniel Hadfield,^{1,2} Louise Rose,^{3,4} Fiona Reid,⁵ Victoria Cornelius,⁶ Nicholas Hart,^{7,8} Clare Finney,¹ Bethany Penhaligon,¹ Clare Harris,¹ Sian Saha,¹ Harriet Noble,¹ John Smith,¹ Philip Anthony Hopkins,¹ Gerrard Francis Rafferty²

To cite: Hadfield D, Rose L, Reid F, *et al*. Factors affecting the use of neurally adjusted ventilatory assist in the adult critical care unit: a clinician survey. *BMJ Open Res* 2020;7:e000783. doi:10.1136/bmjresp-2020-000783

PAH and GFR contributed equally.

This study was presented at the 2019 European Society of Intensive Care Medicine congress in Berlin, Germany.

Received 24 September 2020
Revised 11 November 2020
Accepted 12 November 2020

ABSTRACT

Background Neurally adjusted ventilatory assist (NAVA) involves an intricate interaction between patient, clinician and technology. To improve our understanding of this complex intervention and to inform future trials, this survey aimed to examine clinician attitudes, beliefs and barriers to NAVA use in critically ill adults within an institution with significant NAVA experience.

Methods A survey of nurses, doctors and physiotherapists in four Intensive Care Units (ICUs) of one UK university-affiliated hospital (75 NAVA equipped beds). The survey consisted of 39 mixed open and structured questions. The hospital had 8 years of NAVA experience prior to the survey.

Results Of 466 distributed questionnaires, 301 (64.6%) were returned from 236 nurses (78.4%), 53 doctors (17.6%) and 12 physiotherapists (4.0%). Overall, 207/294 (70.4%) reported clinical experience. Most agreed that NAVA was safe (136/177, 76.8%) and clinically effective (99/176, 56.3%) and most perceived 'improved synchrony', 'improved comfort' and 'monitoring the diaphragm' to be key advantages of NAVA. 'Technical issues' (129/189, 68.3%) and 'NAVA signal problems' (94/180, 52.2%) were the most cited clinical disadvantage and cause of mode cross-over to Pressure Support Ventilation (PSV), respectively. Most perceived NAVA to be more difficult to use than PSV (105/174, 60.3%), although results were mixed when compared across different tasks. More participants preferred PSV to NAVA for initiating ventilator weaning (93/171 (54.4%) vs 29/171 (17.0%)). A key barrier to use and a consistent theme throughout was 'low confidence' in relation to NAVA use.

Conclusions In addition to broad clinician support for NAVA, this survey describes technical concerns, low confidence and a perception of difficulty above that associated with PSV. In this context, high-quality training and usage algorithms are critically important to the design and of future trials, to clinician acceptance and to the clinical implementation and future success of NAVA.

BACKGROUND

Neurally adjusted ventilatory assist (NAVA) uses the electrical activity of the diaphragm (Edi), obtained via a specialised nasogastric feeding catheter (Getinge, Solna, Sweden), as a measure of neural respiratory drive to

Key messages

- ▶ What are the factors affecting the use of neurally adjusted ventilatory assist (NAVA) in the adult critical care unit?
- ▶ Despite broad clinician support for NAVA, technical concerns, low staff confidence and a perception of technical difficulty are relevant factors that affect the clinical use and adoption of this technology.
- ▶ This is the first study to describe the lived experience of clinicians using NAVA, describing factors that are relevant both to future research trials and general clinical implementation.

control the delivery of inspiratory support by a mechanical ventilator (MV).¹ In 12 years of clinical use, numerous clinical studies have suggested important physiological benefits^{2,3} with recent trials suggesting reduced weaning time⁴ and increased ventilator-free days.^{5,6} Despite this, no trial has yet definitively demonstrated improved patient outcomes and NAVA has not been widely adopted into clinical care.

The implementation of any new technology-based healthcare treatment is complex. Evidence from clinical trials is important, but the success of trials and the subsequent implementation into practice is dependent on a number of other factors. In the case of NAVA, such factors potentially include contextual issues (eg, prevalent culture, cost and access to the technology), human issues (eg, user skill level, training requirements, beliefs and attitudes) and technological issues (eg, performance and limitations of the technology). Feasibility studies are recommended to investigate such factors prior to the conduct of large, resource intensive randomised controlled trials (RCTs), to optimise efficiency and chances of success.⁷

In a recently published trial, we demonstrated the feasibility of evaluating NAVA



© Author(s) (or their employer(s)) 2020. Re-use permitted under CC BY. Published by BMJ.

For numbered affiliations see end of article.

Correspondence to

Dr Daniel Hadfield;
daniel.hadfield@nhs.net



in an RCT compared with PSV for patients at risk of prolonged ventilation support, with adequate compliance to the assigned ventilatory mode over prolonged durations.⁵ Despite satisfactory mode compliance, some non-adoption and mode cross-over from NAVA to PSV were still observed. To improve our understanding of these cross-over events and wider factors affecting the use of NAVA, we developed and conducted a survey aimed at exploring clinician attitudes, beliefs, perceived barriers and other factors that potentially affect the implementation of NAVA in critically ill adult patients. These data are critical to the design and interpretation of subsequent trials and may help to explain the slow progress towards efficacy and effectiveness trials and clinical adoption of NAVA worldwide.

METHODS

An anonymous, self-administered cross-sectional survey of nurses, doctors and physiotherapists was undertaken in four ICUs (surgical, general medical, neuro/trauma and liver) totalling 75 beds at a university affiliated hospital in London, UK. The survey was conducted alongside a randomised controlled feasibility trial (NCT01826890) comparing NAVA to PSV in adults at risk of prolonged ventilation. The purpose of the study, assurance of anonymity and voluntary nature of the survey was outlined in the participant information sheet. Informed consent was assumed on completion of the questionnaire.

Survey development and testing

Questionnaire items generated following an evidence review and previous qualitative work,⁸ were refined in three phases through (1) expert review, (2) cognitive interviews and (3) pilot survey distribution. Items were then revised and reduced by an expert team composed of an ICU nurse, a consultant intensivist and an academic physiologist. Questions were formatted into four domains: (1) baseline demographics, NAVA experience and training, (2) beliefs, attitudes and barriers, (3) perceived advantages and disadvantages, and (4) views on NAVA research. Paper-based and web-based (SurveyMonkey) surveys were designed and trialled by a senior statistician and two ICU clinical research nurses in addition to the core expert panel, assessing the clarity, acceptability, time to completion and face validity of the instrument.

Following the process described above, individual cognitive interviews were conducted with three consultant level intensivists, one senior physiotherapist, two senior ICU nurses and two junior ICU nurses. The cognitive interviewing technique is a qualitative method designed to investigate whether a survey question achieves its intended purpose.⁹ Interview notes were collated and used to further revise the draft survey. Finally, electronic and paper surveys were sent to five clinical ICU research nurses, one consultant physiotherapist, and one consultant intensivist to further confirm the time to complete, face and content validity.

Context

NAVA became a treatment option at the study site in 2008 and the RCT ran between 2012 and 2018. During the trial period, NAVA was used in an estimated 4%–7% of all ventilated patients admitted to the ICU, approximately 60–100 patients per year, 10 of which were recruited to the trial. PSV remained the predominant choice for ventilatory weaning, reflecting ventilatory weaning practice worldwide.¹⁰ NAVA was applied mostly in patients with risk factors for prolonged weaning (study inclusion criteria), but also in a range of clinical circumstances outside of the trial. Treatment plans were made during daily, physician-led, multidisciplinary ward rounds; changes to ventilation (eg, mode or support-level changes) were made by both nurses and doctors. Physiotherapists were involved in assessment, planning and adjustment of ventilation in complex and difficult to wean patients. NAVA education was offered to all staff during scheduled, general training events and individual sessions were provided as needed. Although not specified in the survey instrument, due to the almost exclusive use of NAVA in intubated patients, it was implicit that the survey was addressing issues around invasive ventilation.

Sampling frame

The sample comprised staff with responsibility for the management of MV at the study site. A database containing contact details, professions and grades was accessed with appropriate local permissions. The target population consisted of 365 nurses, 89 doctors and 12 physiotherapists.

Instrument administration

The electronic questionnaire (online supplemental file 1) was administered to all eligible staff on the 15 May 2017, towards the end of the clinical trial that ended in January 2018. Non-responders received up to three additional reminder emails, with the final reminder sent on 12 July 2017. The final response was received on 24 July 2017 when the survey was closed. Paper questionnaires were delivered to all ICU staff who had not completed an electronic survey; participants were checked off a master list to avoid duplication. All questionnaires were self-administered. No incentive was offered to encourage participation.

Statistics and analysis

As the project was primarily descriptive in nature a power calculation was not appropriate as hypothesis testing was not a main aim. Statistical analyses were performed using GraphPad Prism (V.8.0 for Windows, GraphPad Software, San Diego, California, USA).¹¹ The response rate was calculated as the total number of returned surveys with answered questions, including those that were partially completed, divided by the total number of eligible participants identified at study start. Descriptive statistics are

presented, including count, percentage and 95% confidence intervals (CIs), or medians with inter quartile ranges (IQRs) as appropriate. Five or seven-point Likert scales¹² and multiple options questions were used to assess attitudes, opinions and agreement with statements. Ordinal Likert data were converted to ranks; descriptive data are presented as median ranks, and Spearman's rank-order correlation was used to assess association. Distributions of categorical variables were compared using the χ^2 test or Fisher exact test. Free-text 'other' options were included in all multiple-options questions from which recurring themes were identified.

Patient and public involvement

Patients or the public were not involved in the design of the study.

RESULTS

Of the 466 questionnaires distributed, 301 (64.6%) were returned including responses from 236 nurses (78.4%), 53 doctors (17.6%) and 12 physiotherapists (4.0%). The response rate expressed as a proportion of the eligible staff was 64.7% for nurses, 59.6% for doctors and 100.0% for physiotherapists. Responses were obtained across all levels of seniority; most participants were junior grade (72.7%), aged 25–34 years (56.0%), and with less than 3 years of ICU experience (60.8%) (table 1), which reflected the general profile of ICU staff.

Of all participants, 157/295 (53.2%) had received NAVA training, 207/294 (70.4%) reported clinical exposure to NAVA (at least one patient treated where NAVA was used), 163/291 (56.0%) indicated that they were familiar (slightly, moderately, very or extremely) with evidence supporting NAVA use in weaning, 269/292 (92.1%) with evidence supporting PSV in weaning, and 289/294 (98.3%) with risk factors for prolonged MV (table 1 and online supplemental figure S1). Of those reporting clinical exposure to NAVA, 125/193 (64.8%) indicated use of Edi monitoring (online supplemental figure S2) and 110/180 (61.1%) had participated in the concurrent clinical trial. Of those who had received NAVA training, the majority (140/157, 89.2%) received individual bedside training from local staff. Doctors were more likely than nurses or physiotherapists to have used NAVA clinically, in greater than five patients and recently (within the month prior to the survey), perhaps reflecting their role in medical management of multiple patients compared with nursing management of individual patients (table 1). Doctors were also more likely to indicate familiarity with NAVA evidence compared with nurses or physiotherapists ($p=0.006$). Overall, very few staff had used NAVA in the week (5/185, 2.6%) or month (16/185, 8.3%) prior to the survey, but most had used NAVA within the past 6 months (113/185, 61.1%) (table 1).

Only those reporting clinical exposure to NAVA (as opposed to training experience) were asked to proceed to subsequent questions.

Most participants agreed or strongly agreed that NAVA was safe (136/177, 76.8%), and clinically effective (99/176, 56.3%), that diaphragm monitoring was clinically effective (101/172, 58.7%), and that ventilator dyssynchrony was a clinically significant issue (133/174, 60.3%) (table 2 and online supplemental figure S3). When asked about general feelings, 94/179 (52.5%) indicated ambivalence, 59/179 (33.0%) 'liked' (slightly, moderately or strongly) and 26/179 (14.5%) 'disliked' (slightly, moderately or strongly) NAVA, suggesting good equipoise for future randomised trials (online supplemental figure S4). In comparison to PSV, similar numbers of participants perceived better (slightly, moderately or significantly) (43/190, 22.6%), worse (slightly, moderately or significantly) (37/190, 19.5%), or equivalent (49/190, 25.8%) NAVA clinical performance, with 61/190 (32.1%) indicating 'don't know', and with no significant differences between professional groups (online supplemental figure S5). More participants indicated a preference for PSV (93/171, 54.4%) compared with NAVA (29/171, 17.0%) for initiation of ventilatory weaning, although a large proportion (49/171, 27.1%) were ambivalent.

Overall most participants indicated that NAVA was more difficult than PSV (105/174, 60.3%) (table 2), and considered it was harder (much/moderately/slightly) rather than easier (much/moderately/slightly) in relation to 'set up and start' (85/116 (73.3%) vs 9/116 (7.8%)), 'ventilation' (49/121 (40.5%) vs 26/121 (21.5%)) and 'reliability' (55/110 (50.0%) vs 23/110 (20.9%)) (figure 1 and online supplemental figure S6). More participants indicated that NAVA was easier than PSV, rather than harder, in relation to 'synchrony' (79/131 (60.3%) vs 34/131 (26.0%)), 'patient comfort' (75/131 (57.3%) vs 22/131 (16.8%)) and 'weaning' (62/126 (49.2%) vs 32/126 (25.4%)). The most frequent response was 'no difference' between the two modes in relation to 'lung protection' (44/120, 36.7%) and 'oxygenation' (92/123, 74.8%). NAVA was not perceived to increase workload by 85/188 (45.2%) of participants, while 74/188 (39.4%) perceived either a slight, moderate, large or substantial workload increase, and with no differences between professional groups (table 2 and online supplemental figure S7).

Participants reported low confidence in performing eight NAVA related tasks (median ranks range from 1 to 2 across all tasks); those with greater NAVA exposure in terms of patients treated, were more likely to indicate greater confidence ($r=0.562$, 95% CI 0.453 to 0.654) (figure 2 and online supplemental figure S8). Level of NAVA exposure was related to years of ICU experience; participants with more years ICU experience were more likely to have used NAVA in a greater number of patients ($r=0.531$, 95% CI 0.442 to 0.610). Doctors expressed greater confidence compared with nurses, who were consistently more likely to reply, 'not at all confident'.

**Table 1** Survey distribution, response, demographics and experience of NAVA

Characteristic	All staff	Nurses	Doctors	Physiotherapists
Years in KCH ICU				
<1	96/301 (31.9)	70/236 (30.0)	18/53 (35.9)	7/12 (58.3)
1–3	87/301 (28.9)	71/236 (30.1)	15/53 (28.3)	1/12 (8.3)
3–5	49/301 (16.3)	41/236 (17.4)	5/53 (9.4)	3/12 (25.0)
>5	69/301 (22.9)	54/236 (22.9)	14/53 (26.4)	1/12 (8.3)
Seniority*				
Junior	193/297 (65.0)	171/235 (72.8)	19/50 (38.0)	3/12 (25.0)
Middle grade	54/297 (18.1)	39/235 (16.6)	11/50 (22.0)	4/12 (33.3)
Senior	50/297 (16.8)	25/235 (10.6)	20/50 (40.0)	5/12 (41.7)
Familiarity with key concepts*†				
Risk factors for PMV	289/294 (98.3)	225/229 (98.3)	52/53 (98.1)	12/12 (100.0)
Evidence for PSV	269/292 (92.1)	211/227 (93.0)	46/53 (86.8)	12/12 (100.0)
Evidence for NAVA	163/291 (56.0)	120/226 (53.1)	36/53 (67.9)	7/12 (58.3)
Training and experience*				
NAVA trained	157/295 (53.2)	124/230 (53.9)	28/53 (52.8)	5/12 (41.6)
Any NAVA use	207/294 (70.4)	153/229 (66.8)	47/53 (88.7)	7/12 (58.3)
Approximate no of patients treated where NAVA was used*				
<5	143/287 (49.8)	115/223 (51.6)	24/53 (45.3)	4/11 (36.4)
5–20	62/287 (21.6)	45/223 (20.2)	15/53 (28.3)	2/11 (18.2)
>20	16/287 (5.6)	5/223 (2.2)	10/53 (18.9)	1/11 (9.1)
None	66/287 (23.0)	58/223 (26.0)	4/53 (7.6)	4/11 (36.4)
Frequency of NAVA use *‡				
Use in past week?	5/193 (2.6)	2/141 (1.4)	3/45 (6.7)	0/12 (0.0)
Use in past month?	16/193 (8.3)	6/141 (4.3)	10/45 (22.2)	0/12 (0.0)
Use >1 month prior	164/193 (85.0)	125/141 (64.8)	32/45 (71.1)	7/12 (58.3)

Data are number (%). Nurse and physiotherapist grade according to UK Agenda for Change (AFC) grades: Junior=AFC band 5; Middle=AFC band 6; Senior=AFC bands seven and above.

Doctors grades according to UK medical role classifications. Junior=foundation year 1/2 or equivalent; Middle=junior or senior middle grades (eg, specialty trainees, clinical fellows, senior house officers); senior=consultant level.

Bold/italic values represent the overall response rate, including all professions

*Numbers do not total 301 because not all participants answered every question. Denominators provided.

†Assessed on a five-point Likert scale: Not at all, slightly, moderately, very and extremely familiar. The presented data are the combined proportions answering either slightly, moderately, very or extremely familiar. See online supplemental file 2 for response numbers in each category.

‡Only staff reporting clinical exposure to NAVA were asked to complete this and subsequent questions.

KCH ICU, King's College Hospital Intensive Care Unit; NAVA, neurally adjusted ventilatory assist; PMV, prolonged mechanical ventilation; PSV, pressure support ventilation.

When asked about the potential benefits of using NAVA and participants could select multiple options, the most common response (140/190, 73.7%) was 'improved synchrony' (figure 3 and online supplemental figure S9). The single most important benefit (single choice) was 'reduced time in ventilation' (57/183, 31.3%). The need to improve ventilator synchrony and weaning were also considered the most common clinical reasons for NAVA use (133/192 (69.3%) and 99/192 (51.6%), respectively). In regard to the disadvantages of NAVA (figure 3 and online supplemental figure S10), most selected 'technical issues' (129/189, 68.3%), with half (88/173, 50.9%) selecting this option as the most important

single disadvantage. The next most commonly selected disadvantage was 'difficult to adjust or regulate', further suggesting the perception of difficulty and/or complexity with this mode. There were no significant differences between nurses and doctors in their perception of clinical advantages or disadvantages.

The most commonly selected barrier to acceptance and implementation of NAVA, was 'low experience, skills or confidence' (172/183, 94.0%) (figure 3 and online supplemental figure S11), which was also selected as the single main barrier (132/171, 78.4%). Most participants believed that in their experience of NAVA, the mode was 'often switched' to PSV (56/182,

Table 2 General beliefs and attitudes towards NAVA (participants with clinical experience only)

Question and response	All staff	Nurses	Doctors	Physio
General feelings towards NAVA				
Slightly/moderately/strongly like	59/179 (33.0)	43/131 (32.8)	13/41 (31.7)	3/7 (42.9)
Ambivalent	94/179 (52.5)	68/131 (51.9)	24/41 (58.5)	2/7 (28.6)
Slightly/moderately/strongly dislike	26/179 (14.5)	20/131 (15.3)	4/41 (9.8)	2/7 (28.6)
Perceived performance compared with PSV				
Slightly/moderately/significantly better	43/190 (22.6)	26/138 (18.8)	16/45 (35.6)	1/7 (14.3)
Equivalent	49/190 (25.8)	38/138 (27.5)	8/45 (17.8)	3/7 (42.9)
Slightly/moderately/significantly worse	37/190 (19.5)	27/138 (19.6)	9/45 (20.0)	1/7 (14.3)
Don't know	61/190 (32.1)	47/138 (34.1)	12/45 (26.7)	2/7 (28.6)
NAVA is safe (agreement)				
Agree/strongly agree	136/177 (76.8)	97/129 (75.2)	32/41 (78.1)	7/7 (100.0)
Ambivalent	32/177 (18.1)	31/129 (24.0)	8/41 (19.5)	0/7 (0.0)
Disagree/strongly disagree	7/177 (4.0)	1/129 (1.0)	1/41 (2.4)	0/7 (0.0)
NAVA is clinically effective (agreement)				
Agree/strongly agree	99/176 (56.3)	70/129 (54.3)	28/40 (70.0)	1/7 (14.3)
Ambivalent	70/176 (39.8)	53/129 (41.1)	11/40 (27.5)	6/7 (85.7)
Disagree/strongly disagree	7/176 (4.0)	6/129 (4.7)	1/40 (2.5)	0/7 (0.0)
Edi monitoring is clinically effective (agreement)				
Agree/strongly agree	101/172 (58.7)	72/125 (57.4)	25/40 (62.5)	4/7 (57.1)
Ambivalent	60/172 (34.9)	44/125 (34.1)	13/40 (32.5)	3/7 (42.9)
Disagree/strongly disagree	11/172 (6.4)	9/125 (7.0)	2/40 (5.0)	0/7 (0.0)
Dysynchrony is clinically significant (agreement)				
Agree/strongly agree	133/174 (76.4)	96/127 (75.6)	31/40 (77.5)	6/7 (85.7)
Ambivalent	32/174 (18.4)	24/127 (18.9)	7/40 (17.5)	1/7 (14.3)
Disagree/strongly disagree	7/174 (4.0)	5/127 (3.9)	2/40 (5.0)	0/7 (0.0)
NAVA is more difficult than PSV (agreement)				
Agree/strongly agree	105/174 (60.3)	69/127 (54.3)	31/40 (77.5)	5/7 (71.4)
Ambivalent	40/174 (23.0)	36/127 (28.3)	3/40 (7.5)	1/7 (14.3)
Disagree/strongly disagree	29/174 (16.7)	22/127 (17.3)	6/40 (15.0)	1/7 (14.3)
Preferred mode in ventilatory weaning				
NAVA	29/171 (17.0)	21/124 (16.9)	8/40 (20.0)	0/7 (0.0)
PSV	93/171 (54.4)	64/124 (51.6)	23/40 (57.5)	6/7 (85.7)
No preference	49/171 (28.8)	39/124 (31.5)	9/40 (22.5)	1/7 (14.3)
Increased workload due to NAVA?				
None	85/188 (45.2)	66/137 (48.2)	14/44 (31.8)	5/7 (71.4)
Slight/moderate increase	69/188 (36.7)	51/137 (37.2)	16/44 (36.4)	2/7 (28.6)
Large/substantial increase	5/188 (2.7)	1/137 (1.0)	4/44 (9.1)	0/7 (0.0)
Don't know	29/188 (15.4)	19/137 (13.9)	10/44 (22.7)	0/7 (0.0)

Data are number (%). See online supplemental file 2 for response numbers in each category.

Bold/italic values represent the overall response rate, including all professions

Edi, electrical activity of the diaphragm; NAVA, neurally adjusted ventilatory assist.

30.8%), with only very few (19/182, 10.4%) believing the mode was 'switched rarely' or 'not at all' (online supplemental figure S12). The reasons most often selected for switching were 'lack of experience' and 'Edi signal problems', which were also the most

selected single main reasons for cross-over (57/168, 33.9% and 39/168, 23.2%, respectively) (figure 3 and online supplemental figure S13). When asked what single initiative would most help acceptance and use of NAVA, the majority (101/167, 60.5%) selected

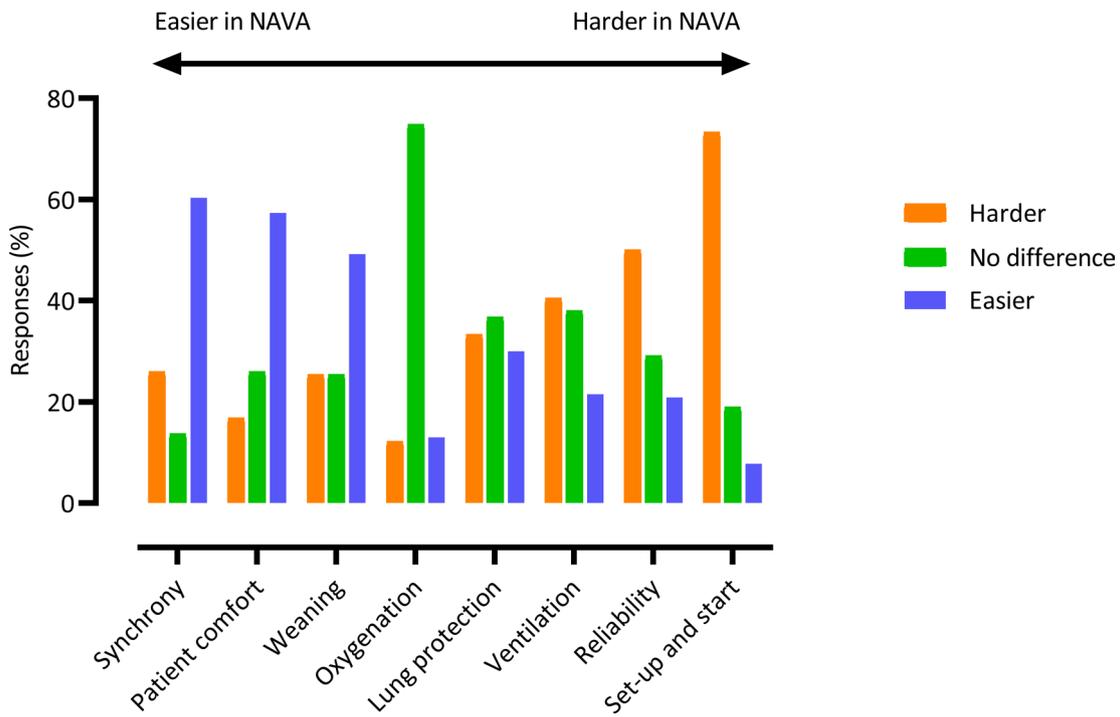


Figure 1 How easy is it to achieve the following aspects of ventilation practice when using the NAVA mode compared to the PSV mode? Assessed on a seven-point Likert scale: much, moderately, slightly easier; no difference; slightly, moderately, much harder. see online supplemental file 2 for individual response rates for each category and a breakdown of responses in each category. For ease of interpretation, activities are ordered from left to right broadly in terms of the difficulty of each activity in NAVA compared with PSV as indicated by the horizontal arrow, with oxygenation and lung protection central due to the relative equipoise of participants. NAVA, neurally adjusted ventilatory assist; PSV, pressure support ventilation.

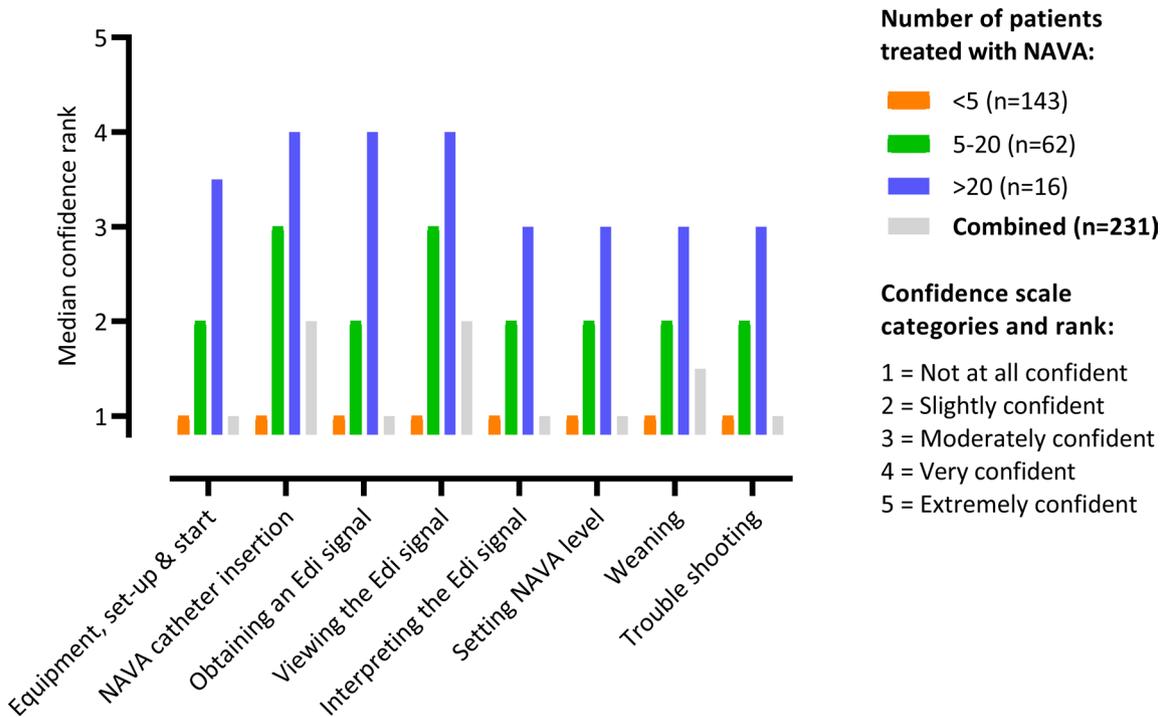


Figure 2 Confidence in performing NAVA tasks compared between staff with differing NAVA experience. Colours refer to different numbers of patients treated where NAVA was used. See online supplemental file 2 for individual response rates for each category. NAVA, neurally adjusted ventilatory assist.

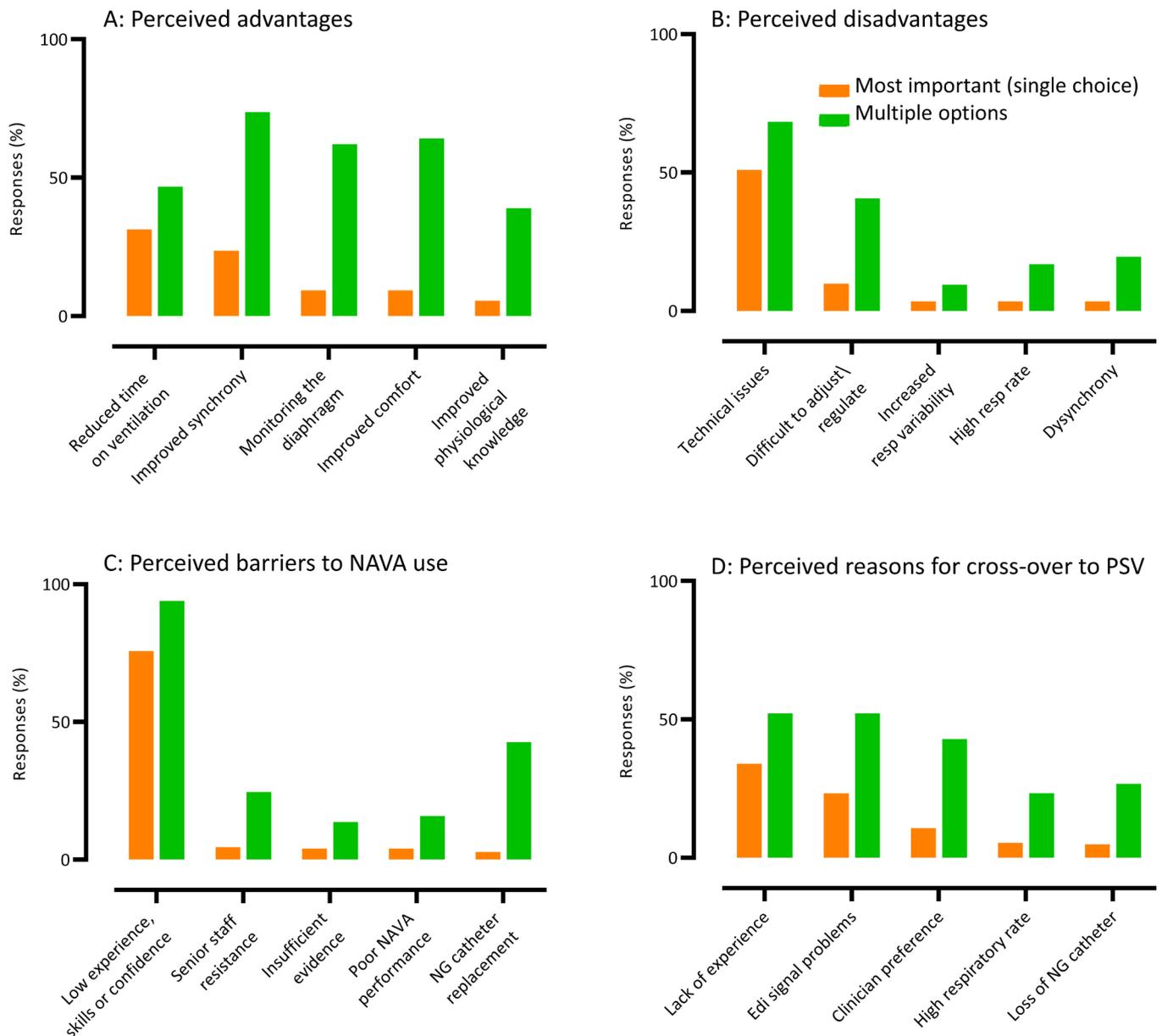


Figure 3 (A). Perceived advantages of neurally adjusted ventilatory assist (NAVA) compared with pressure support ventilation (PSV): multiple options (response rate 190) and single most important (response rate 182). (B) Perceived disadvantages of NAVA compared with PSV: multiple options (response rate 189) and single most important (response rate 173). (C) Barriers to the acceptance and implementation of NAVA: multiple options (response rate 183) and single most important (response rate 177). (D) Reasons for mode cross-over: multiple options (response rate 180) and single most important (response rate 168). Top five single most important reasons presented only. Please see online supplemental file 2 for full a full list of responses.

‘improved training and sharing of the evidence base’ (online supplemental figure S14).

When asked about research aimed at investigating the use of NAVA in prolonged MV weaning, 161/180 (89.4%) of participants were supportive, 3/180 (1.7%) were not supportive, and 16/180 (8.9%) did not know. ‘Research evidence’ was considered to be less influential on clinicians’ views on NAVA (21/174, 12.1%) compared with ‘personal, clinical experience’ (93/174, 53.5%) and ‘colleague/peer recommendation’ (54/174, 31.0%).

Out of 42 participants who made a qualitative comment, 10 (23.8%) were supportive of NAVA and/

or NAVA research, 23 (54.8%) stated that more training was needed, and 18 (42.9%) highlighted a lack of clinical experience. The following statements are representative and support other data presented above:

1. Nurse: ‘I found that when I had more exposure to NAVA, I became more confident and could see the benefits. But I found the inconsistency made it difficult to use. Sometimes I found it worked really well and other times very difficult to use.’
2. Doctor: ‘In theory NAVA sounds great. I wish I had more experience with it.’



3. Nurse: 'NAVA is a great idea, but more training needed for staff confidence and being able to troubleshoot.'

DISCUSSION

This multidisciplinary survey of 301 critical care clinicians, conducted across four adult ICUs at one academic hospital with >8 years of NAVA experience, sought to describe attitudes, beliefs, barriers and other factors relating to NAVA use, both in usual clinical practice and within clinical trials. In summary, the results indicate: (1) supportive attitudes to NAVA and belief that NAVA is safe and clinically effective with broadly equivalent performance to PSV, indicating good equipoise for future research; (2) a perception of increased difficulty and/or complexity compared with PSV; (3) a perception of low experience, skills and confidence, despite reasonable numbers of participants having clinical exposure to NAVA, and a need for improved training and (4) broad support for future NAVA research. These results provide context and feasibility data that can benefit clinical implementation of NAVA and the design of future trials. Perceived increased difficulty of using NAVA, low confidence of clinical staff and technical issues, may help to understand the current disparity between NAVA studies suggesting clinical benefit to patients, and the slow progress towards adequately powered efficacy or effectiveness trials and potential clinical adoption worldwide.

Technical difficulties with the equipment and/or Edi signal acquisition was the most often selected clinical disadvantage and reason for mode cross-over from NAVA to PSV. Cross-over is potentially a significant issue in future trials; low compliance will impact on statistical power and interpretation of results in a definitive study.¹³ Similar issues were not reported in three recent clinical trials,^{3,6,14} however, Edi signal problems were a stated cause of cross-over in 10 out of 36 participants (27.8%) in the concurrent feasibility RCT,⁵ and Edi synchrony and/or signal issues were reported in seven out of 20 (35%) patients recruited to an RCT by Di Mussi *et al.*¹⁵ Reasons for the differences between these trials remain unclear. Issues may relate to one or a combination of component failures, usability issues, or user behaviour, and these factors may be further influenced by environmental factors, such as staff to patient ratios, admission rates, availability of hardware, and varying levels of clinical expertise across a large staff group. In the NAVA mode, the Edi triggers and controls the support during the respiratory cycle, therefore acquisition of a stable, reliable signal is essential. Although trials have tested the catheter positioning technique,^{16,17} and demonstrated the reliability of the signal in different clinical situations,^{1,18-22} these studies were small (n<20), using cross-over designs with observations of limited durations—usually around 30 min. The results of this survey suggest a need to explore catheter positioning, signal acquisition and signal stability over prolonged periods and in a variety of clinical

situations where the neural respiratory drive may become unreliable.

Despite >70% of participants indicating clinical exposure to NAVA and despite >8 years of NAVA use at the study site, perceived low confidence and/or inexperience of clinicians when using NAVA is a central theme within the survey. This finding was also detected in the concurrent feasibility RCT, where clinical inexperience of the mode was considered the reason for cross-over to PSV for four (11%) of 36 participants randomised to NAVA.⁵ Infrequent application within the difficult-to-wean patient group may be relevant to these perceptions, but the results also suggest perceived difficulty and/or complexity of NAVA and technical difficulties in relation to the Edi signal, both likely to impact use and confidence with NAVA.

In the context of a clinical trial, these issues may reduce the fidelity of the NAVA intervention and threaten the internal validity of the results. They may also significantly impact clinical implementation; clinical staff are unlikely to choose a more complex, difficult and novel treatment where an easier existing option that is not perceived to cause harm is available. A comparison may be made to the exponential increase in use of nasal high flow internationally despite a lack of evidence of efficacy,²³ which may be partially due to its ease of use.

Critically, these factors are potentially modifiable. Our results suggest a need for high quality algorithms, training and standard operating procedures to mitigate these risks. Such approaches would improve the potential of demonstrating a true effect in a clinical trial by ensuring the fidelity of the intervention.^{13,24} Given the limited influence of current NAVA evidence on clinician behaviour suggested here, combined with growing evidence of clinical benefit from NAVA in preliminary studies,^{4,5} a well-designed clinical trial to demonstrate efficacy and/or effectiveness is required and is necessary to prompt behavioural change and clinical implementation.

Strengths and weaknesses

To the authors' knowledge, this is the first detailed clinician survey to assess the use of NAVA in the context of a randomised feasibility trial, with a representative range of doctor, nurse, and physiotherapist knowledge and a high response rate from all groups. The main limitation of this work relates to its conduct at a single site meaning results may not be generalisable. In particular, variability in MV management and practice across professions, institutions and healthcare systems, may affect the generalisability of these results. At the study site and in common with other UK sites,²⁵ the management of MV is collaborative, with nurses, doctors and physiotherapists requiring knowledge of ventilation modes. This may not be true in other healthcare systems outside of the UK.

As a self-report survey, various biases may have affected participant responses. Despite 44% of the participants being unfamiliar with evidence to support the use of

NAVA in weaning, participant opinions may have been biased towards a more positive view (information bias) regarding the efficacy and potential advantages of NAVA due to a largely positive body of literature. Social desirability bias may have influenced participants towards responses that were supportive of NAVA. Data relating to disadvantages and barriers are less susceptible to these biases and are, therefore, potentially of greater value.

In addition, the survey did not assess the knowledge of the participants in relation to the use of PSV or NAVA. Frequent use and greater familiarity with PSV may have affected staff attitudes towards that mode of ventilation. However, despite its relative ease of application²⁶ and ubiquitous use, it does not necessarily follow that PSV is correctly understood and applied, or that NAVA was poorly understood and applied. It may be argued that PSV requires less deliberate thought, whereas NAVA demands an understanding of underlying physiology, which may enhance the performance of the user and increase quality of the intervention.²⁷

Finally, although this study used 'the number of patients treated where NAVA was used' as a surrogate for clinical experience, the level of involvement of each participant in the management of NAVA was not assessed.

CONCLUSIONS

This study presents the experience and views of a large and diverse staff group, describing attitudes, beliefs, perceived barriers and other factors that potentially affect NAVA use in critically ill adult patients. The central perceptions of low confidence, increased difficulty and technical issues reported here, may partially explain slow progress towards clinical implementation and large trials. In this context, high-quality training, algorithms and evidence from large definitive trials are even more critical to clinician acceptance and the future success of NAVA.

Author affiliations

- ¹Critical Care Research, King's College Hospital, London, UK
- ²King's College London, Centre for Human and Applied Physiological Sciences, London, UK
- ³King's College London Florence Nightingale School of Nursing and Midwifery, London, UK
- ⁴Faculty of Nursing, Midwifery and Palliative Care, King's College, London, UK
- ⁵King's College London School of Population Health and Environmental Sciences, London, UK
- ⁶Imperial College London School of Public Health, London, UK
- ⁷Centre for Human and Applied Physiological Sciences, King's College London School of Biomedical Sciences, London, UK
- ⁸Lane Fox Respiratory Unit, Guy's and St Thomas' Hospitals NHS Trust, London, UK

Acknowledgements The authors would like to thank the Anaesthetics, Critical Care, Emergency Medicine and Trauma (ACET) research team, the London Respiratory Muscle Research Group, and all staff within the Critical Care Units at King's College Hospital for their support of this project.

Contributors Conception and design of the study: DH, PAH, GFR and NH. Data acquisition: DH, CF, BP, CH, SS, JS and HN. Analysis and interpretation of data: DH, FR, VC, LR, GFR and PAH. Manuscript preparation: DH, LR, GFR, FR and PAH. Review of manuscript: all authors. GFR and PAH contributed equally.

Funding This report is independent research supported by a National Institute for Health Research and Health Education England Clinical Doctoral Research Fellowship (DH, CDRF-2014-05-056). Additional funding received from The Moulton Foundation and Guy's and St Thomas' Biomedical Research Centre.

Disclaimer The funding sources had no role in the study design, data collection, analysis, interpretation, or manuscript preparation. The views expressed are those of the author(s) and not necessarily those of the NIHR or the Department of Health and Social Care.

Competing interests DH received funds from Maquet/Getinge to cover the travel, accommodation and registration for conferences and meetings prior to 2016.

Patient consent for publication Not required.

Ethics approval The study was approved by London Westminster ethics committee (13/L0/0012).

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement All data relevant to the study are included in the article or uploaded as online supplemental information. Data are available on reasonable request from corresponding author.

Supplemental material This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution 4.0 Unported (CC BY 4.0) license, which permits others to copy, redistribute, remix, transform and build upon this work for any purpose, provided the original work is properly cited, a link to the licence is given, and indication of whether changes were made. See: <https://creativecommons.org/licenses/by/4.0/>.

REFERENCES

- 1 Sinderby C, Navalesi P, Beck J, *et al*. Neural control of mechanical ventilation in respiratory failure. *Nat Med* 1999;5:1433–6.
- 2 Colombo D, Cammarota G, Bergamaschi V, *et al*. Physiologic response to varying levels of pressure support and neurally adjusted ventilatory assist in patients with acute respiratory failure. *Intensive Care Med* 2008;34:2010–8.
- 3 Demoule A, Clavel M, Rolland-Debord C, *et al*. Neurally adjusted ventilatory assist as an alternative to pressure support ventilation in adults: a French multicentre randomized trial. *Intensive Care Med* 2016;42:1723–32.
- 4 Liu L, Xu X, Sun Q, *et al*. Neurally adjusted ventilatory assist versus pressure support ventilation in difficult weaning: a randomized trial. *Anesthesiology* 2020;132:1482–93.
- 5 Hadfield DJ, Rose L, Reid F, *et al*. Neurally adjusted ventilatory assist versus pressure support ventilation: a randomized controlled feasibility trial performed in patients at risk of prolonged mechanical ventilation. *Crit Care* 2020;24:220.
- 6 Kacmarek RM, Villar J, Parrilla D, *et al*. Neurally adjusted ventilatory assist in acute respiratory failure: a randomized controlled trial. *Intensive Care Med* 2020. doi:10.1007/s00134-020-06181-5
- 7 Thabane L, Lancaster G. Improving the efficiency of trials using innovative pilot designs: the next phase in the conduct and reporting of pilot and feasibility studies. *Pilot Feasibility Stud* 2018;4:14.
- 8 Hadfield D, Colorado L, Vercueil A, *et al*. The introduction of neurally adjusted ventilatory assist (Nava) into a central London teaching hospital and a comparison with conventional pressure support. *Intensive Care Med* 2010;36:S108.
- 9 Willis GB, Artino AR. What do our Respondents think we're asking? using cognitive interviewing to improve medical education surveys. *J Grad Med Educ* 2013;5:353–6.
- 10 Esteban A, Anzueto A, Frutos F, *et al*. Characteristics and outcomes in adult patients receiving mechanical ventilation: a 28-day International study. *JAMA* 2002;287:345–55.
- 11 GraphPad prism version 7 for windows.
- 12 Elder JP, Artz LM, Beaudin P, *et al*. Multivariate evaluation of health attitudes and behaviors: development and validation of a method for health promotion research. *Prev Med* 1985;14:34–54.
- 13 Chan A-W, Tetzlaff JM, Altman DG, *et al*. Spirit 2013 statement: defining standard protocol items for clinical trials. *Ann Intern Med* 2013;158:200–7.



- 14 Kuo N-Y, Tu M-L, Hung T-Y, *et al.* A randomized clinical trial of neurally adjusted ventilatory assist versus conventional weaning mode in patients with COPD and prolonged mechanical ventilation. *Int J Chron Obstruct Pulmon Dis* 2016;11:945–51.
- 15 Di Mussi R, Spadaro S, Mirabella L, *et al.* Impact of prolonged assisted ventilation on diaphragmatic efficiency: NAVA versus PSV. *Crit Care* 2016;20:1.
- 16 Barwing J, Pedroni C, Quintel M, *et al.* Influence of body position, PEEP and intra-abdominal pressure on the catheter positioning for neurally adjusted ventilatory assist. *Intensive Care Med* 2011;37:2041–5.
- 17 Barwing J, Ambold M, Linden N, *et al.* Evaluation of the catheter positioning for neurally adjusted ventilatory assist. *Intensive Care Med* 2009;35:1809–14.
- 18 Jolley CJ, Luo Y-M, Steier J, *et al.* Neural respiratory drive in healthy subjects and in COPD. *Eur Respir J* 2009;33:289–97.
- 19 Beck J, Sinderby C, Lindström L, *et al.* Effects of lung volume on diaphragm EMG signal strength during voluntary contractions. *J Appl Physiol* 1998;85:1123–34.
- 20 Sinderby C, Beck J, Spahija J, *et al.* Voluntary activation of the human diaphragm in health and disease. *J Appl Physiol* 1998;85:2146–58.
- 21 Beck J, Gottfried SB, Navalesi P, *et al.* Electrical activity of the diaphragm during pressure support ventilation in acute respiratory failure. *Am J Respir Crit Care Med* 2001;164:419–24.
- 22 Sinderby C, Spahija J, Beck J, *et al.* Diaphragm activation during exercise in chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 2001;163:1637–41.
- 23 Helviz Y, Einav S. A systematic review of the high-flow nasal cannula for adult patients. *Crit Care* 2018;22:71.
- 24 Friedman LM, Furberg C, DeMets DL, *et al.* *Fundamentals of clinical trials*. New York: Springer International Publishing, 2015.
- 25 Rose L, Blackwood B, Egerod I, *et al.* Decisional responsibility for mechanical ventilation and weaning: an international survey. *Crit Care* 2011;15:R295.
- 26 Brochard L, Lellouch F. Pressure-support ventilation. In: Tobin MJ, ed. *Principles and Practice of mechanical ventilation*. New York: McGraw-Hill Education, 2013: 199–225.
- 27 Rasmussen J. Skills, rules, and knowledge; signals, signs, and symbols, and other distinctions in human performance models. *IEEE Trans Syst, Man, Cybern IEEE Transactions on Systems, Man, and Cybernetics* 1983;13.

Supplemental file 1: Survey instrument

Manuscript title

Factors affecting the use of Neurally Adjusted Ventilatory Assist in the adult critical care unit: A clinician survey

Corresponding Author

Daniel Hadfield. Email: daniel.hadfield@nhs.net.

BACKGROUND

1. What is your professional group?

Nurse	Doctor	Physiotherapist
<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

2. Current Agenda for Change grade/band (nurses and physiotherapists)

5	6	7	8+
<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

3. Current medical grade / level (if applicable)

FY1/2	Junior resident	Senior Resident (809,927,699)	Consultant
<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

4. Current job title

5. What is your age?

<25	25-34	35-44	45-54	55-64	>64
<input type="radio"/>					

6. For how long have you worked in the ICU at KCH?

< 1 year	1 - 3 years	3-5 years	>5 years
<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

TRAINING AND KNOWLEDGE

7. What form of NAVA training have you received? *Tick ALL that apply*

A	None	<input type="radio"/>
B	Group training from the Maquet representative	<input type="radio"/>
C	Group training from local staff	<input type="radio"/>
D	Bedside training from local staff / peer to peer	<input type="radio"/>
E	Maquet website tutorial	<input type="radio"/>

F	Other (Please state)	<input type="radio"/>
---	----------------------	-----------------------

8. What form of training did you find most useful? **Please enter the relevant letter from the list above**

9. How would you rate the strength of your general understanding of mechanical ventilation in critically ill adults?

Tick ONE option

Much below average	Below average	Average	Above average	Much above average
<input type="radio"/>				

10. Please indicate your level of familiarity with the following:

	Not at all familiar	Slightly familiar	Moderately familiar	Very familiar	Extremely familiar
A: Evidence supporting the use of sedation holds and breathing trials	<input type="radio"/>				
B: Risk factors for prolonged ventilatory weaning	<input type="radio"/>				
C: The current RESTUS / NAVA trial	<input type="radio"/>				
D: Evidence supporting the use of Pressure Support (PS) during weaning	<input type="radio"/>				
E: Evidence supporting NAVA use during weaning	<input type="radio"/>				

11. Have you been involved directly or indirectly in the care of a patient where NAVA was used?

Yes	No
<input type="radio"/>	<input type="radio"/>

12. Of all the patients that you have cared for at KCH, in how many was NAVA used?

≤ 5 patients	5 to 20 patients	> 20 patients	NONE
<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

13. Have you had experience of NAVA in any other hospital?

Yes	No	If yes, approximately how many patients?
<input type="radio"/>	<input type="radio"/>	

CLINICAL EXPERIENCE OF NAVA

14. How recently have you cared for a patient where NAVA was used?

Within the last week	Within the last month	Within the last 6 months	Within the last year	>1 year	Don't know
<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

15. How confident are you in performing the following NAVA related tasks?

	Not at all confident	Slightly confident	Moderately confident	Very confident	Extremely confident	NA
A: Equipment selection, set-up and initiation	<input type="radio"/>					
B: NAVA catheter insertion	<input type="radio"/>					
C: NAVA catheter positioning to obtain diaphragm signal	<input type="radio"/>					
D: Viewing the diaphragm signal (Edi)	<input type="radio"/>					
E: Interpreting the diaphragm signal (Edi)	<input type="radio"/>					
F: Setting the support level	<input type="radio"/>					
G: Weaning	<input type="radio"/>					
H: Trouble shooting	<input type="radio"/>					

16. How have you used the EDI signal? *Tick ALL that apply*

Viewed as a trend across a period of time	<input type="radio"/>
To monitor and/or improve synchrony	<input type="radio"/>
To evaluate readiness for a spontaneous ventilation mode	<input type="radio"/>
To evaluate readiness for extubation	<input type="radio"/>
To measure response to interventions (e.g. spontaneous breathing trials, sedation holds, physiotherapy interventions)	<input type="radio"/>
I have not used the Edi signal	<input type="radio"/>
Other (Please state)	<input type="radio"/>

17. Outside of the current trial, what are the most common reasons for NAVA catheter insertion?

Tick ALL that apply

To accelerate weaning	<input type="radio"/>
-----------------------	-----------------------

To improve synchrony	<input type="radio"/>
To diagnose muscle dysfunction	<input type="radio"/>
To monitor the diaphragm	<input type="radio"/>
Clinician preference	<input type="radio"/>
Don't know	<input type="radio"/>
Other (Please state)	<input type="radio"/>

18. Outside of the current trial, **WHEN** are NAVA catheters most often inserted at KCH? *Tick ONE option*

At or soon after intubation	At the start of weaning	In later weaning	Don't know
<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

19. How **easy** is it to achieve the following when using the **NAVA** mode compared to the **PS** mode?

Please record NA if you have no experience of the specific practice

	Much harder in NAVA	Moderately harder in NAVA	Slightly harder in NAVA	No difference	Slightly easier in NAVA	Moderately easier in NAVA	Much easier in NAVA	NA/ don't know
A: Set-up and start the mode	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
B: Ventilation (adequate MV and CO ₂ clearance)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
C: Lung protection (TV 6-8mls/kg)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
D: Oxygenation	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
E: Synchrony	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
F: Patient comfort	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
G: Weaning	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
H: Maintaining the mode without switching (reliability)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

20. **In your experience**, how did NAVA perform clinically in comparison to Pressure Support?

Significantly worse	Moderately worse	Slightly worse	Equivalent	Slight better	Moderately better	Significantly better	Don't know / NA
<input type="radio"/>							

ADVANTAGES AND DISADVANTAGES

21. What do you consider are the potential clinical **benefits** of using NAVA in comparison to Pressure Support? *Tick ALL that apply*

A	Reduced time on ventilation	<input type="radio"/>
B	Improved patient ventilator interaction (synchrony)	<input type="radio"/>
C	Improved gas exchange	<input type="radio"/>
D	Improved patient sleep	<input type="radio"/>
E	Improved patient comfort	<input type="radio"/>
F	Reduced sedation	<input type="radio"/>
G	Increased variability in pressure and tidal volume	<input type="radio"/>
H	Reduced need for external PEEP to off-set intrinsic PEEP	<input type="radio"/>
I	Monitoring of diaphragm activity and respiratory drive	<input type="radio"/>
J	Reduced risk of diaphragm dysfunction	<input type="radio"/>
K	Improved understanding of respiratory physiology and the patient's condition	<input type="radio"/>
L	Don't know	<input type="radio"/>
M	No clinical benefits	
N	Other (Please state)	<input type="radio"/>

22. What do you consider to be the **ONE** most important potential clinical benefit?

Please enter ONE letter from the list above. Record NA if you answered 'M'

23. What do you consider are the potential clinical **disadvantages** of using NAVA in comparison to Pressure Support? *Tick ALL that apply*

A	Technical issues (signal quality / equipment malfunction / reliability)	<input type="radio"/>
B	Increased variability in pressure and tidal volume	<input type="radio"/>
C	High tidal volumes and/or inspiratory pressures	<input type="radio"/>
D	Inferior gas exchange	<input type="radio"/>
E	Difficult to adjust or regulate	<input type="radio"/>
F	High respiratory rate	<input type="radio"/>

G	Inferior patient ventilator interaction (synchrony)	<input type="radio"/>
H	Don't know	<input type="radio"/>
I	No clinical disadvantages	<input type="radio"/>
J	Other (Please state)	<input type="radio"/>

24. What do you consider is the **SINGLE** most important clinical disadvantage?

Please enter ONE letter from the list above. Record NA if you answered 'I'

25. In your experience, is there an increase in **your personal** workload associated with the use of NAVA?

None	Slight	Moderate	Large	Substantial	Don't know
<input type="radio"/>					

ATTITUDES AND BARRIERS

26. What do you consider are the main barriers to the acceptance and implementation of NAVA?

Tick ALL that apply

A	Poor general performance of NAVA	<input type="radio"/>
B	Lack of experience / insufficient skill set / lack of confidence	<input type="radio"/>
C	Insufficient evidence base	<input type="radio"/>
D	Inadequate clinical guidelines	<input type="radio"/>
E	Increased workload	<input type="radio"/>
F	The need to replace the existing NG catheter	<input type="radio"/>
G	Cost	<input type="radio"/>
H	Resistance from senior clinicians / management	<input type="radio"/>
I	No barriers	<input type="radio"/>
J	Other (Please state)	<input type="radio"/>

27. What do you consider to be the **MAIN** barrier from the list above?

Please enter ONE letter from the list above. Record NA if you answered 'I'

28. What initiative do you think would most help the acceptance and use of NAVA at KCH? *Tick ONE option*

Well conducted research	Improved training	Improved clinical guidelines and protocol	The creation of a 'Super User' group	NAVA catheters in all patients	Other (please write below)
<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
					Other

29. Assuming no contraindications to either mode, which mode would you prefer when initiating ventilator weaning? *Tick one option*

NAVA	Pressure Support	No preference	Other
<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
			Other

30. Please indicate your general feelings towards NAVA on the scale below. *Tick ONE option*

Strongly dislike NAVA	Moderately dislike NAVA	Slightly dislike NAVA	Ambivalent	Slightly like NAVA	Moderately like NAVA	Strongly like NAVA
<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

31. Thinking about your experience, how frequently is the NAVA mode 'switched' to the PS mode?

Never	Rarely	Sometimes	Often	Always	Don't know / NA
<input type="radio"/>					

32. In your experience, what are the main reasons for switching from NAVA to the PS mode? *Tick all that apply*

A	Lack of knowledge / unfamiliarity with the NAVA technology	<input type="radio"/>
B	Personal clinician preference	<input type="radio"/>
C	Problems with the diaphragm signal	<input type="radio"/>
D	Inappropriate pressures or volumes	<input type="radio"/>
E	High respiratory rate	<input type="radio"/>
F	Inadequate gas exchange	<input type="radio"/>
G	Loss of NG catheter	<input type="radio"/>
H	Don't know	<input type="radio"/>

I	Other (Please state)	<input type="radio"/>
---	----------------------	-----------------------

33. What is the main reason that the NAVA mode may be switched to the PS mode?

Please enter ONE letter from the list above. Record NA if you answered 'J'

34. What do you consider to have had the biggest impact on your personal practice and views towards NAVA?

Tick ONE option

Published research	Your own experience	Colleague / peer recommendation	Other (please state below)
<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Other			

35. Please indicate your level of agreement with the following statements:

	Strongly disagree	Disagree	No opinion	Agree	Strongly agree	NA
A: NAVA is safe	<input type="radio"/>					
B: The NAVA mode is clinically effective	<input type="radio"/>					
C: Diaphragm monitoring is clinically effective	<input type="radio"/>					
D: Ventilator dyssynchrony is a clinically significant issue	<input type="radio"/>					
E: The NAVA mode is more difficult to use than PS	<input type="radio"/>					

RESTUS / NAVA RESEARCH

The current RESTUS feasibility study aims to investigate our use of NAVA in prolonged weaning to help us to understand if a larger trial is possible.

36. Are you broadly supportive of the aim of the current research study as stated above?

Yes	No	Don't know
<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

37. Have you cared for a patient who was recruited to the NAVA/ RESTUS trial?

Yes	No	Can't remember / don't know
<input type="radio"/>	<input type="radio"/> (Go to Question 39)	<input type="radio"/>

38. If you have cared for a patient in the RESTUS / NAVA study, how acceptable did you find the research protocol and research process?

Not acceptable	Slightly acceptable	Moderately acceptable	Very acceptable	Completely acceptable	Don't know
<input type="radio"/>					

a. Please comment below.

.....

FINAL COMMENTS

39. Thank you for completing the survey. Please feel free to add any comments or suggestions below.

.....

.....

.....

Supplemental file 2: Response rates and additional data

Manuscript title

Factors affecting the use of Neurally Adjusted Ventilatory Assist in the adult critical care unit: A clinician survey

Corresponding Author

Daniel Hadfield. Email: daniel.hadfield@nhs.net.

Contents

Table 1. Response rates for each question.....	2
Figure S1. Familiarity with the RESTUS trial and aspects of weaning care.....	3
Figure S2. Application of Edi monitoring	4
Figure S3. Staff level of agreement with four statements.....	5
Figure S4. General feelings towards NAVA.....	6
Figure S5. Perceived clinical performance of NAVA versus PSV.....	6
Figure S6. Ease of use, NAVA versus PSV.....	7
Figure S7. Workload associated with NAVA by profession	8
Figure S8. Confidence of NAVA use	9
Figure S9. Perceived advantages of NAVA.....	10
Figure S10. Perceived disadvantages of NAVA	11
Figure S11. Barriers to the acceptance and implementation of NAVA	12
Figure S12. Frequency of mode cross-over	13
Figure S13. Reasons for mode cross-over.....	14
Figure S14. Initiatives to help with clinician acceptance of NAVA	15

Table 1. Response rates for each question

Item number	Number of responders	Item number	Number of responders
1	301	19c	120
2	247	19d	123
3	50	19e	131
4	285	19f	131
5	300	19g	126
6	301	19h	110
7	295	20	190
8	231	21	190
9	294	22	182
10a	293	23	189
10b	294	24	173
10c	293	25	188
10d	292	26	183
10e	291	27	177
11	294	28	167
12	287	29	181
13	29	30	179
14	193	31	182
15a	191	32	180
15b	193	33	168
15c	193	34	174
15d	193	35a	181
15e	190	35b	181
15f	192	35c	181
15g	192	35d	179
15h	192	35e	181
16	191	36	180
17	192	37	121
18	193	38	125
19a	116	39	42
19b	121		

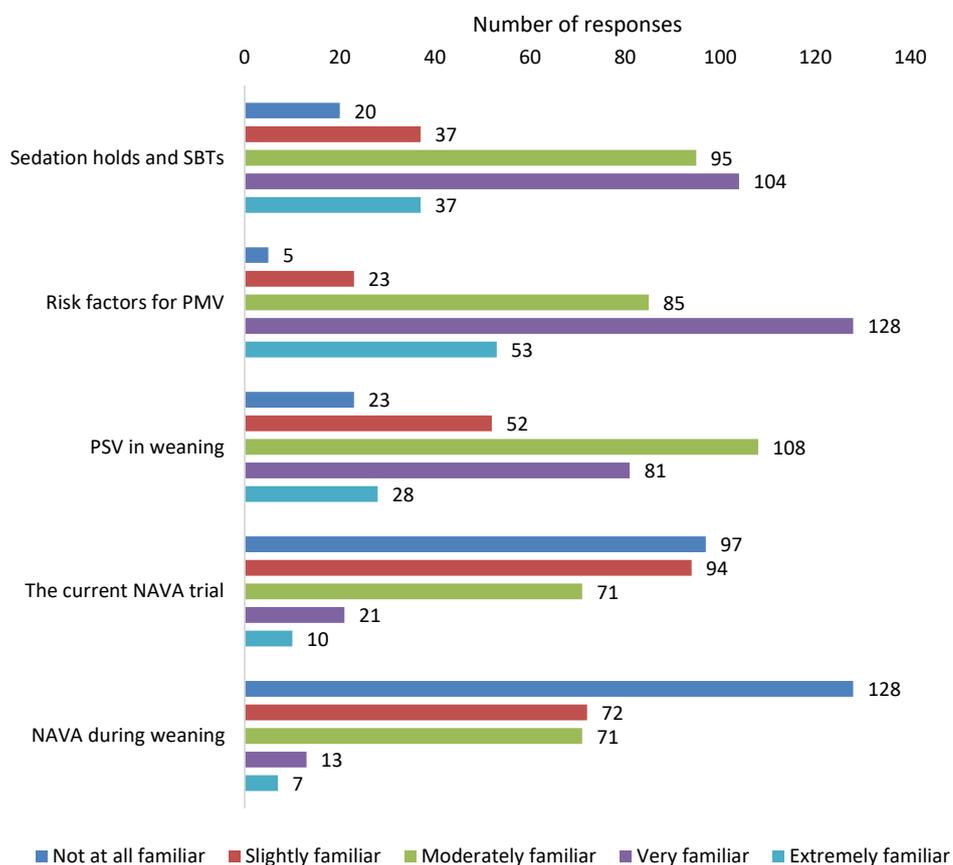


Figure S1. Familiarity with the RESTUS trial and aspects of weaning care.

Please indicate your familiarity with the following: A. Evidence supporting the use of sedation holds and spontaneous breathing trials (SBTs); response rate = 293

B. Risk factors for prolonged MV (PMV); response rate = 294

C. Evidence supporting the use of PSV during weaning; response rate = 292

D. The current NAVA trial; response rate = 292

E. Evidence supporting the use of NAVA during weaning; response rate = 291

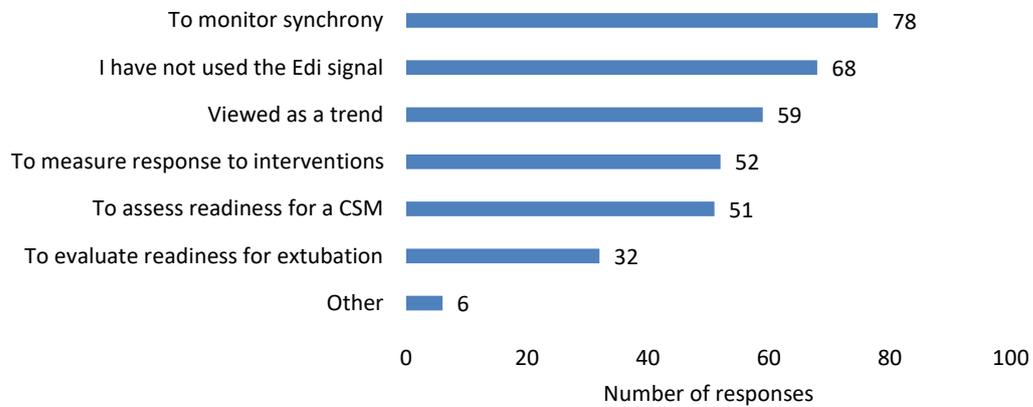


Figure S2. Application of Edi monitoring

How have you used the Edi (electrical activity of the diaphragm) signal? Participants could select multiple items. Response rate: 191. 'Other' responses (n=6), were categorised as 'Invalid' (n=5) and one referred to the use of Edi monitoring in post-extubation support. CSM = Continuous Spontaneous Mode (NAVA or PSV)

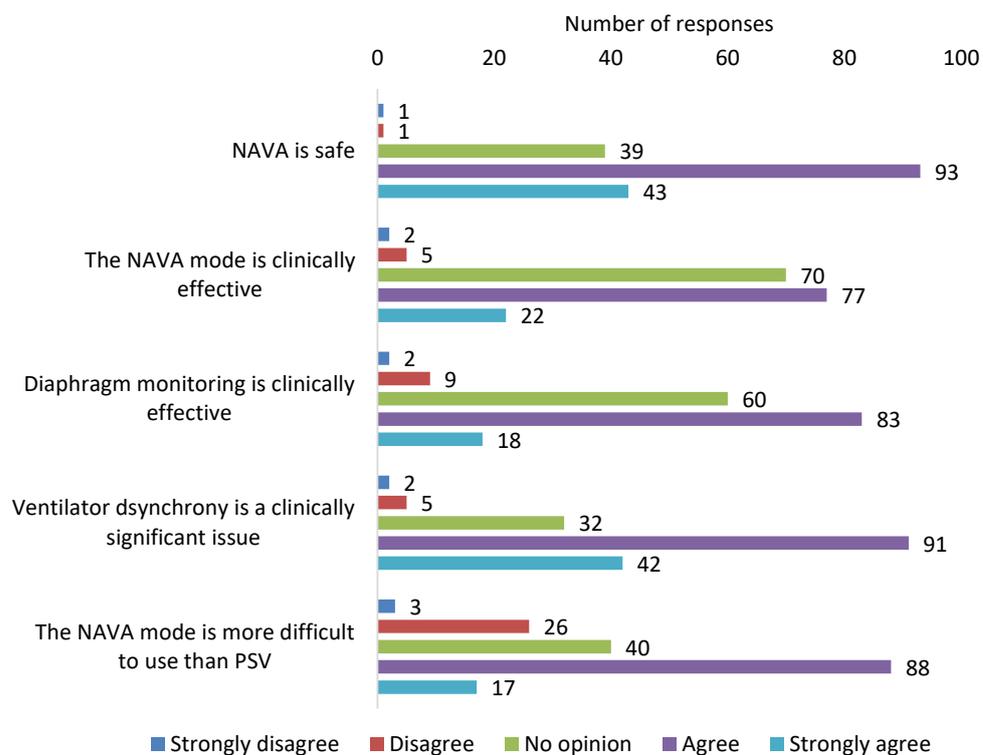


Figure S3. Staff level of agreement with four statements

Please indicate your level of agreement with the following statements: A. NAVA is safe; total responses 181; NA = 4

B. The NAVA mode is clinically effective; total responses 181; NA = 5

C. Diaphragm monitoring is clinically effective; total responses 181; NA = 9

D. Ventilator dysynchrony is a clinically significant issue; total responses 179; NA = 7

E. The NAVA mode is more difficult to use than PSV; total responses 181; NA = 7

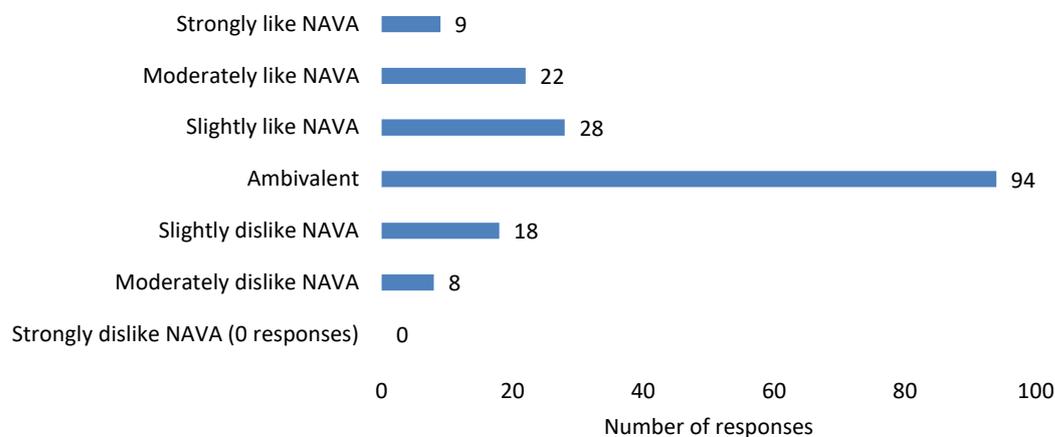


Figure S1. General feelings towards NAVA

Please indicate your general feelings towards NAVA. Total respondents 179. Participants could select one option only. Total respondents = 179

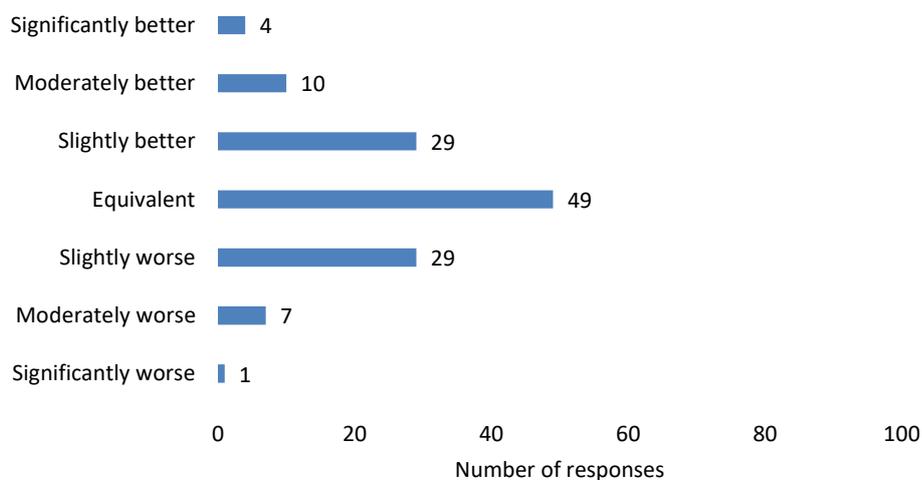


Figure S5. Perceived clinical performance of NAVA versus PSV

In your experience, how did NAVA perform clinically in comparison to Pressure Support? Total respondents = 190. Participants could answer one option only. 'Don't know' (n=61)

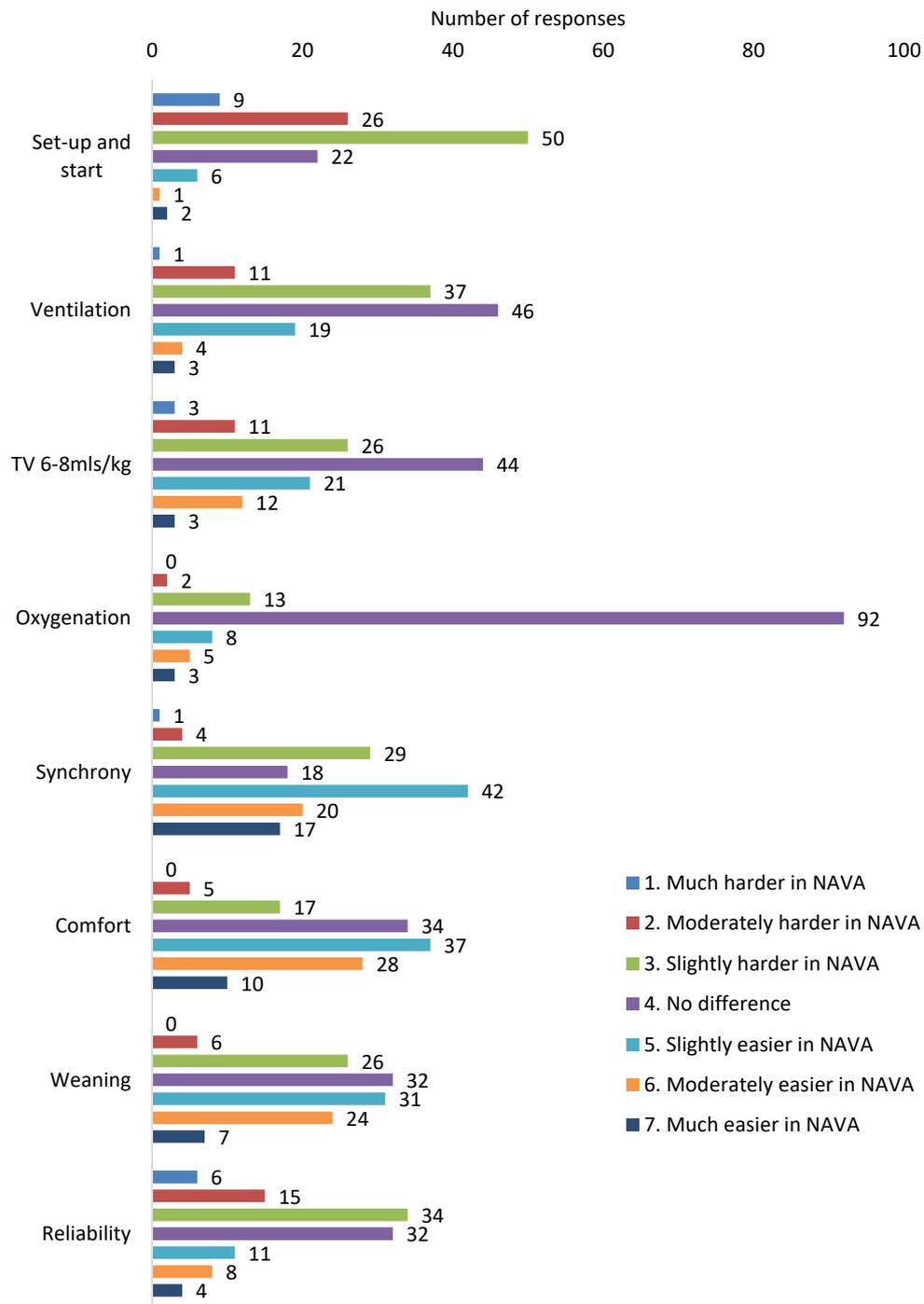


Figure S6. Ease of use, NAVA versus PSV

How easy is it to achieve the following aspects of ventilation practice when using the NAVA mode compared to the PSV mode? Response rates and 'Not Applicable' (NA) responses:

A. Set-up and start; response rate 116, NA 75

B. Ventilation (adequate minute volume and CO₂ clearance); response rate 121, NA 70

C. Lung protection (achieving tidal volumes of 6-8mls/kg); response rate 120, NA 71

D. Oxygenation; response rate 123, NA 68

E. Synchrony; response rate 131, NA 60

F. Patient comfort; response rate 131, NA 61

G. Weaning; response rate 126, NA 66

H. Maintaining the mode without 'switching' (reliability); response rate 110, NA 82

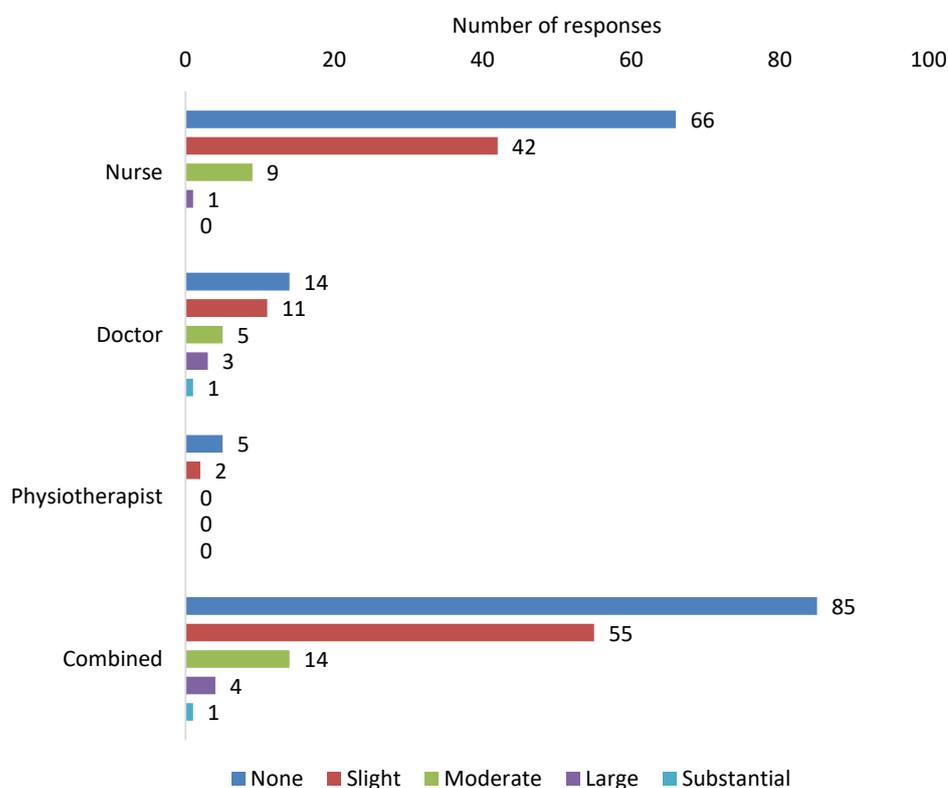


Figure S7. Workload associated with NAVA by profession

In your experience, is there an increase in your personal workload associated with the use of NAVA?

Total respondents 188, comprised of 118 nurses, 34 doctors and 7 physiotherapists. Don't know/NA responses (n=29)

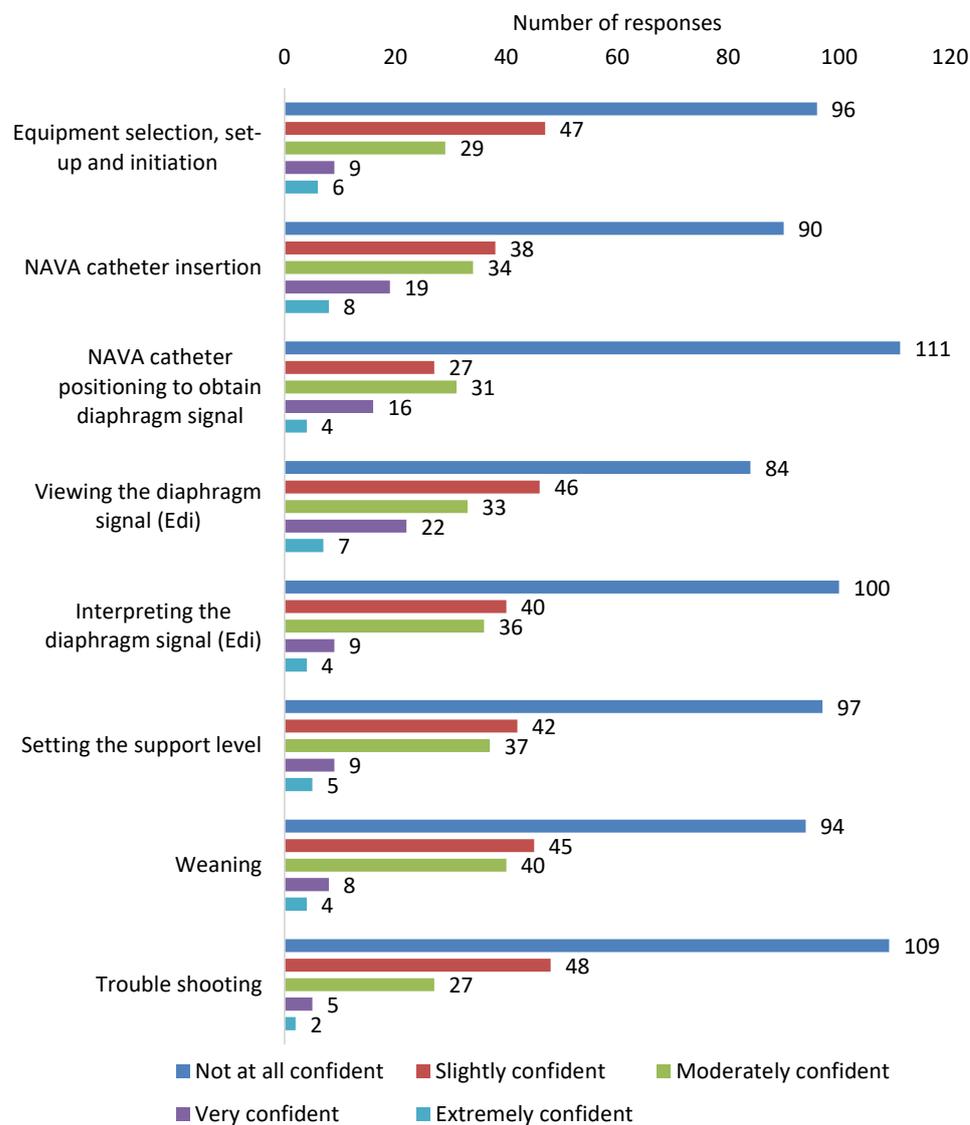


Figure S2. Confidence of NAVA use

How confident are you in performing the following NAVA related tasks? Response rates and 'Not Applicable' (NA) responses:

A: Equipment selection, set-up and initiation; response rate 191, NA 4

B: NAVA catheter insertion; response rate 193, NA 4

C: NAVA catheter positioning to obtain diaphragm signal; response rate 193, NA 4

D: Viewing the diaphragm signal (Edi); response rate 193, NA 1

E: Interpreting the diaphragm signal (Edi); response rate 190, NA 1

F: Setting the support level; response rate 192, NA 2

G: Weaning; response rate 192, NA 1

H: Trouble shooting; response rate 192, NA 1

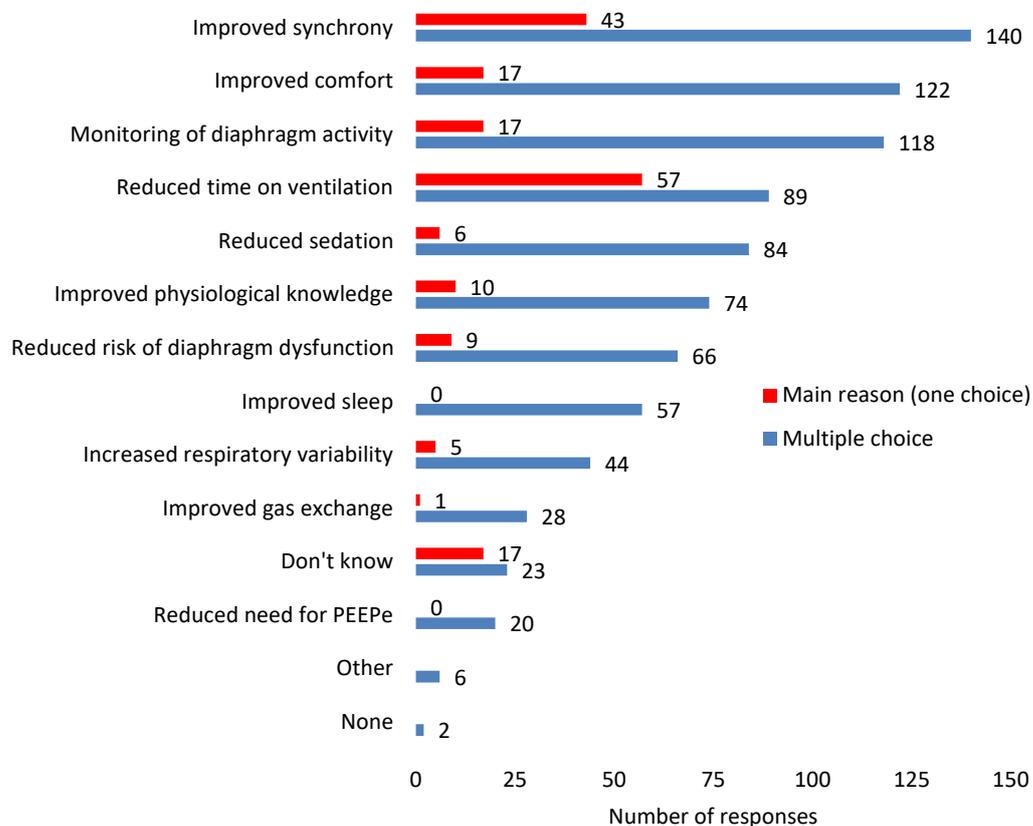


Figure S9. Perceived advantages of NAVA

What do you consider are the potential clinical benefits of using NAVA technology in comparison to PSV? Participants could select multiple options (blue line). Other responses were categorised as 'invalid' (n=5) and 'Monitoring of diaphragm activity' (n=1). Response rate: 190. Participants were also asked 'What do you consider to be the one most important potential clinical benefit from the list above?' Participants were asked to tick one option only. Response rate = 182. Two 'Other' responses were categorised as 'invalid'.

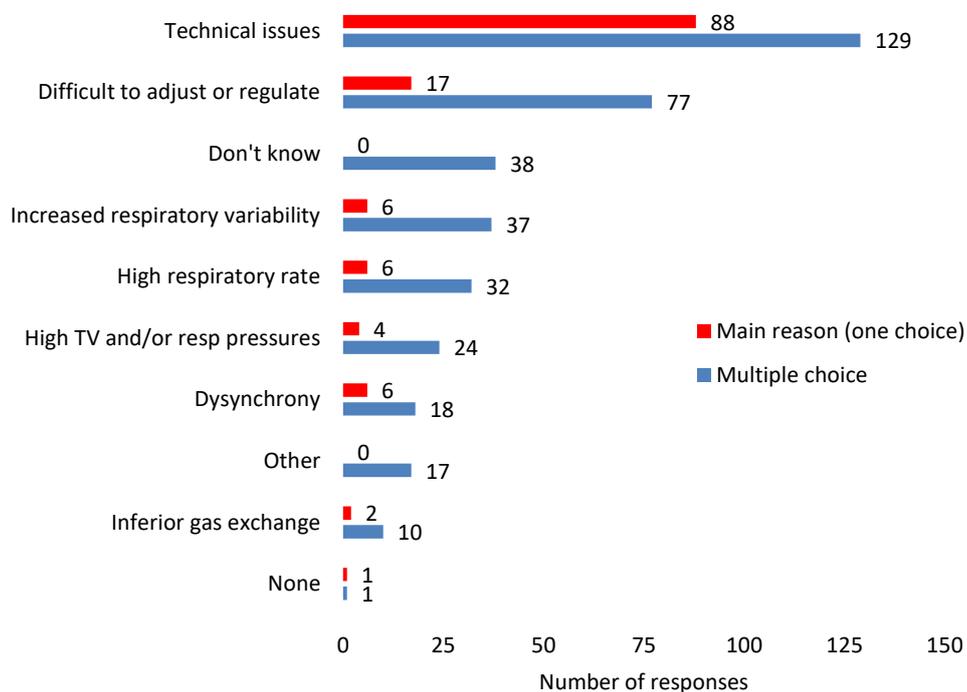


Figure S10. Perceived disadvantages of NAVA

What do you consider are the potential clinical disadvantages of using NAVA in comparison to Pressure Support? Participants could select multiple items. Participants were informed that the response 'Technical issues' included signal quality, equipment malfunction, reliability. Response rate: 189. Free text other responses referred to difficulty with NAVA catheter connections (n=1), increased complexity in trouble shooting (n=1), and lack of staff training/familiarity (n=2). Participants were also asked 'What do you consider is the one most important clinical disadvantage?'. Response rate: 173. Participants were asked to tick one option only (red line). Other responses included 'NA' (n=8), lack of knowledge/familiarity (n=6) and the possibility of accidental NAVA catheter removal (n=1)

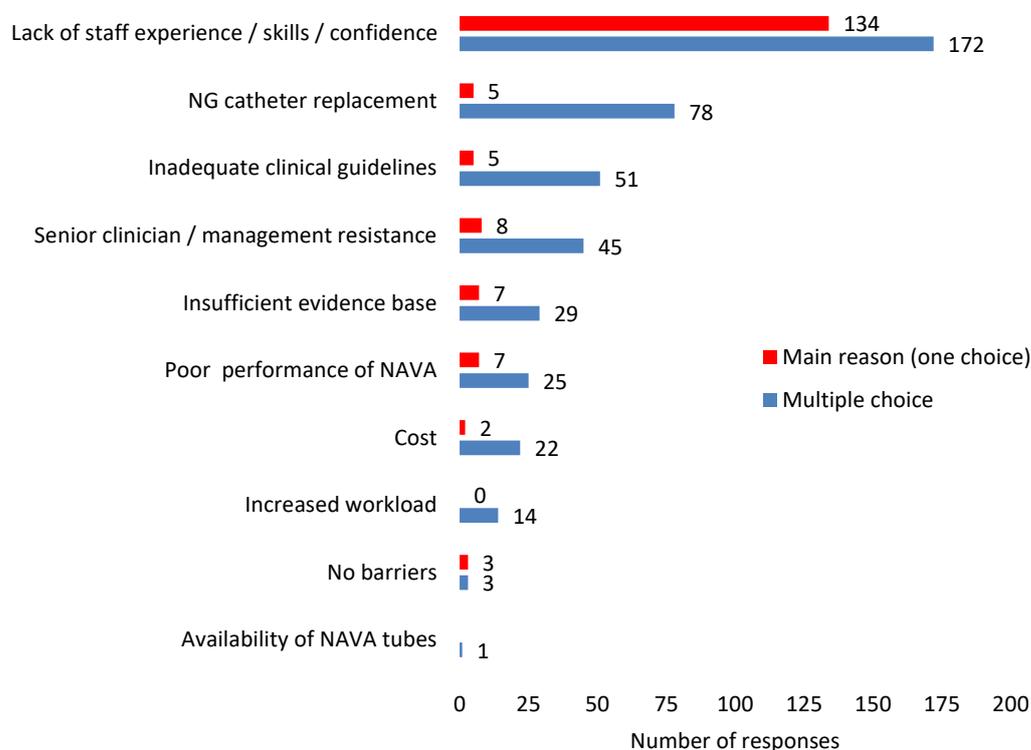


Figure S11. Barriers to the acceptance and implementation of NAVA

What do you consider are the main barriers to the acceptance and implementation of NAVA? Total respondents = 183. Participants could select multiple options (blue line). Other responses (n=12) were categorised post hoc as 'Lack of staff experience / skills confidence' (n=4), availability of NAVA tubes (n=1), 'Poor performance of NAVA' (n=1), and 'invalid' (n=6). Participants were also asked, 'What do you consider to be the main barrier from the list above?'. Participants were asked to select one option only (red line). Total respondents = 177. 'Other' responses (n=8) were categorised post hoc to 'Lack of staff experience / skills confidence' (n=2) and 'invalid' (n=6).

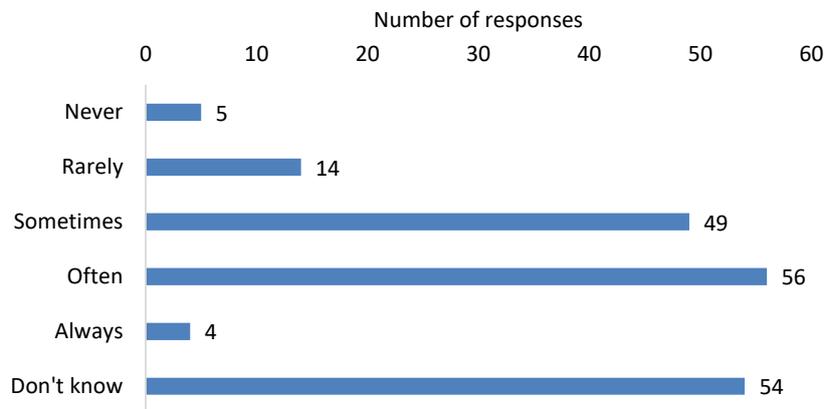


Figure S12. Frequency of mode cross-over

Thinking about your experience, how often is the NAVA mode 'switched' to the PSV mode? Total respondents = 182. Participants could select one option only

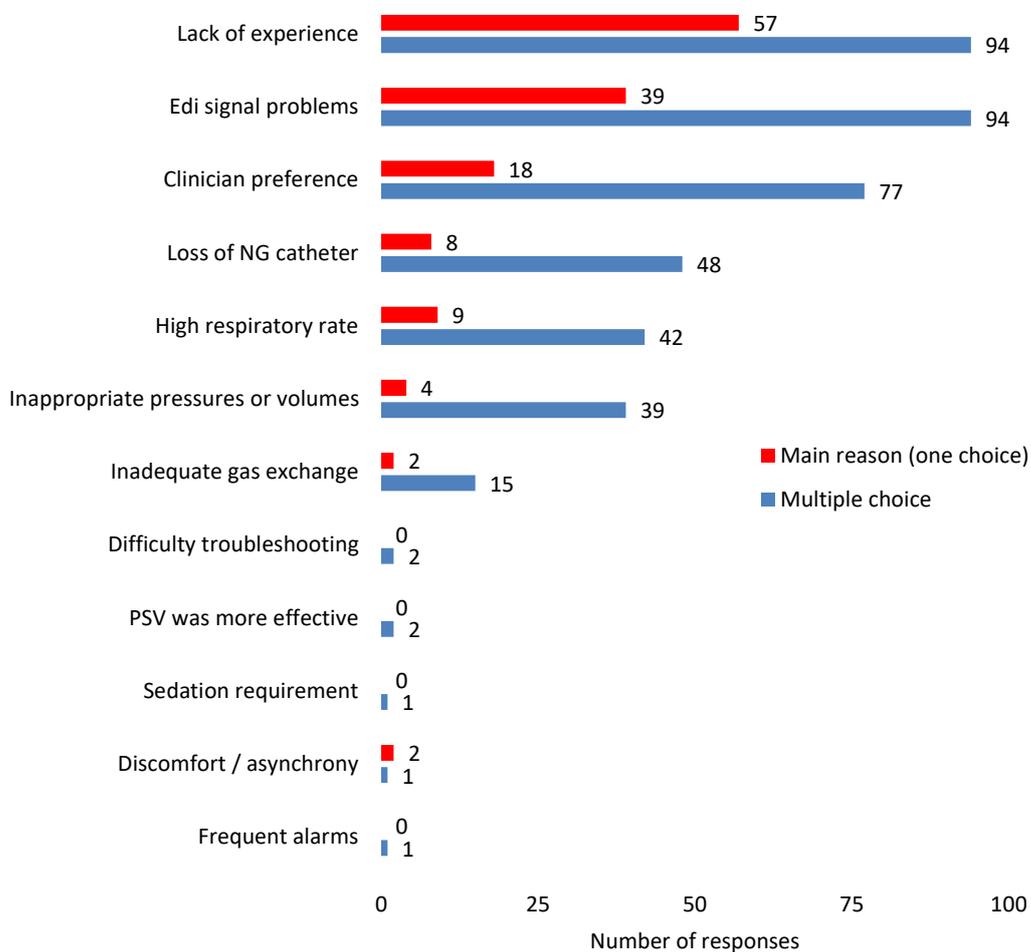


Figure S3. Reasons for mode cross-over

In your experience, what are the main reasons for switching from NAVA to the PSV mode? Total respondents = 180. Participants could select multiple options (blue line). Participants were also asked to select what they considered to be the main reason (one option only) for switching from NAVA to the PSV mode (red line); total respondents = 168. 'Other' responses (n=6) were categorised as 'Sedation requirement' (n=1), 'Lack of experience' (n=3), 'Patient discomfort / dysynchrony' (n=2), PSV more effective (n=3), difficulty troubleshooting (n=2), 'invalid' (n=4).

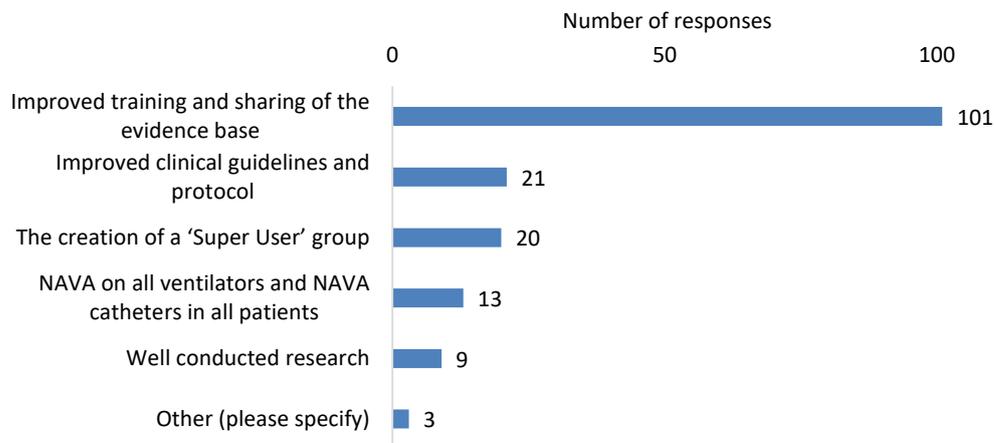


Figure S14. Initiatives to help with clinician acceptance of NAVA

What initiative do you think would most help the acceptance and use of NAVA at KCH? Participants could answer one option only. 'Other' responses were classified as invalid