XXX International Symposium of Mechanical Ventilation of Albert Einstein Hospital Albert Einstein Hospital, São Paolo, Brazil August 17, 2023; 14:50–15:10 am BRT

Noninvasive Ventilation and (/versus) High–flow Nasal Oxygen

when (not) to start, and when to stop?



University of Amsterdam, The Netherlands

Oxford University, UK



Disclosures

- until January 2021 Xenios/Fresenius, Germany
- until January 2023 Hamilton Medical AG, Switzerland

less experience with NIV than with HFNO



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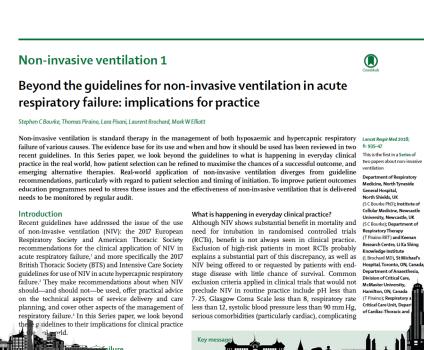
Limitations

- NIV and HFNO in patients in non-hypercaphic AHRF
- 'beyond the guidelines'

Bourke Lancet Resp Med 2018; 6:935

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two papers about non-invasiv (S C Bourke PhD); Institute o Cellular Medicine, Newcastl

Series



- changing landscapes?
- noninvasive ventilation
- high—flow oxygen
- pandemics
- conclusions



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Increase in Use of Noninvasive Ventilation in Hypercapnic Patients

	Number of hospitals	Number of patients	Mean age	Consolidation on radiograph	Median initial PaCO ₂	Median Initial pH	NIV unsuccessful	IMV	Died		Proportion discharged from hospital
									All causes	Respiratory	
2010	61	925	71	30%	10.2	7.30	27%	2.3%	29%	22%	67%
2011	122	2187	71	38%	10.1	7.26	33%	3.8%	30%	25%	66%
2012	130	2490	72	40%	10.2	7.25	31%	2.7%	31%	26%	65%
2013	148	2693	72	40%	10.2	7.24	33%	3.0%	34%	27%	66%

Table shows data for adult patients admitted to hospital with COPD exacerbation receiving NIV. NIV=non-invasive ventilation. PaCO₂=arterial partial pressure of carbon dioxide. IMV=invasive mechanical ventilation. COPD=chronic obstructive pulmonary disease.

Table: British Thoracic Society national audits of NIV, 2010–13

Bourke Lancet Resp Med 2018; 6:935



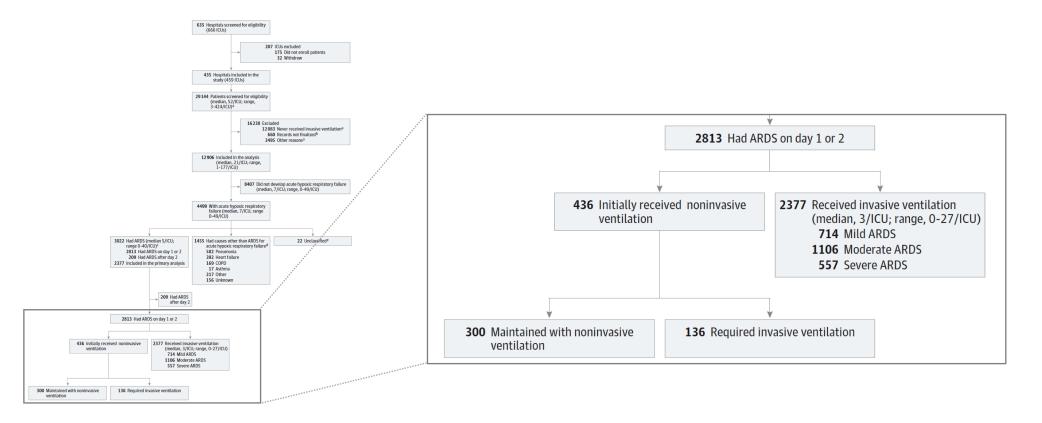
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Use of Noninvasive Ventilation in ARDS



Bellani JAMA 2016; 315:788



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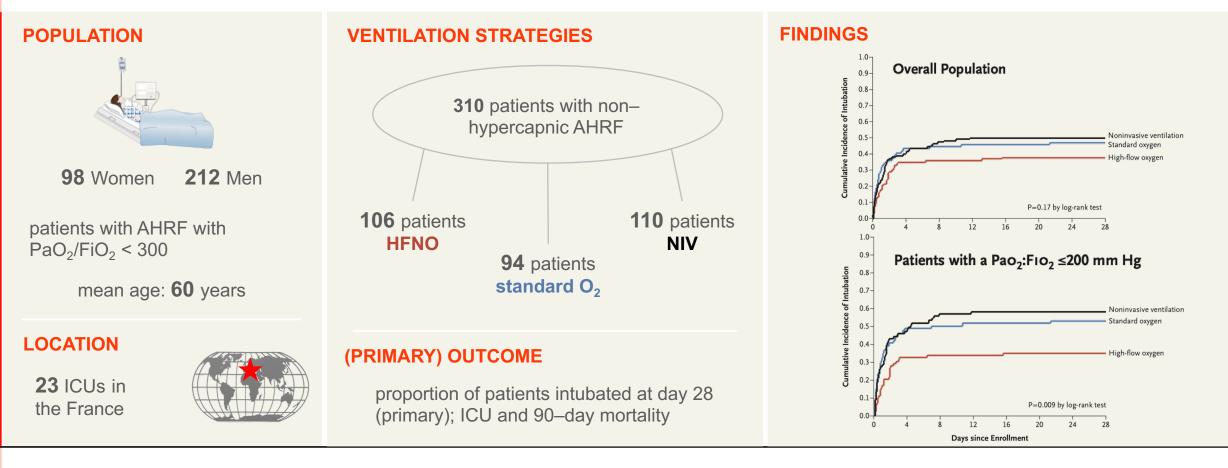
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QUESTION In patients with non–hypercapnic acute hypoxemic respiratory failure (AHRF), what are the effects of treatment with high–flow nasal oxygen (HFNO), standard oxygen, or noninvasive ventilation on need for intubation??

CONCLUSION In patients with non–hypercapnic AHRF, treatment with HFNO, standard oxygen, or noninvasive ventilation did not result in different intubation rates; there was a significant difference in favor of HFNO in 90–day mortality.

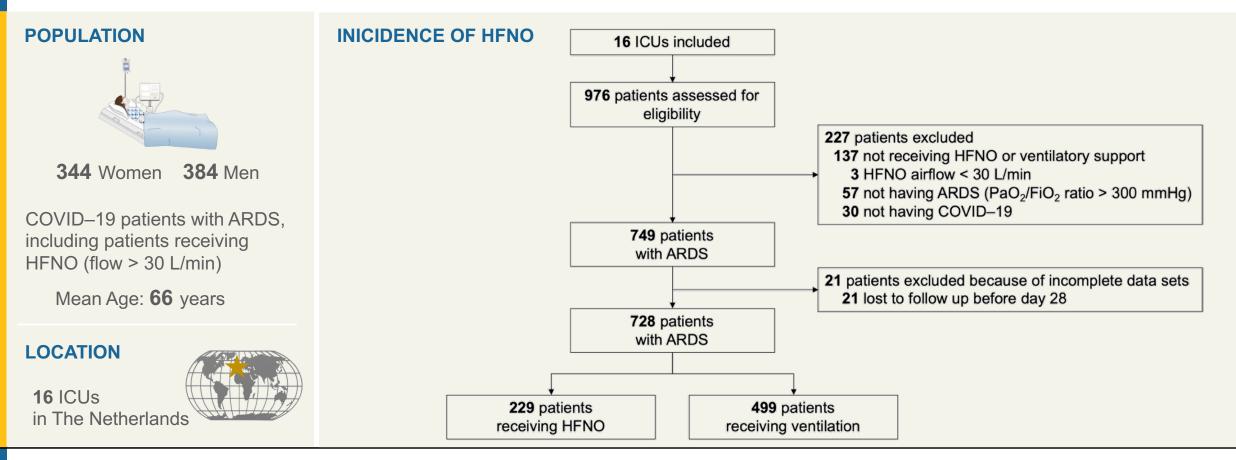


FLORALI–investigators. High–flow Oxygen through Nasal Cannula in Acute Hypoxemic Respiratory Failure. [*New Eng J Med* 2015; **372**:2185 doi: 10.1056/NEJMoa1503326]



QUESTION Does a broadened Berlin definition of ARDS, in which ARDS can be diagnosed in patients who are not receiving ventilation, results in similar groups of patients receiving HFNO as in patients receiving ventilation?

CONCLUSION Using a broadened definition of ARDS may facilitate an earlier diagnosis of ARDS in patients receiving HFNO; however, ARDS patients receiving HFNO and ARDS patients receiving ventilation have distinct baseline characteristics and mortality rates.

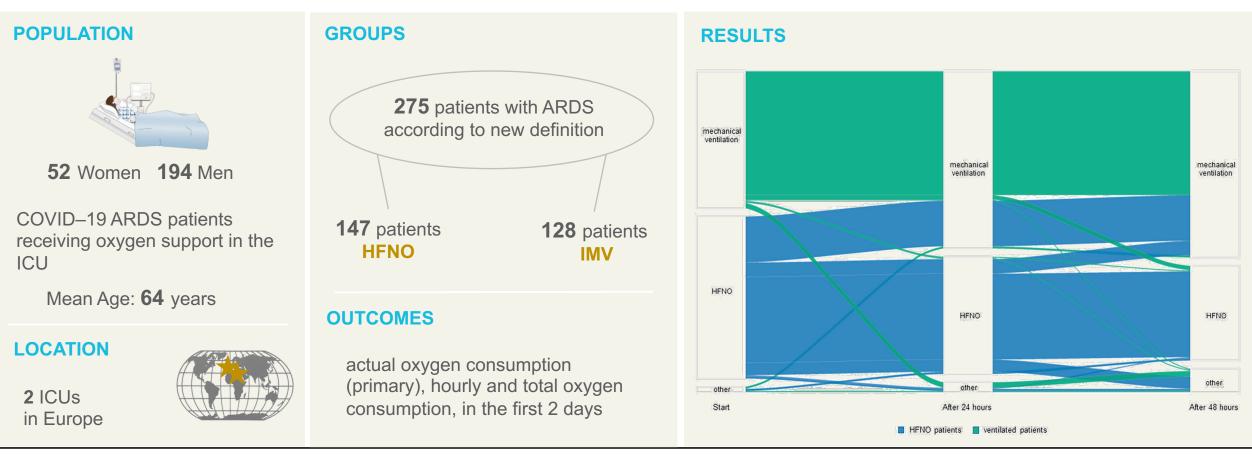


PRoAcT–COVID study investigators. Broadening the Berlin definition of ARDS to patients receiving high-flow nasal oxygen: an observational study in patients with acute hypoxemic respiratory failure due to COVID–19. [*AoIC* 2023; **13**:64; doi:10.1186/s13613-023-01161-6]



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CONCLUSION Actual oxygen consumption, hourly oxygen consumption, and total oxygen consumption are substantially higher in patients that start with HFNO.



PROXY–COVID study investigators. Oxygen Consumption with High-Flow Nasal Oxygen versus Mechanical Ventilation—An International Multicenter Observational Study in COVID–19 Patients (PROXY–COVID). [Am J Trop Med Hyg 2023; **108**:1035; doi:10.4269/ajtmh.22-0793]

Noninvasive Ventilation vs Standard or High–flow Oxygen

- equipment
- interfaces
- PEEP
- cooperation
- failures



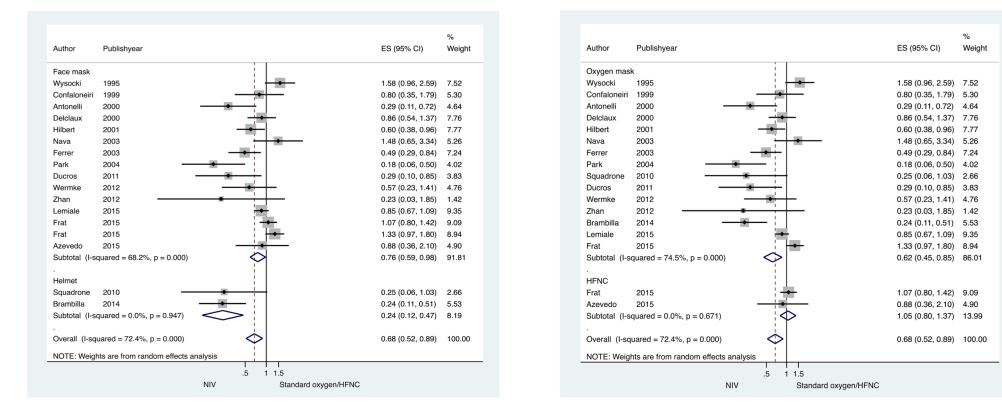
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Intubation Rate with Noninvasive Ventilation vs Standard or High–flow Oxygen



Aswanetmanee Scientific Reports 2023; 13:8283



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JAMA

QUESTION What is the effect of noninvasive ventilation delivered by helmet vs usual respiratory support on the risk of mortality among adults with acute hypoxemic respiratory failure due to COVID-19?

CONCLUSION Helmet noninvasive ventilation did not significantly reduce 28-day mortality compared with usual respiratory support in patients with acute hypoxemic respiratory failure due to COVID-19; however, study interpretation is limited by imprecise effect estimate.

POPULATION INTERVENTION FINDINGS Mortality rate at 28 days **187** Men Helmet noninvasive **Usual respiratory** 133 Women 322 Patients randomized ventilation support **320** Patients analyzed 43 of 159 patients 42 of 161 patients Adults with acute hypoxemic respiratory failure due to 159 161 suspected or confirmed COVID-19 Helmet noninvasive **Usual respiratory** 27.0% 26.1% ventilation support Median age: **58** years Oxygen delivered noninvasively Mask noninvasive ventilation. via a helmet device high-flow nasal oxygen, and standard oxygen LOCATIONS The between-group difference was not significant: 8 ICUs **PRIMARY OUTCOMES Risk difference**, **1.0%** (95% CI, -8.7% to 10.6%) in Saudi Arabia and Kuwait 28-day all-cause mortality **Relative risk**, **1.04** (95% CI, 0.72 to 1.49); *P* = .85

Arabi YM, Aldekhyl S, Al Qahtani S, et al; Saudi Critical Care Trials Group. Effect of helmet noninvasive ventilation vs usual respiratory support on mortality among patients with acute hypoxemic respiratory failure due to COVID-19. JAMA. Published September 20, 2022. doi:10.1001/jama.2022.15599

Agenda

- changing landscapes?
- noninvasive ventilation
- high–flow oxygen
- pandemics
- conclusions

Series Non-invasive ventilation 1 Beyond the guidelines for non-invasive ventilation in acute respiratory failure: implications for practice Stephen C Bourke, Thomas Piraino, Lara Pisani, Laurent Brochard, Mark W Elliott Non-invasive ventilation is standard therapy in the management of both hypoxaemic and hypercapnic respiratory Lancet Respir Med 2018; failure of various causes. The evidence base for its use and when and how it should be used has been reviewed in two 6:935-47 recent guidelines. In this Series paper, we look beyond the guidelines to what is happening in everyday clinical This is the first in a Series of two papers about non-invasiv practice in the real world, how patient selection can be refined to maximise the chances of a successful outcome, and ventilation emerging alternative therapies. Real-world application of non-invasive ventilation diverges from guideline recommendations, particularly with regard to patient selection and timing of initiation. To improve patient outcomes Department of Respirator education programmes need to stress these issues and the effectiveness of non-invasive ventilation that is delivered General Hospital, needs to be monitored by regular audit. North Shields, UK (S C Bourke PhD): Institute of Cellular Medicine, Newcastl Introduction What is happening in everyday clinical practice? University, Newcastle, UK Recent guidelines have addressed the issue of the use Although NIV shows substantial benefit in mortality and (SC Bourke) Repartment of of non-invasive ventilation (NIV): the 2017 European need for intubation in randomised controlled trials Respiratory Therapy (T Piraino RRT) and Keena Respiratory Society and American Thoracic Society (RCIs), benefit is not always seen in clinical practice. recommendations for the clinical application of NIV in Exclusion of high-risk patients in most RCTs probably Knowledge Institute Research Centre, Li Ka Shino acute respiratory failure,¹ and more specifically the 2017 explains a substantial part of this discrepancy, as well as (Brochard MD) St Michael's British Thoracic Society (BTS) and Intensive Care Society NIV being offered to or requested by patients with end-Hospital, Toronto, ON, Canada guidelines for use of NIV in acute hypercapnic respiratory stage disease with little chance of survival. Common Department of Anaesthesia Division of Critical Care, exclusion criteria applied in clinical trials that would not failure.2 They make recommendations about when NIV should-and should not-be used, offer practical advice preclude NIV in routine practice include pH less than Hamilton. (A anda on the technical aspects of service delivery and care 7.25, Glasgow Coma Scale less than 8, respiratory rate (TPiraino); Respiratory planning, and cover other aspects of the management of less than 12, systolic blood pressure less than 90 mm Hg, of Cardiac Thoracic an piratory failure.² In this Series paper, we look beyond serious comorbidities (particularly cardiac), complicating

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idelines to their implications for clinical practice

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Panel 2: Prognostic indices to be considered prior to initiation of acute non-invasive ventilation in acute hypercapnic respiratory failure

Cause of acute hypercapnic respiratory failure

- Favourable: chronic obstructive pulmonary disease, extra-pulmonary restriction, and cardiogenic pulmonary oedema
- Adverse: pulmonary fibrosis and isolated pneumonia

Stable state

- Poor performance status
 - Unable to leave home unassisted
 - Requires help washing and dressing
- High comorbidity burden
- Low body-mass index

Severity of acute illness

- Blood gas abnormalities
 - Late development of acute hypercapnic respiratory failure after admission
 - Coexistent metabolic acidaemia or low base excess
 - Severe acidaemia (pH <7.25)
- Other organ failure or impairment
- Consolidation
- Observations including: respiratory rate >30, hypotension (particularly if unresponsive to fluid resuscitation), and low Glasgow Coma Scale (<11)
- Blood results including: eosinopenia (<50 cells per μL), raised urea, and hypoalbuminaemia
- Inability to clear secretions

Indices listed are associated with worse outcome unless otherwise stated. No single index in isolation should preclude a trial of non-invasive ventilation.

Panel 4: Prognostic factors for successful non-invasive ventilation in acute hypoxaemic respiratory failure

Cause of acute hypoxaemic respiratory failure

- Favourable: cardiogenic pulmonary oedema, post-operative, and PaO₂/FiO₂ > 200 mm Hg
- Adverse: $PaO_2/FiO_2 < 200 \text{ or } 150 \text{ mm Hg}$

Predictors of failure

- $PaO_2/FiO_2 < 150 \text{ mmHg}$
- Tidal volume (exhaled) under non-invasive ventilation (NIV) ≥9.0 or 9.5 mL/kg
- High severity score (eg, Acute Physiology And Chronic Health Evaluation II or Sequential Organ Failure Assessment)
- Heart rate, acidosis, consciousness, oxygenation, respiratory rate score >5 after 1 h of NIV

Considerations

- Trial of high-flow nasal cannula for $PaO_2/FiO_2 < 200 \text{ mm Hg}$
- Avoid delaying intubation

Text: Use of NIV in ARDS

Limited information in the literature

Uncertain whether NIV should be used

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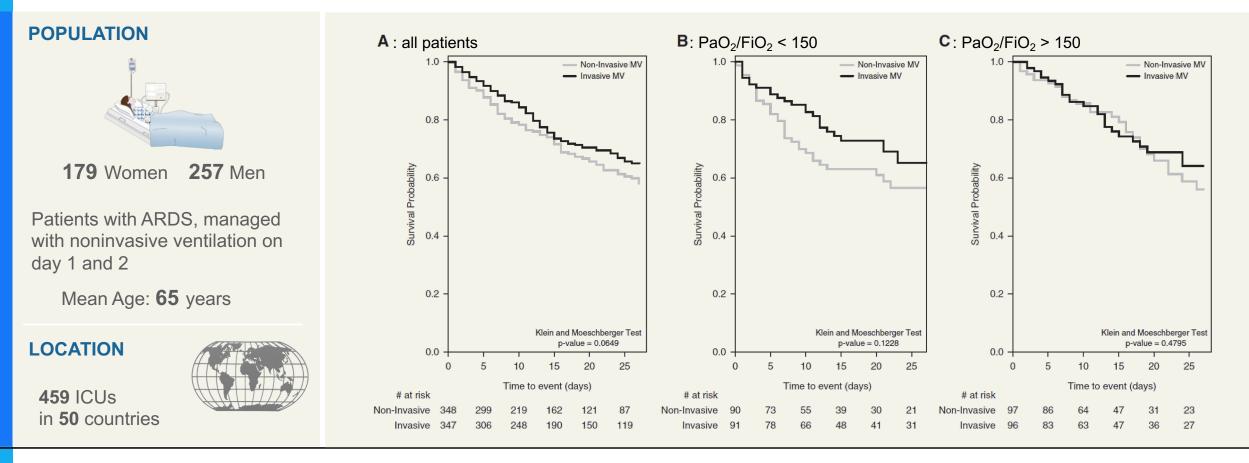
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QUESTION Is during NIV the categorization of ARDS severity based on the PaO₂/FiO₂ Berlin criteria is useful?

CONCLUSION NIV was used in 15% of patients with ARDS, irrespective of severity category; NIV seems to be associated with higher ICU mortality in patients with a PaO_2/FiO_2 lower than 150 mm Hg.



LUNG SAFE investigators. Noninvasive Ventilation of Patients with Acute Respiratory Distress Syndrome - Insights from the LUNG SAFE Study. [*Am J Resp Crit Care Med* 2016; **175**:67; doi:10.1164/rccm.201606-1306OC]

Key messages

- Guidelines for the use of non-invasive ventilation (NIV) in acute or chronic hypercaphic respiratory failure and acute hypoxaemic respiratory failure are evidence based and should be followed
- The right patient
 - The cause of respiratory failure is important in determining the likelihood of a successful outcome with NIV
 - NIV should not be used when it is very unlikely to succeed or when a purely palliative approach would be more appropriate—prediction tools should inform decision making
 - The right time
 - Physiological criteria should be used to determine the timing of NIV
 - NIV should be discontinued in a timely manner if the patient is deteriorating on the basis of worsening pH and respiratory rate (for acute hypercapnic respiratory failure) or exhaled tidal volume >9.5 mL/kg and heart rate, acidosis, consciousness, oxygenation, respiratory rate score >5 after 1 h (for hypoxaemic respiratory failure)
- The right equipment
 - The correct interface should be used and should fit well
 - Condition-specific settings should be used, and adjusted according to response
- The right environment
 - The unit or ward should be properly staffed and resourced
 - Staff should be NIV trained and competency assessed
 - Training should be updated regularly
- Ongoing audits and quality assurance should be done



- changing landscapes?
- noninvasive ventilation
- high—flow oxygen
- pandemics
- conclusions



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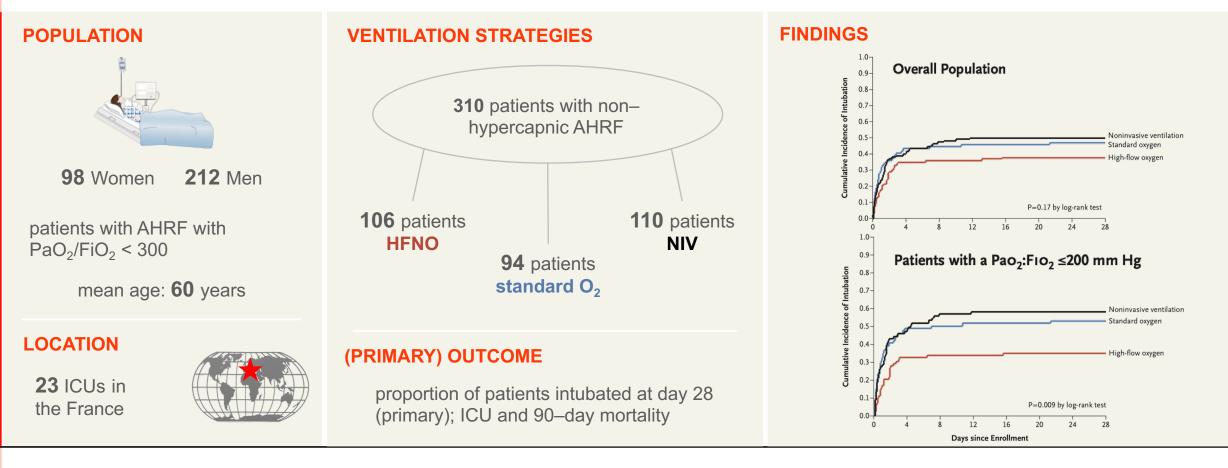


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Benefits and Risks of HFNO

- comfortable
- failures less well recognized
- oxygenation
- in case of hypoxic failure little reserves
- lower risk of P–SILI?



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ROX index = SpO_2/FiO_2 to RR

- predictors of HFNC failure
 - ROX < 2.85 at 2 hours
 - ROX < 3.47 at 6 hours
 - ROX < 3.85 at 12 hours



Roca AJRCCM 2019; 119:1368



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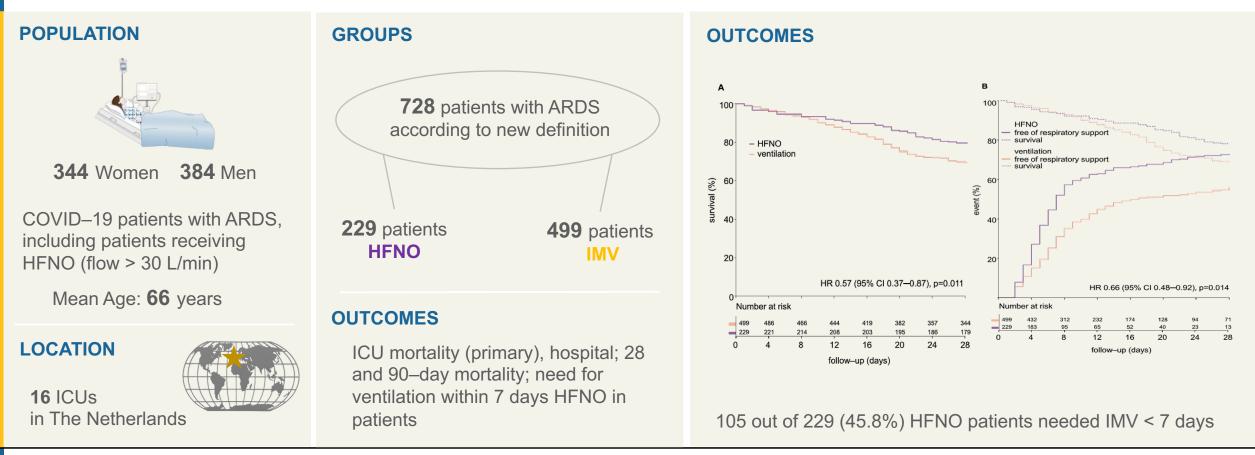


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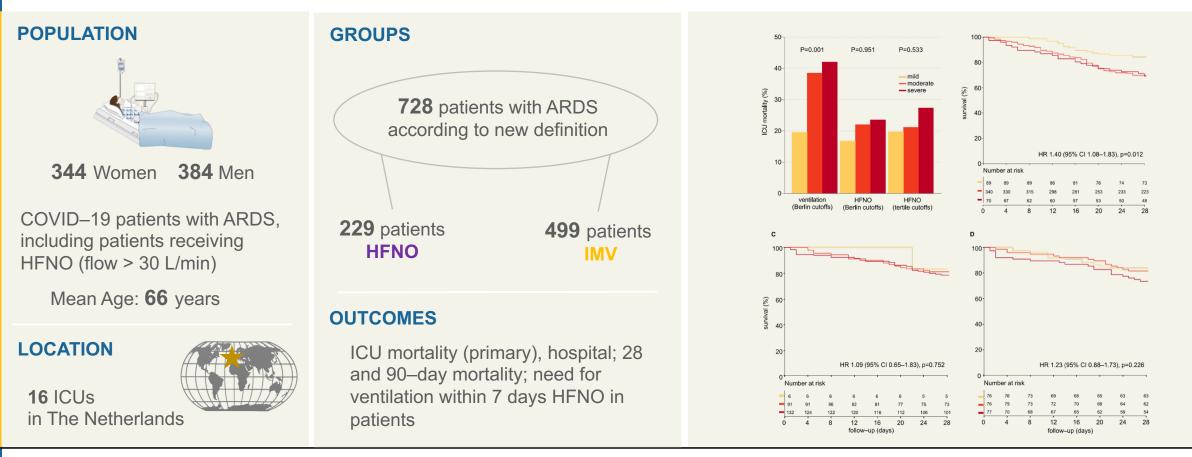


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Oxygen Scarcity

• running out of oxygen = realistic scenario



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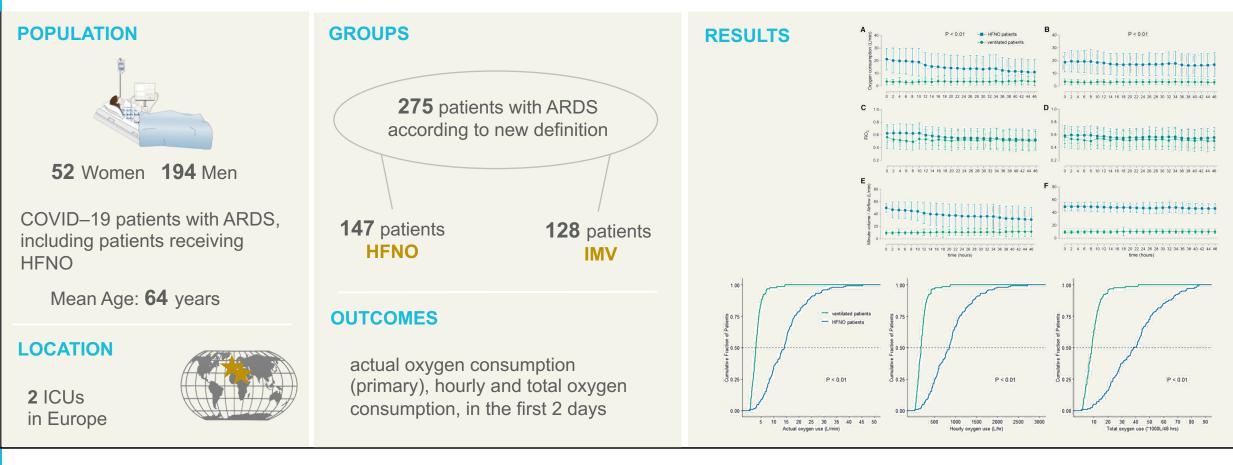


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Oxygen Use with High–flow Oxygen

- 850 vs 176 L/hour, or 20.000 vs 4.200 L/day
 - 10.000 liquid oxygen tank ~ 400 HFNO days
 - pressure swing adsorption plant 1.000 l/min ~ 70 HFNO patients
 - 50 L steel cylinder ~ 9 HFNO days
 - oxygen generator ~ 1 HFNO patient

Botta A J Trop Med Hyg 2023; 108:1035



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Oxygen Use with High–flow Oxygen

- oxygen-sparing strategies
 - automated oxygen titration
 - better bedside use of guidelines with strict cutoffs for SpO₂
 - prone positioning
 - noninvasively or invasive ventilation

Botta A J Trop Med Hyg 2023; 108:1035



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QUESTION What is the efficacy of a closed–loop oxygen control in critically ill patients with moderate to severe acute hypoxemic respiratory failure (AHRF) treated with high flow nasal oxygen (HFNO).

CONCLUSION Closed–loop oxygen control improves oxygen administration in patients with moderate-to-severe AHRF treated with HFNO, increasing the percentage of time in the optimal oxygenation range and decreasing the workload of healthcare personnel.

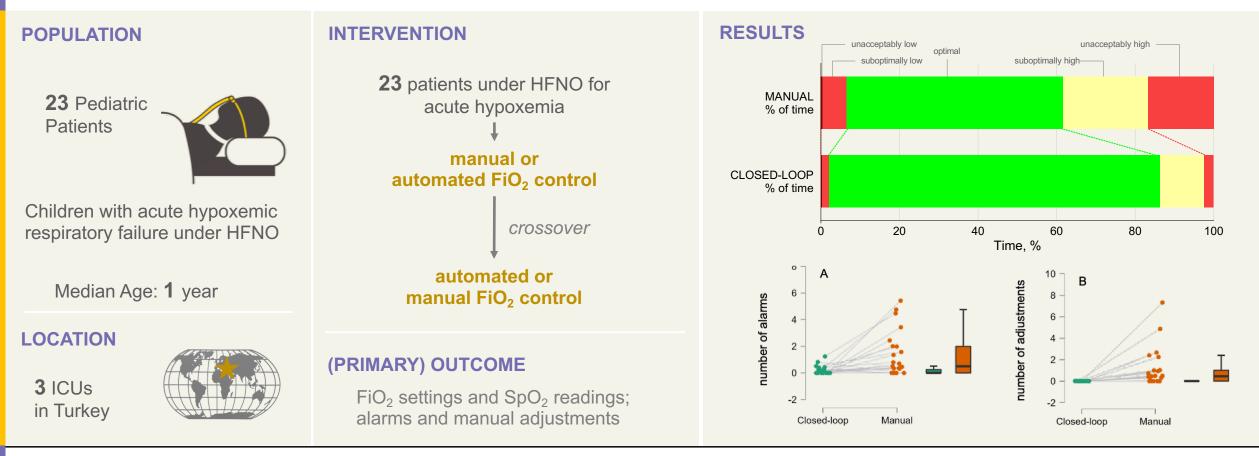
INTERVENTION RESULTS POPULATION SpO2 in optimal range 45 patients under HFNO 100% **45** patients p<0.0001 under HFNO 100-4 hours manual or **90** 75% automated FiO₂ control 80of time Percentage of time 70patients with moderate to crossover severe ARF, including patients 60-Percentage 50% with COVID-19 50-4 hours automated or 40-Median Age: 49 year manual FiO₂ control **30** FiO2 range 25% LOCATION 20-SpO2 out of range (high (PRIMARY) OUTCOME Mode Suboptimal SpO2 (high) **Optimal SpO2** 10-Closed-loop oxygen control Suboptimal SpO2 (low) 1 ICU percentage of time spent in pO2 out of range (low) Manual oxygen titration in Spain the individualized optimal SpO₂ ranges Closed-loop Manual oxygen titration Closed-loop Manual oxygen titration

Roca O. Closed–loop oxygen control improves oxygen therapy in acute hypoxemic respiratory failure patients under high flow nasal oxygen (HILOOP): a randomized cross-over study. [*Crit Care* 2022; **26**:108; doi10.1186/s13054-022-03970-w]

frontiers Frontiers in Medicine

QUESTION What is the effect of HFNO with closed–loop control of the fraction of inspired oxygen (FiO₂), compared to HFNO with manual titrations of the FiO₂, on time spent in predefined pulse oximetry (SpO₂) zones in pediatric critically ill patients?

CONCLUSION In this randomized crossover trial in pediatric critically ill patients under HFNO, the percentage of time spent within in optimal SpO₂ zones increased with the use of closed–loop FiO₂ control.

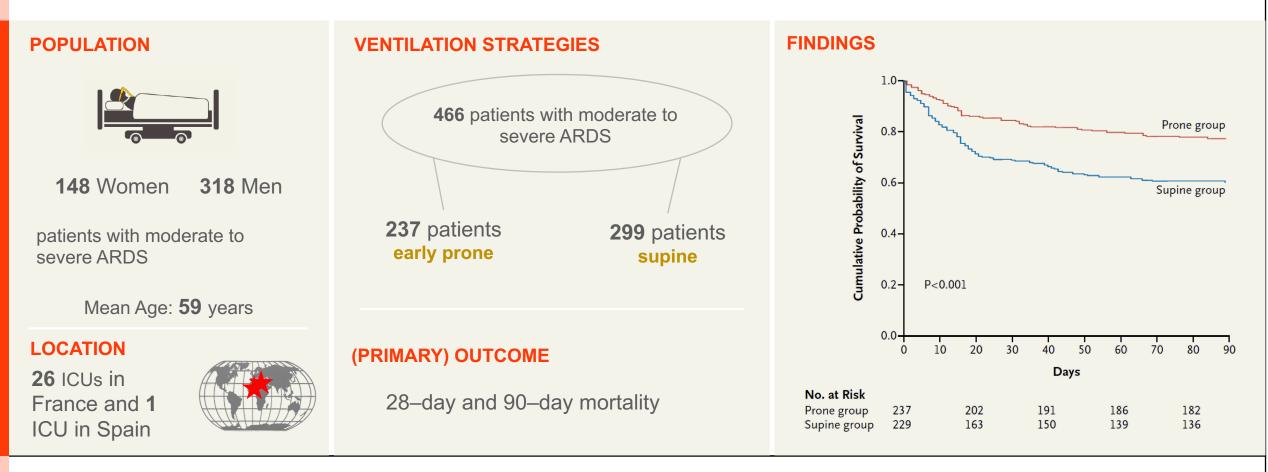


Sandal O. Closed–loop Oxygen Control Improves Oxygenation in Pediatric Patients Under High–flow Oxygen Therapy – a randomized crossover study. [Frontiers Med 2022; **9**:1046902; doi: 0.3389/fmed.2022.1046902]



QUESTION Does early application of prone positioning improves outcomes in patients with severe ARDS?

CONCLUSION In patients with severe ARDS, early application of prolonged prone-positioning sessions significantly decreased 28–day and 90– day mortality.

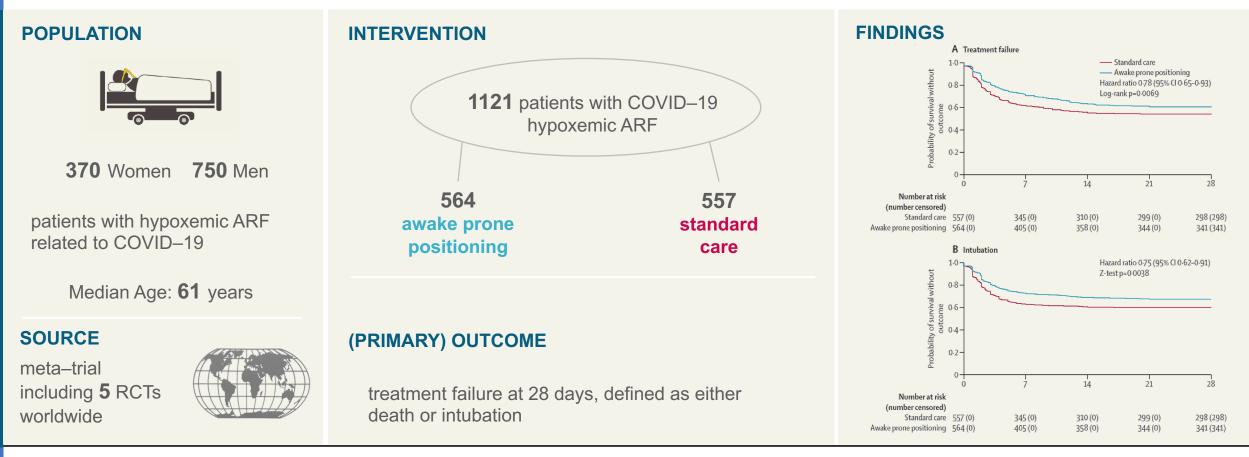


The PROSEVA study group. Prone Positioning in Severe Acute Respiratory Distress Syndrome. [*New Eng J Med* 2013; **368**:2159; doi:10.1056/NEJMoa1214103]

THE LANCET Respiratory Medicine

QUESTION Does awake prone positioning reduce the rate of treatment failure at 28 days, defined as either death or intubation, in patients with severe COVID–19 acute hypoxemic respiratory failure who require respiratory support with HFNO?

CONCLUSION Awake prone positioning of patients with hypoxemic respiratory failure due to COVID–19 reduces the incidence of treatment failure and the need for intubation without any signal of harm.



The Awake Prone Positioning Meta–Trial Group. Awake prone positioning for COVID-19 acute hypoxemic respiratory failure: a randomized, controlled, multinational, open-label meta–trial. [*Lancet Respir Med* 2021; **9**:1387; doi:10.1016/S2213-2600(21)00356-8]

Conclusions

- landscape is changing
- HFNO seems to outperform NIV in ARDS patients
- do not use NIV when $PaO_2/FiO_2 < 150 \text{ mm Hg}$
- be aware of limitations of your oxygen resources when applying HFNO



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