



OUTCOMEREA

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Oxygénothérapie à haut débit (vs. VNI) en réanimation

Etat des connaissances



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**Réanimation des Détrésses Respiratoires
et Infections Sévères**

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Aucun conflit d'intérêt !

Dispositifs médicaux disponibles

Non Humidifié - Non Réchauffé
Contrôle FiO2 peu précis

Humidification - Réchauffement
Contrôle FiO2 précis



Sonde nasale

Canule nasale

Masque facial

MHC

OHD

VNI ou CPAP

Intubation

0.5 à 6 l/min

Max 10 l/min

Max 15 l/min

35 à 60 l/min



Plan

- Effets physiologiques - Tolérance
- HFNC dans l'IRA
- HFNC en post-extubation
 - En réanimation
 - En post-op
- HFNC en pré-intubation

HFNC (high-flow nasal cannula) = OHD (oxygène à haut débit)

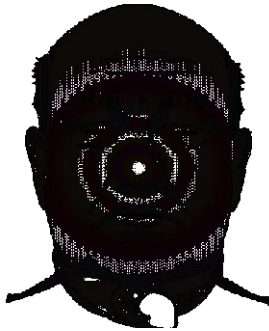
Effets physiologiques -
Tolérance

OHD = Oxygénothérapie à Haut Débit

effets physiologiques

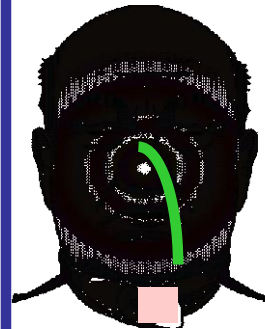
1. Effet de l'OHD sur la FiO₂
2. Effet de l'OHD sur les pressions des voies aériennes (VA)
 - Amélioration des échanges gazeux
 - Amélioration du confort
 - Diminution du travail inspiratoire
3. Conditionnement des Gaz (humidification et réchauffement)
4. « Lavage » de l'espace mort naso-pharyngé

Trachéotomie

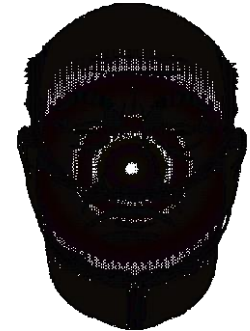


**Systemes
d'oxygénothérapie**

Sonde
nasale



Lunettes
nasales



Masque O2



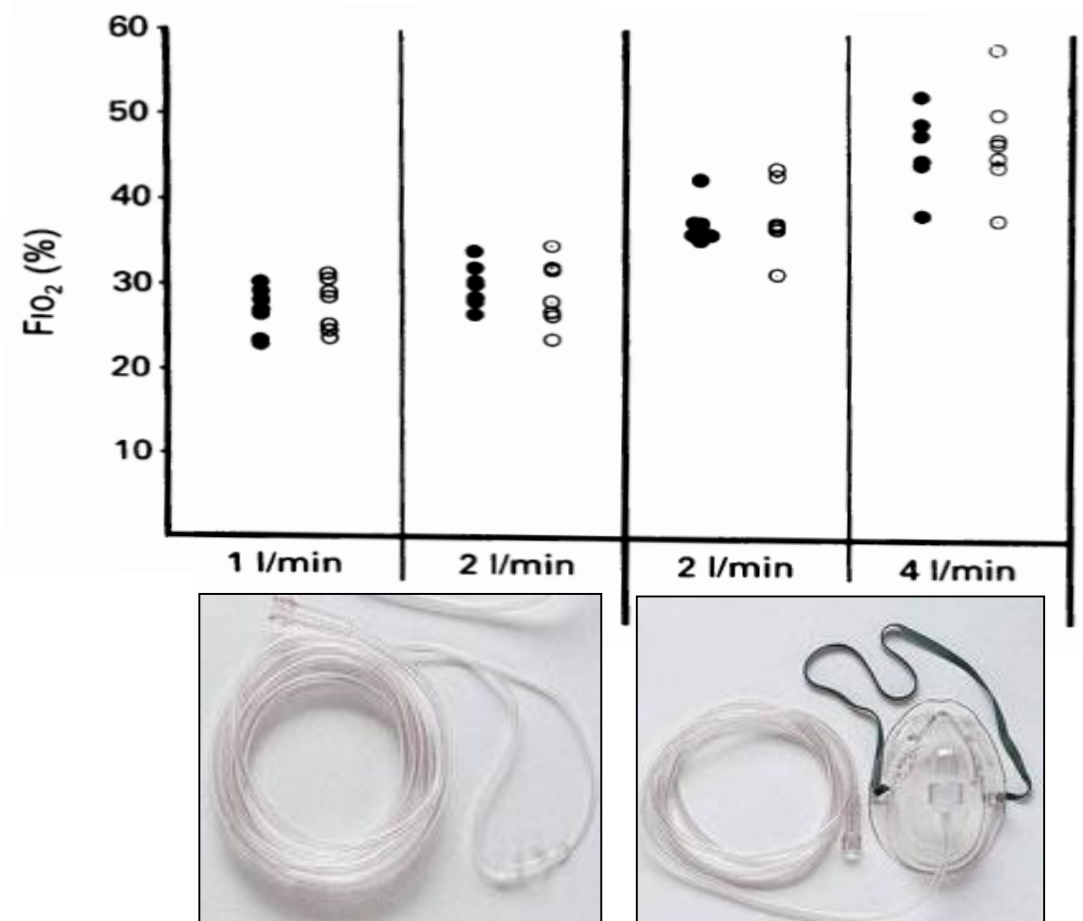
Masque
à venturi



Masque à haute
concentration



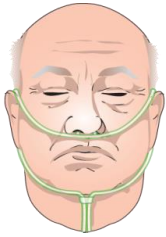
Canule nasale « classique » – Limites d'efficacité



Importante variabilité interindividuelle de la FiO₂ en fonction du débit avec les dispositifs usuels

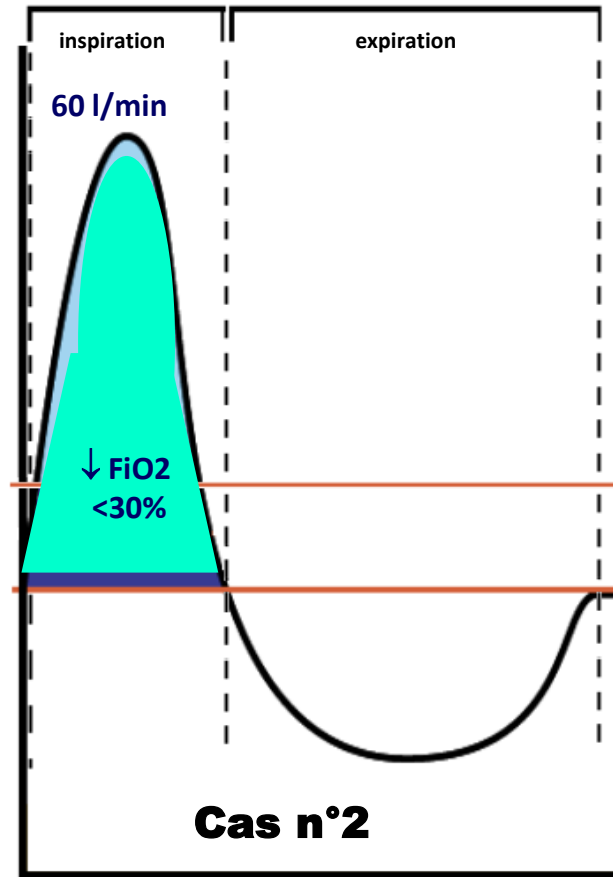
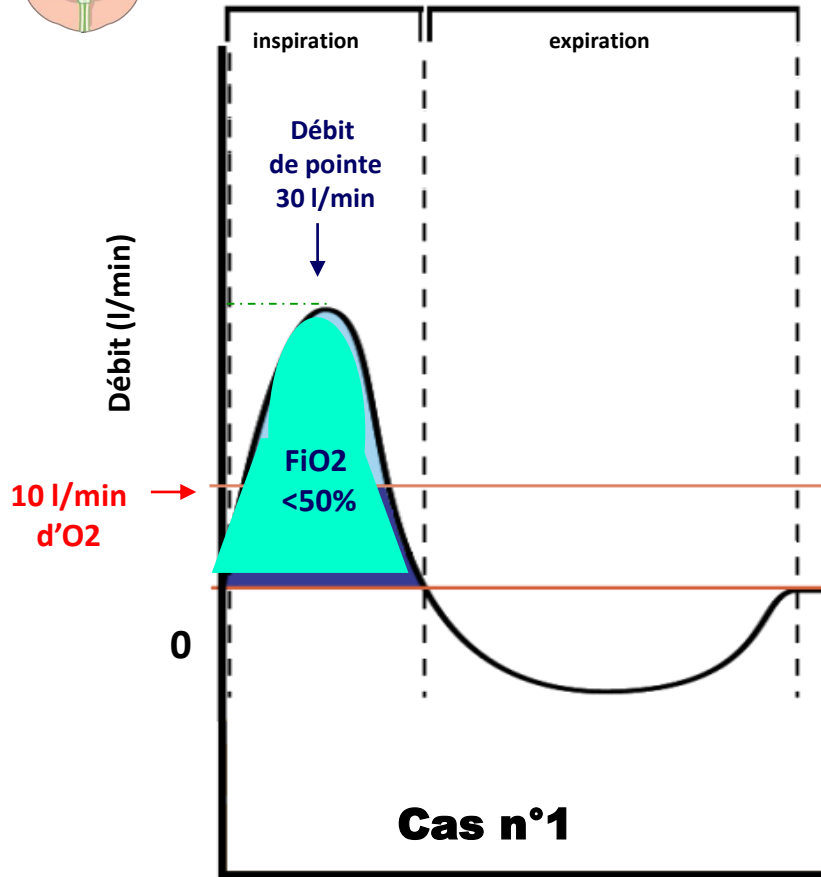
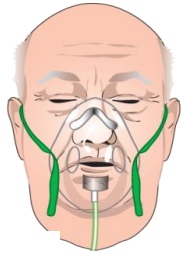


Bazuaye *et al.* Thorax 1992



La « vraie » FiO_2 dépend :

1. débit d'O₂ mural
2. des fuites autour de l'interface
3. ventilation minute du patient



Un mélange à haut débit (→ 60 l/min)



30
|
15
L/min

Fonction oxygénothérapie
sur certains ventilateurs



Débitmètres Air/O2 ou
un mélangeur à haut débit



Oxygénothérapie en cours !

EvitaXL Veille

Marche / Veille

Oxygénothérapie

Attention !
Lors de séances d'oxygénothérapie, n'utilisez que les masques à oxygène. Ne pas utiliser les masques VNI. Durant l'oxygénothérapie, seul un monitoring patient restreint n'est possible. Les paramètres mesurés à partir de débit expiratoire, ex. ventilation minute ou apnée, ne sont plus monitorés.

O2 36 Débit 38

FiO2 36 % Vol.
SpO2 98 %
Pouls 75 cpm

Veille

Oxygénothérapie

P _{mo}	PEP
% Vol.	FI _{O2}
L/min	VM
L/min	VM _{pp}
cpm	f _{res}
L	V _T

FiO2 36 % Vol.
SpO2 98 %
Pouls 75 cpm

VM 15.0
VM_{pp} 2.50
f_{res}
V_T 800

OHD = Oxygénothérapie à Haut Débit

effets physiologiques

1. Effet de l'OHD sur la FiO₂

2. Effet de l'OHD sur les pressions des voies aériennes

- **Amélioration des échanges gazeux**

- **Amélioration du confort**

- **Diminution du travail inspiratoire**

3. Conditionnement des Gaz (humidification et réchauffement)

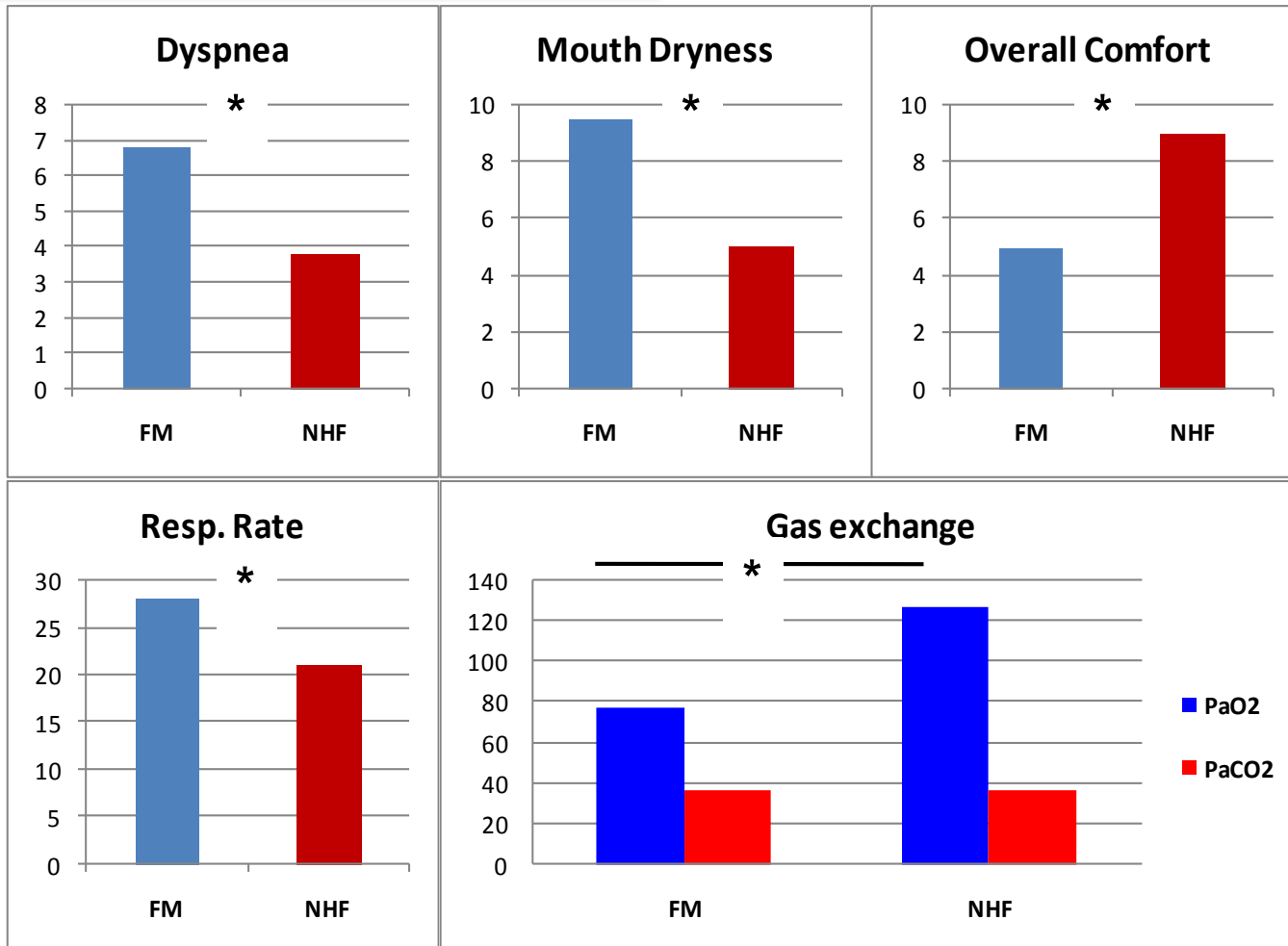
4. « Lavage » de l'espace mort naso-pharyngé

Amélioration oxygénation et confort

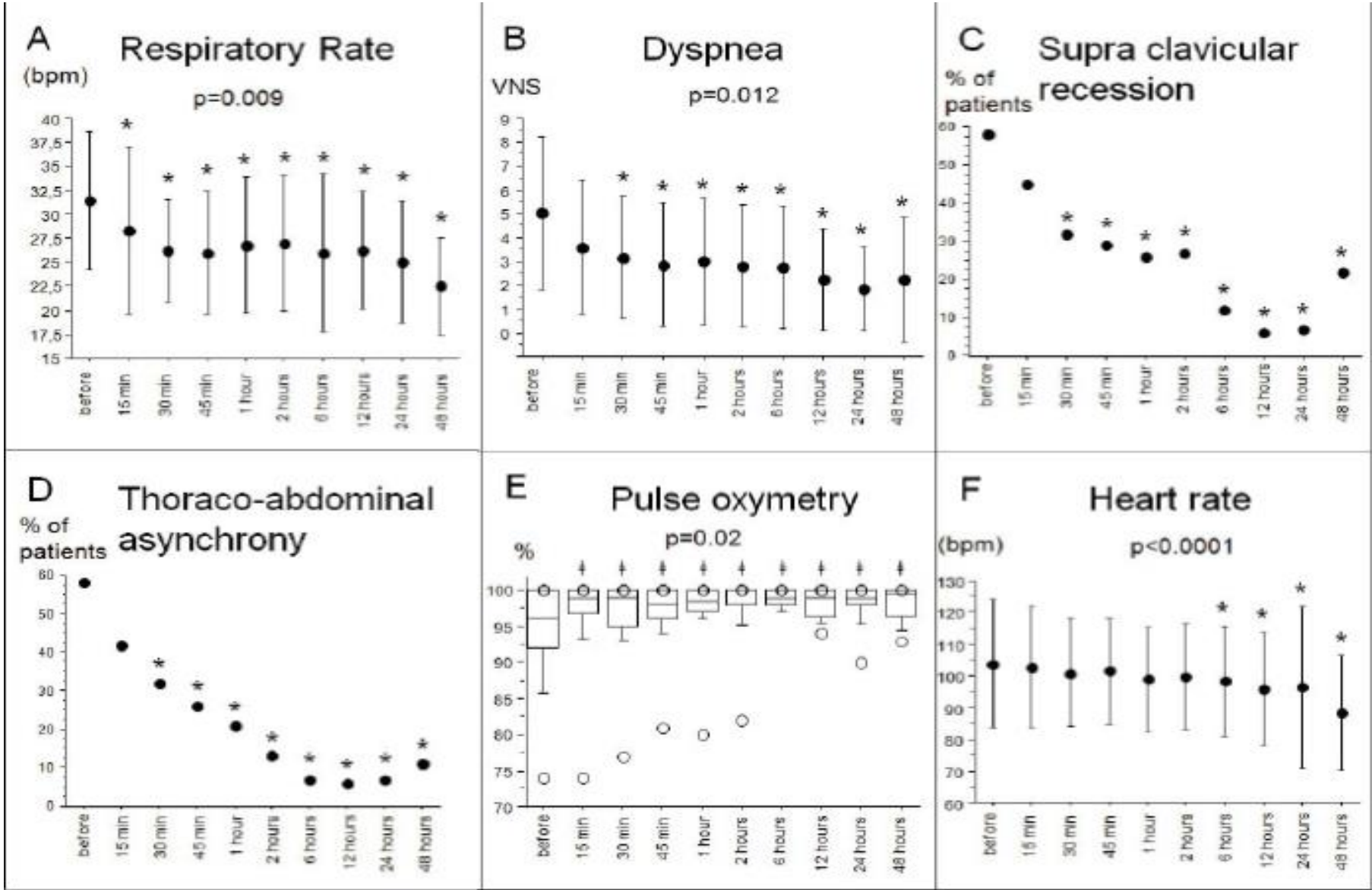
20 patients - SpO₂ < 96% à FiO₂ > 0.5

Venturi Mask vs NHF /30 min

Roca *et al.* Respir Care 2010



ARF, N=38

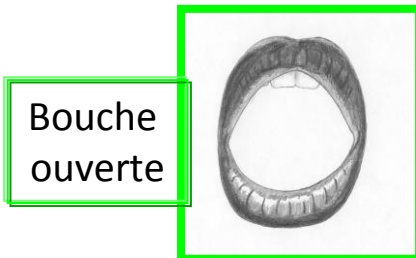
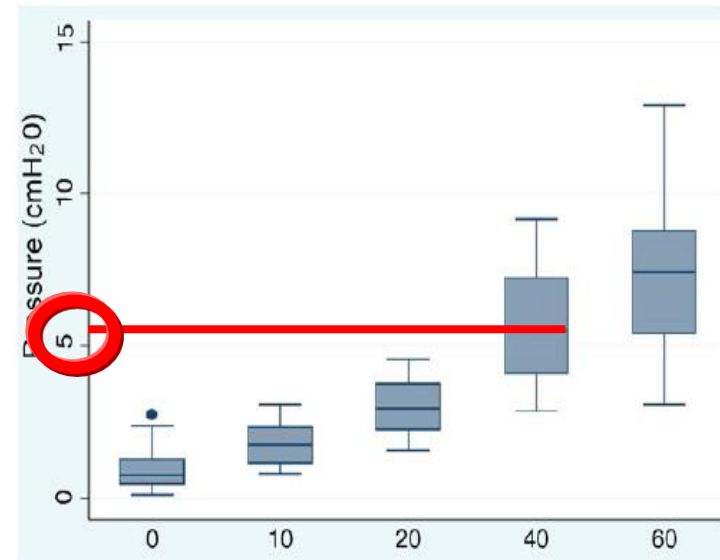
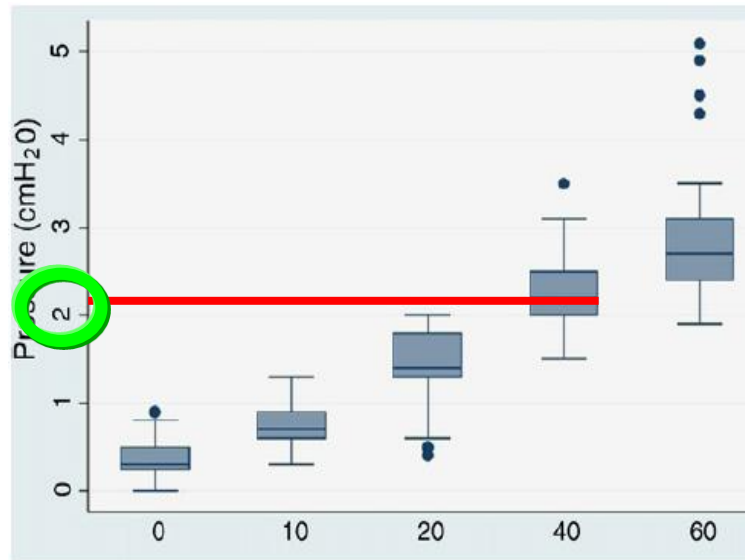


‡ p<0.005 vs before value * p<0.05 vs before value



High flow nasal oxygen generates positive airway pressure in adult volunteers

Nicole Groves Bachelor of Nursing, Post-Graduate Diploma in Advanced Clinical Nursing (Critical Care)*,
Antony Tobin FRACP FJFICM



cathéter nasopharyngé

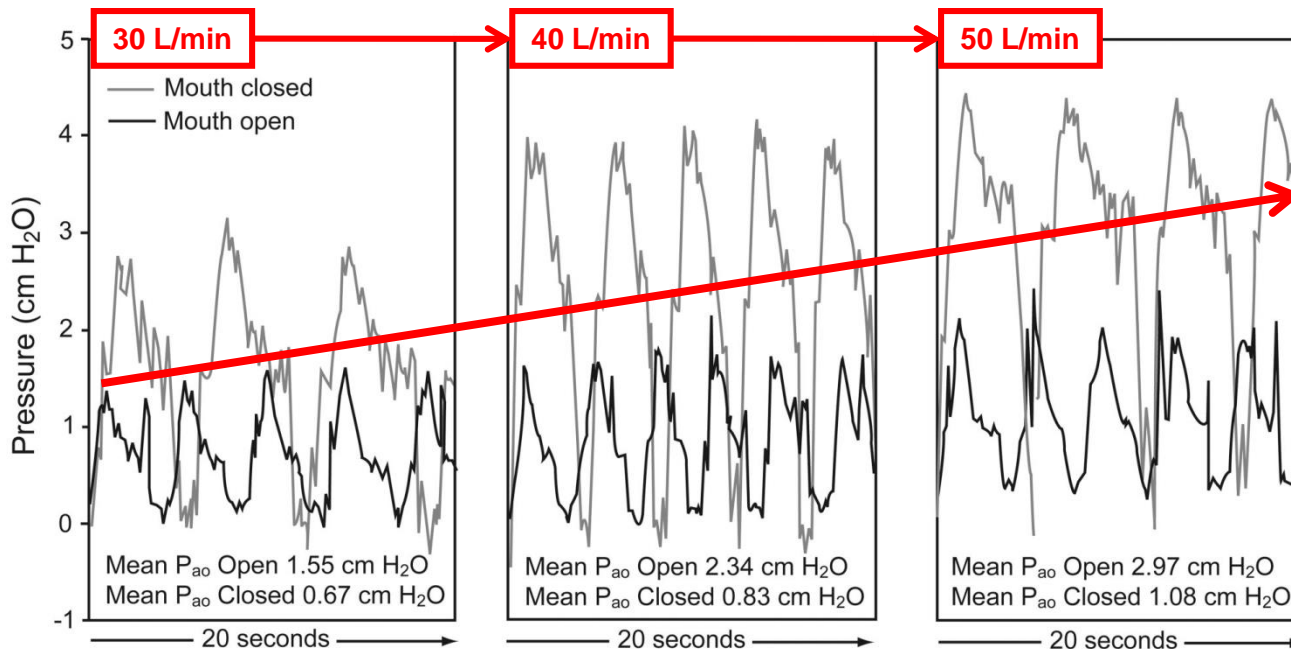


Pression positive proportionnelle au débit

Nasopharyngeal pressure (cmH₂O) [Flow 35 L/min]

	NHF Mouth Closed	NHF Mouth Open	FM Mouth Closed	FM Mouth Open
M	2.7	1.2	0.2	0.1
	±	±	±	±
SD	1.0	0.8	0.6	0.4

Parke et al. Br J Anaesth 2009

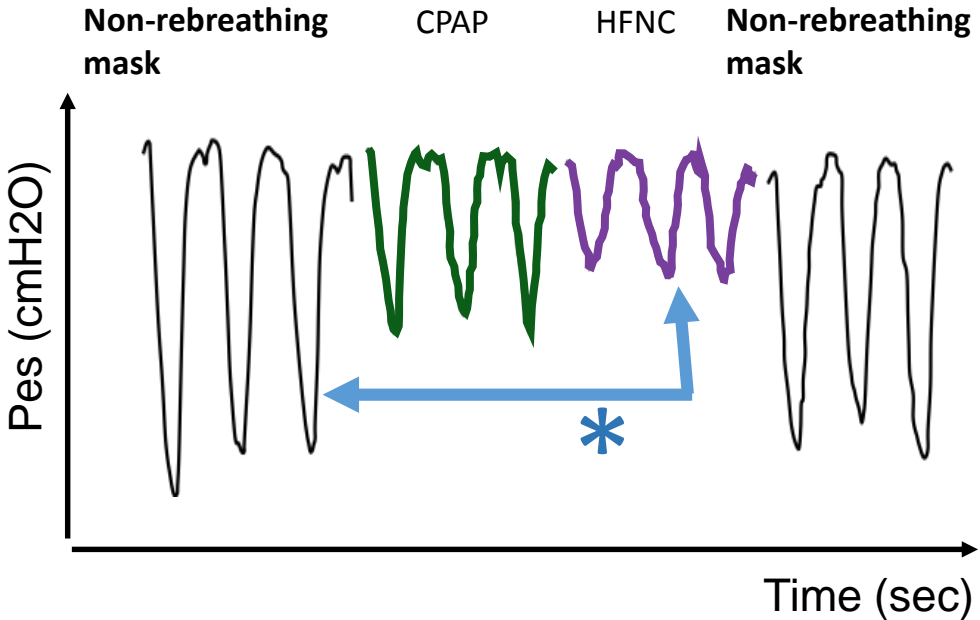


Parke et al. Respir Care 2011

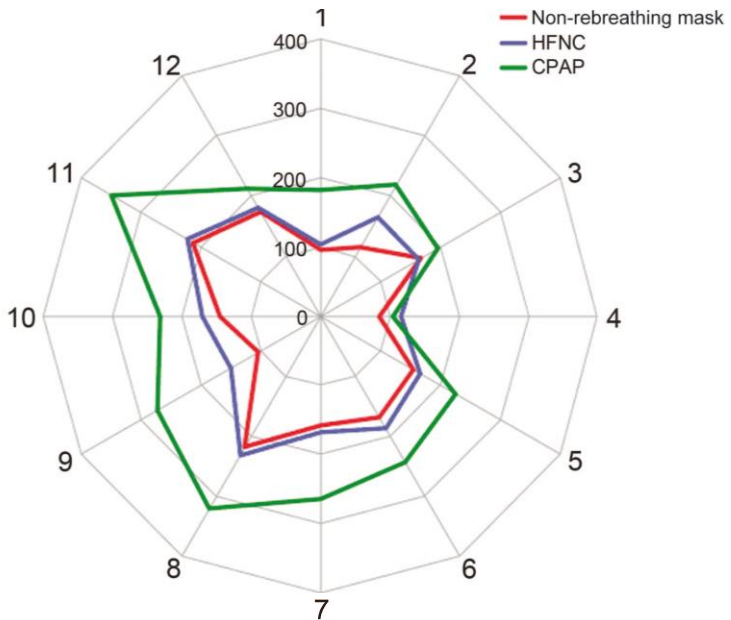
Diminution du travail inspiratoire

Mesure pression oesophagienne (N=12 ARF)

Inspiratory effort



Individual changes in PaO2/FiO2



OHD = Oxygénothérapie à Haut Débit

effets physiologiques

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 - Amélioration des échanges gazeux
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- 3. Conditionnement des Gaz** (humidification et réchauffement)
4. « Lavage » de l'espace mort naso-pharyngé

Importance du réchauffement

Masque facial >
5 L/min



**Humidificateur
Chauffant**

ACTIF

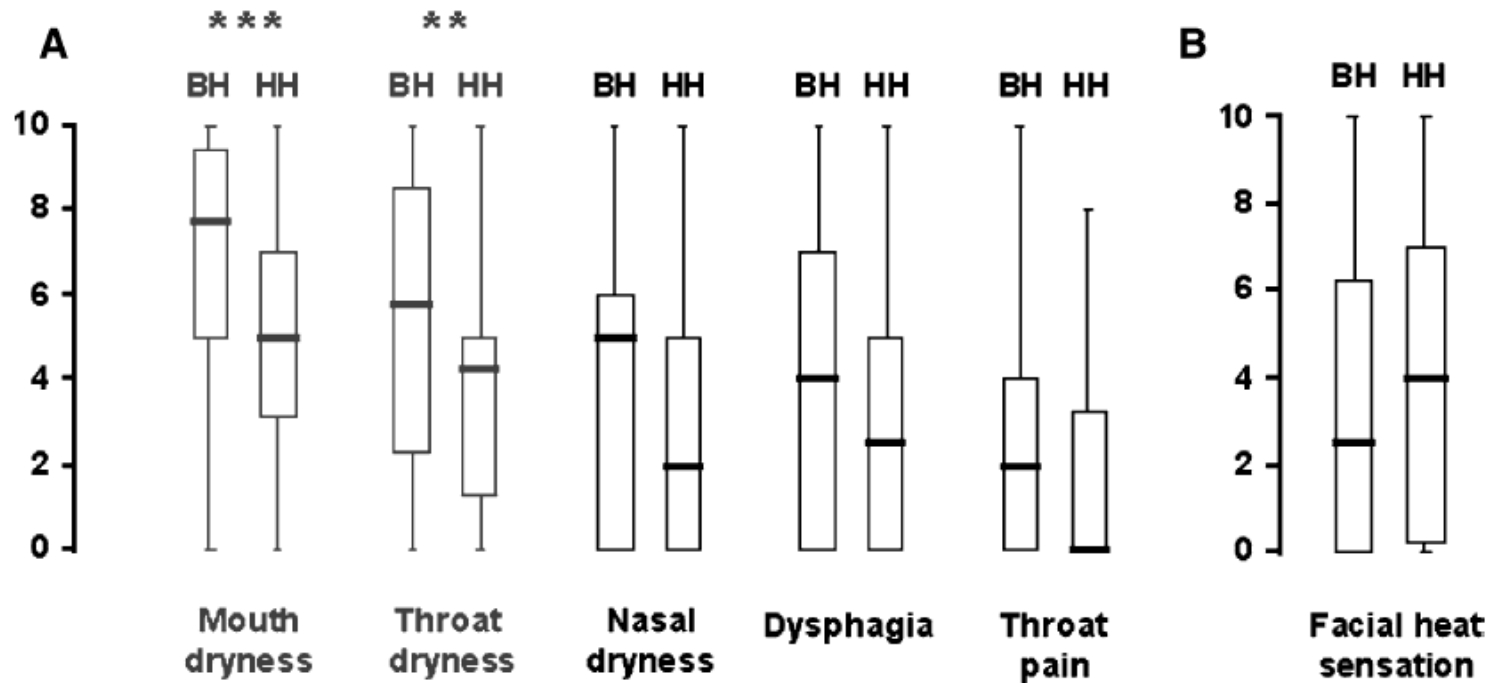


Barboteur

PASSIF



50% des patients ont un inconfort modéré à sévère



Intensité des symptômes pour chaque système

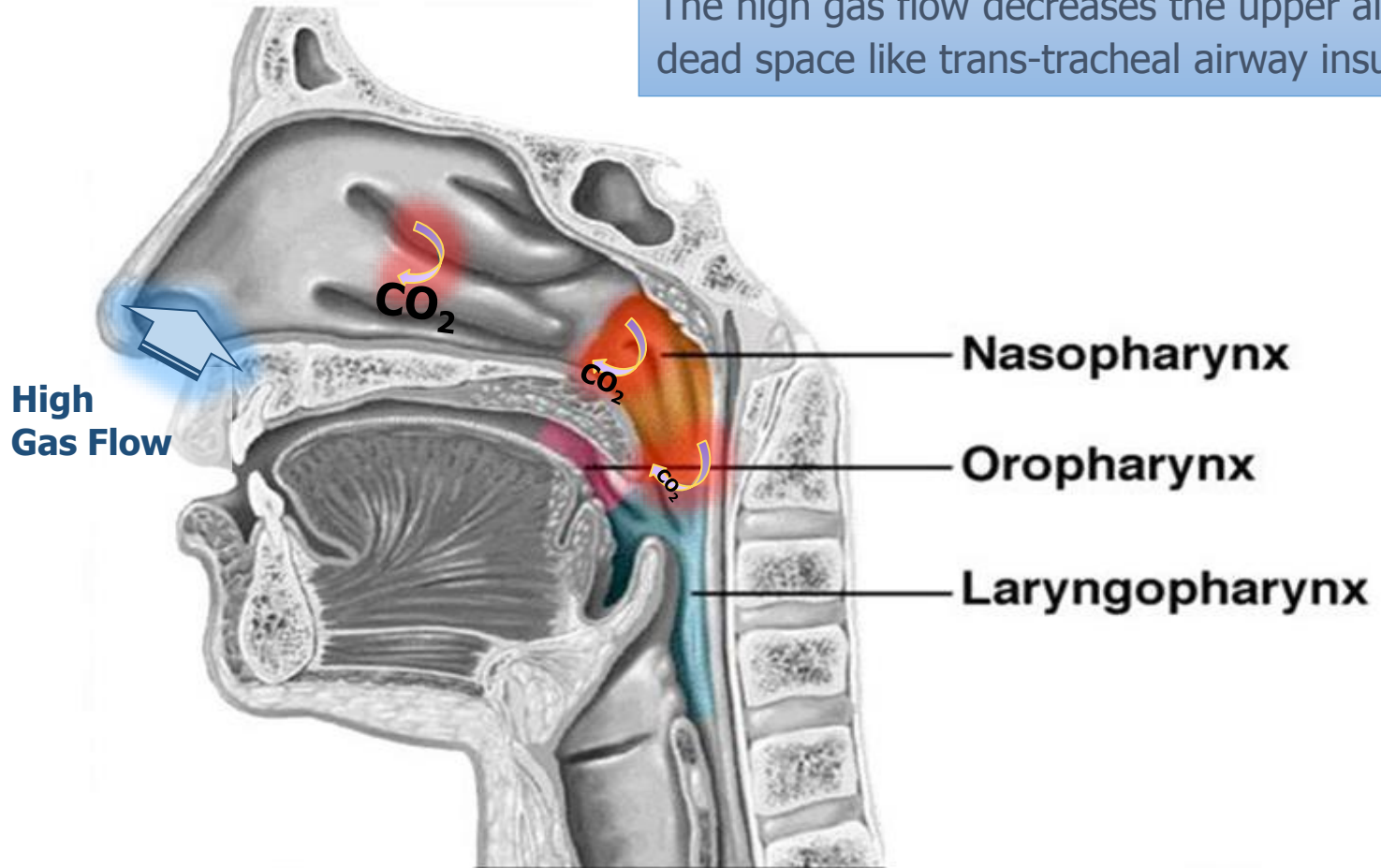
OHD = Oxygénothérapie à Haut Débit

effets physiologiques

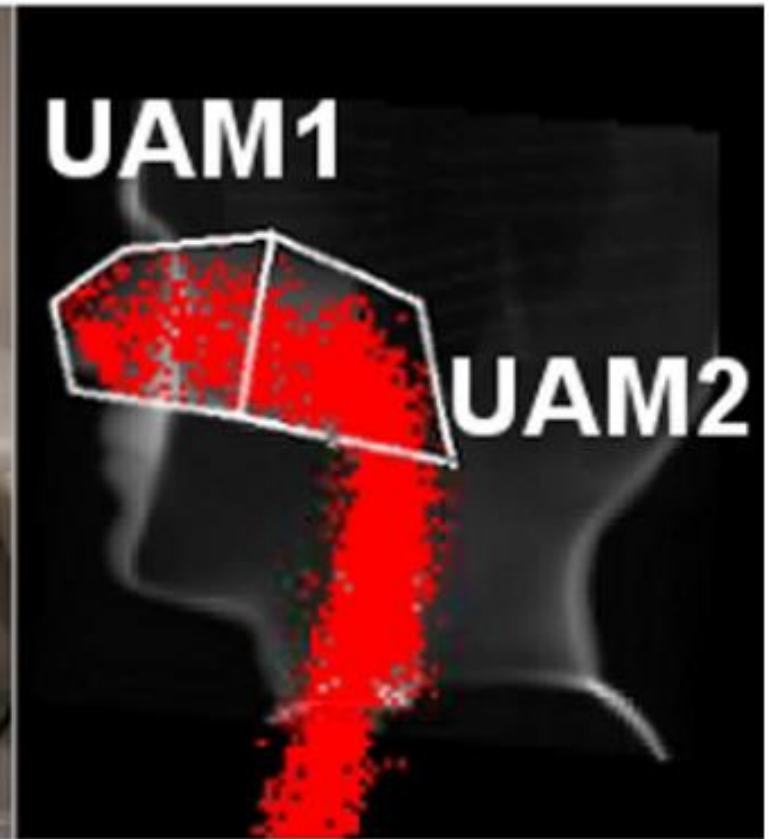
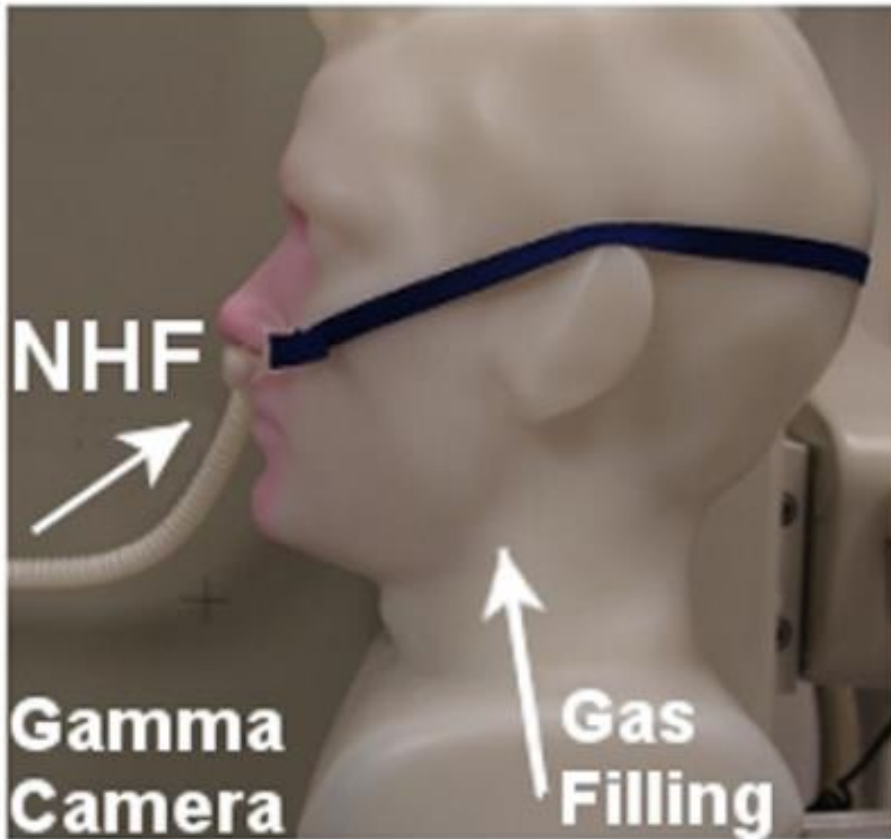
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Lavage espace-mort

The high gas flow decreases the upper airway dead space like trans-tracheal airway insufflation



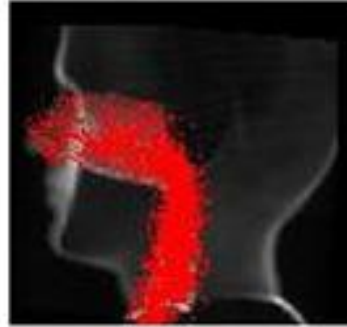
Lavage espace-mort



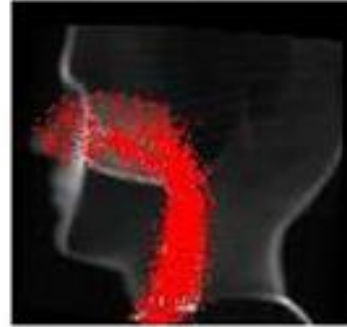
**Elimination
Du CO2**

**15
L/min**

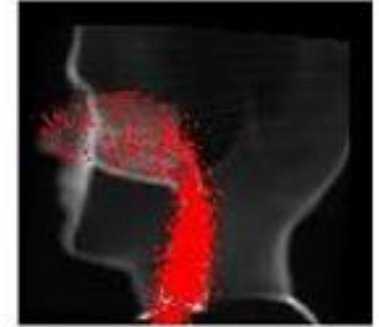
0.5s



1.0s



2.0s



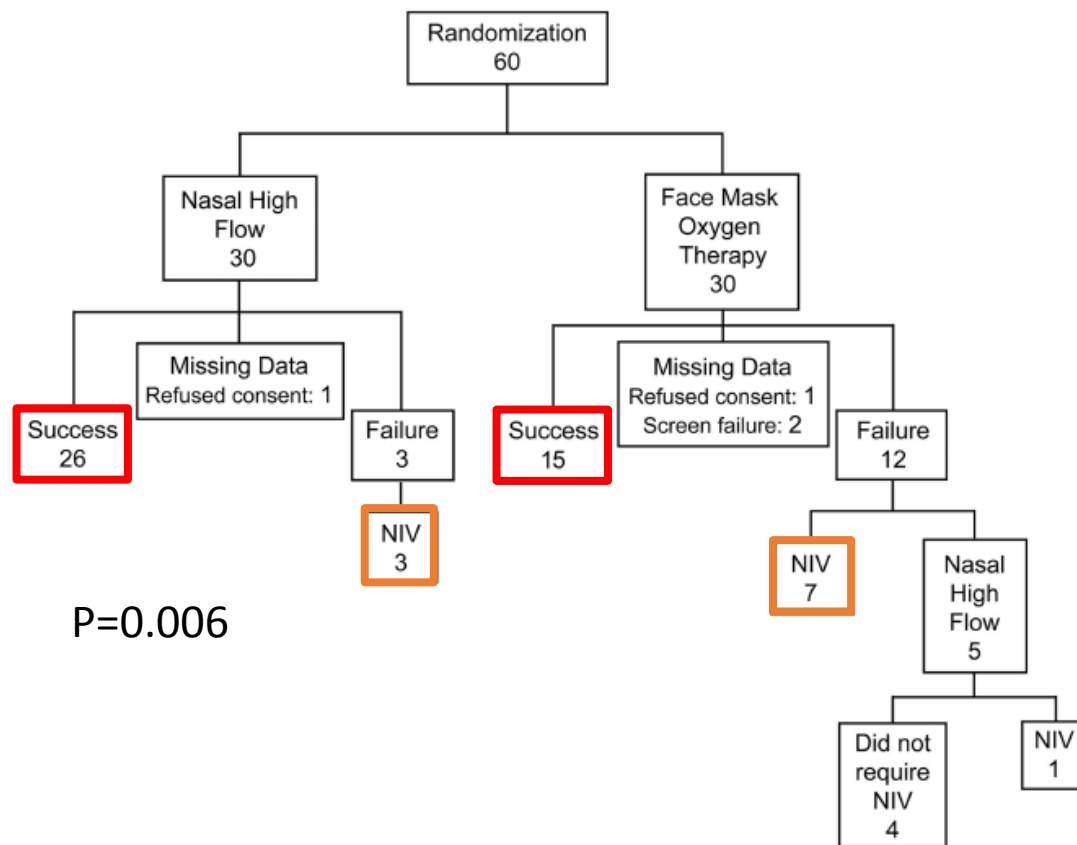
HFNC et IRA

Reference	Study design	Population	N patients	Main results
Hypoxemic acute respiratory failure in the ICU				
Sztrymf ICM 2011	Cohort, unselected patients. HFNC 50 L/min vs. face mask oxygen	Hypoxemic ARF	38	Improved oxygenation Decreased respiratory rate
Sztrymf JCC 2012	Cohort, unselected patients. HFNC 20-30 L/min vs. face mask oxygen	Hypoxemic ARF	20	Improved oxygenation Decrease in respiratory/heart rates, dyspnoea, respiratory distress, and thoraco-abdominal asynchrony
Parke Respir Care 2011	HFNC vs. face mask oxygen	Hypoxemic ARF	60	Decreased treatment failure (defined as need for non-invasive ventilation) from 30% to 10%. Fewer desaturation episodes
Roca Respir Care 2010	Cohort study, HFNC 20-30 L/min vs. face mask oxygen	Hypoxemic ARF	20	Improved comfort; Improved oxygenation
Rello JCC 2012	Cohort study (post hoc)	Hypoxemic ARF (2009 A/H1N1v outbreak)	20	9/20 (45%) success (no intubation). All 8 patients on vasopressors required intubation within 24 hours. After 6 hours of HFNC, non-responders had lower PaO ₂ /FiO ₂ values
Messika Respir Care 2015	Observational, single-centre study	ARDS	45	40% intubation rate. HFNC failure associated with higher SAPSII, development of additional organ failure, and trends toward lower PaO ₂ /FiO ₂ values and higher respiratory rate
Frat NEJM 2015	Multicentre, open-label RCT with 3 groups: HFNC, usual oxygen therapy (face mask), or non-invasive ventilation.	Hypoxemic ARF, PaO ₂ /FiO ₂ ≤ 300	310	Intubation rate was 38% with HFNC, 47% with standard oxygen, and 50% with non-invasive ventilation. Decreased day-90 mortality with HFNC
Nagata Respir Care 2015	Retrospective before/after study of HFNC	Hypoxemic ARF	172	Reduced need for intubation (100% vs 63%, P<0.01)
Kang ICM 2015	Patients intubated after HFNC	Hypoxemic ARF	175	In patients intubated early, lower mortality (39.2 vs. 66.7 %), higher extubation success (37.7% vs. 15.6 %) and more ventilator-free days. Early intubation was associated with decreased ICU mortality.
Hypoxemic acute respiratory failure in the emergency department				
Lenglet Respir Care 2012	Patients with ARF (>9 L/min oxygen or clinical signs of respiratory distress)	Hypoxemic ARF	17	Decreased dyspnoea and respiratory rate and improved oxygenation
Rittayamai Respir Care 2015	RCT of HFNC vs. standard oxygen for 1 h	Hypoxemic ARF	40	Decreased dyspnoea and improved comfort

RCT

A Preliminary Randomized Controlled Trial to Assess Effectiveness of Nasal High-Flow Oxygen in Intensive Care Patients

60 pts (54 cardiac/vacular surgery) with mild to moderate hypoxemic ARF
High-flow face mask vs NHF for 24 h



ORIGINAL ARTICLE

High-Flow Oxygen through Nasal Cannula in Acute Hypoxemic Respiratory Failure

Jean-Pierre Frat, M.D., Arnaud W. Thille, M.D., Ph.D., Alain Mercat, M.D., Ph.D.,
Christophe Girault, M.D., Ph.D., Stéphanie Ragot, Pharm.D., Ph.D.,
Sébastien Perbet, M.D., Gwénael Prat, M.D., Thierry Boulain, M.D.,
Elise Morawiec, M.D., Alice Cottereau, M.D., Jérôme Devaquet, M.D.,
Saad Nseir, M.D., Ph.D., Keyvan Razazi, M.D., Jean-Paul Mira, M.D., Ph.D.,
Laurent Argaud, M.D., Ph.D., Jean-Charles Chakarian, M.D.,
Jean-Damien Ricard, M.D., Ph.D., Xavier Wittebole, M.D., Stéphanie Chevalier, M.D.,
Alexandre Herbland, M.D., Muriel Fartoukh, M.D., Ph.D.,
Jean-Michel Constantin, M.D., Ph.D., Jean-Marie Tonnelier, M.D., Marc Pierrot, M.D.,
Armelle Mathonnet, M.D., Gaëtan Béduneau, M.D., Céline Delétage-Métreau, Ph.D.,
Jean-Christophe M. Richard, M.D., Ph.D., Laurent Brochard, M.D.,
and René Robert, M.D., Ph.D., for the FLORALI Study Group and the REVA Network*

Objectif: déterminer si HFNC \pm la VNI comparées à l'O₂ standard (masque haute concentration) permettent de diminuer le risque d'intubation (J28) et d'améliorer le pronostic des patients admis en réanimation pour IRA

Etude prospective, randomisée, multicentrique

Inclusion : patients avec une IRA

- FR > 25
- P/F < 300 (débit \geq 10 lpm)
- PaCO₂ < 46 mmHg
- Sans pathologie pulmonaire chronique

N = 310

Exclusion dont OAP, neutropénie, vasopresseurs, LATA...

O₂ Standard

Masque haute concentration

Débit O₂ \geq 10 lpm
pour SpO₂ \geq 92%

VS

HFNC

Optiflow®

Canules nasales

Débit 50L/min

FiO₂ pour SpO₂ \geq 92%
Relais O₂ standard

VS

Optiflow®

+ VNI

AI pour Vte = 7-10 ml/kg
et PEEP = 2-10 cmH₂O
FiO₂ pour SpO₂ \geq 92%

Au moins 8h par jour durant
les deux premiers jours

Critères d'intubation

- hemodynamic instability
- deterioration of neurologic status
- signs of persisting or worsening respiratory failure as defined by at least two of the following criteria:
 - a respiratory rate of more than 40 breaths per minute
 - a lack of improvement in signs of high respiratory-muscle workload
 - the development of copious tracheal secretions
 - acidosis with a pH of less than 7.35
 - an SpO₂ of less than 90% for more than 5 minutes without technical dysfunction
 - a poor response to oxygenation techniques
- VNI pré-intubation possible dans les 3 groupes

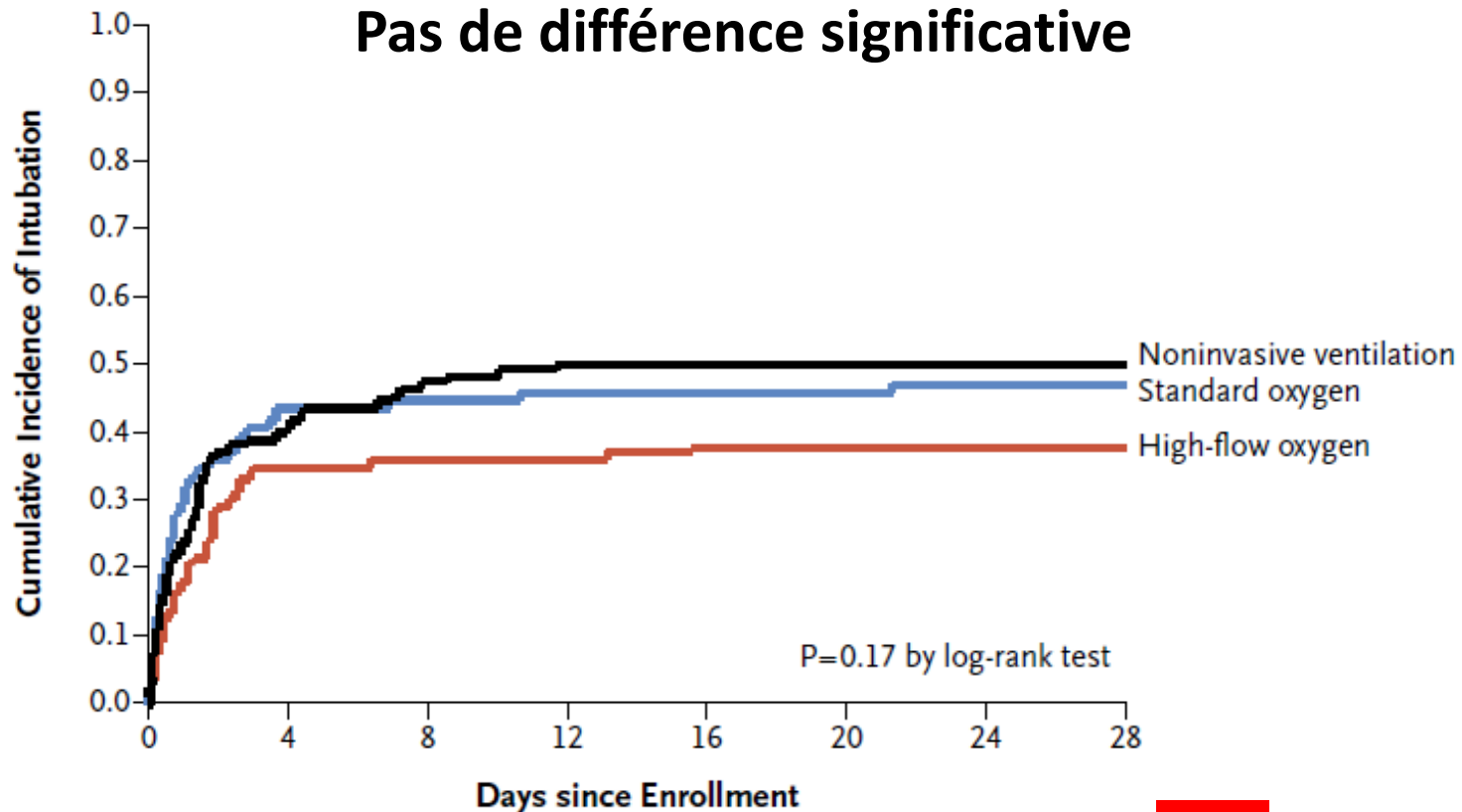
Characteristic	High-Flow Oxygen (N=106)	Standard Oxygen (N=94)	Noninvasive Ventilation (N=110)
Age — yr	61±16	59±17	61±17
Male sex — no. (%)	75 (71)	63 (67)	74 (67)
Body-mass index†	25±5	26±5	26±6
SAPS II‡	25±9	24±9	27±9
Current or past smoking — no. (%)	34 (32)	36 (38)	40 (36)
Reason for acute respiratory failure — no. (%)			
Community-acquired pneumonia	71 (67)	57 (61)	69 (63)
Hospital-acquired pneumonia	12 (11)	13 (14)	12 (11)
Extrapulmonary sepsis	4 (4)	5 (5)	7 (6)
Aspiration or drowning	3 (3)	1 (1)	2 (2)
Pneumonia related to immunosuppression	6 (6)	4 (4)	10 (9)
Other	10 (9)	14 (15)	10 (9)
Arterial blood gas			
pH	7.43±0.05	7.44±0.06	7.43±0.06
Pao ₂ — mm Hg	85±31	92±32	90±36
Fio ₂ §	0.62±0.19	0.63±0.17	0.65±0.15
Pao ₂ :Fio ₂ — mm Hg	157±89	161±73	149±72
Paco ₂ — mm Hg	36±6	35±5	34±6

Résultats

- Réglages initiaux
 - standard-oxygen group 13 ± 5 lpm
 - HFNC 48 ± 11 lpm ($FiO_2: 0,82 \pm 0,21$)
 - VNI
 - AI 8 ± 3 cmH₂O
 - PEEP 5 ± 1 cmH₂O
 - $FiO_2: 0,67 \pm 0,24$
 - $V_t 9,2 \pm 3$ ml/kg
- Durée VNI
 - 8h (IQR 4-12) à J1
 - 8h (IQR 4-13) à J2

Critère de jugement principal = intubation à J28

A Overall Population



No. at Risk

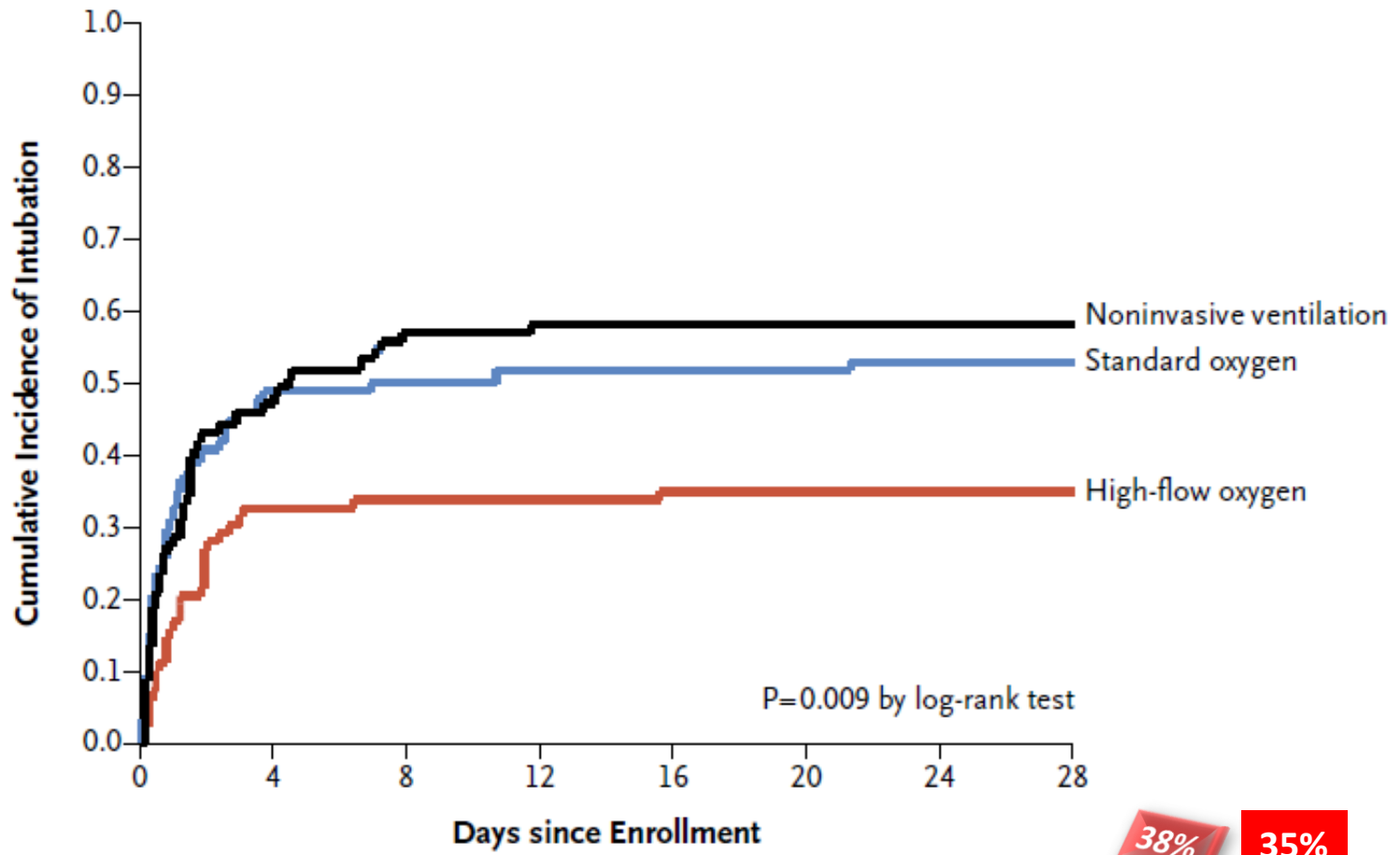
High-flow oxygen	106	68	67	67	65	65	65	65
Standard oxygen	94	52	50	49	49	49	48	48
Noninvasive ventilation	110	64	57	53	53	53	53	52

38%

47%

50%

P/F < 200



No. at Risk

High-flow oxygen	83	55	54	54	53	53	53	53
Standard oxygen	74	37	35	34	34	34	33	33
Noninvasive ventilation	81	41	34	32	32	32	32	32

38%

47%

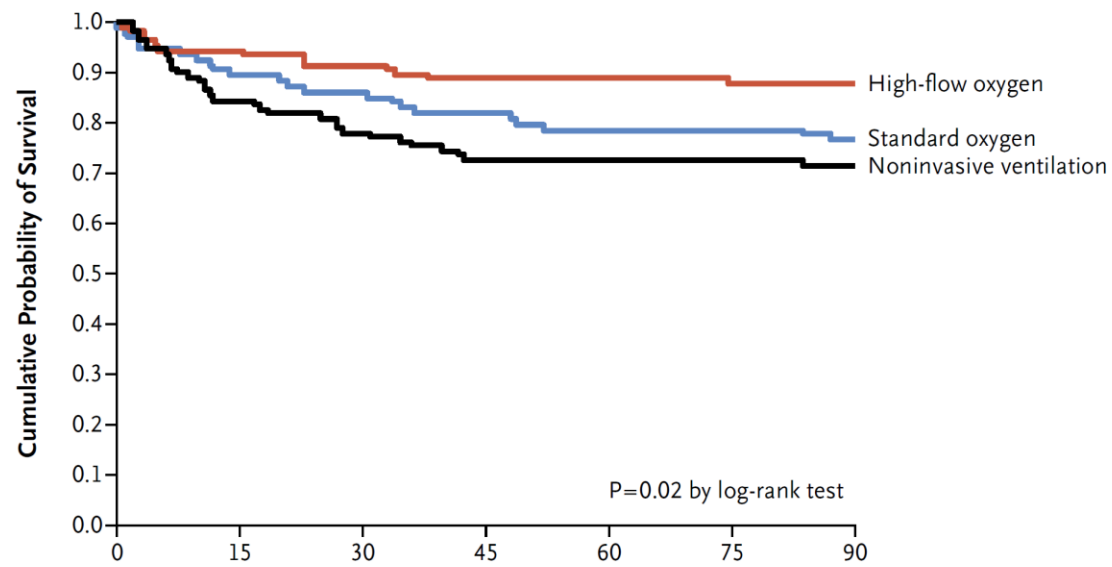
50%

35%

53%

58%

Mortalité



Outcome	Study Group			P Value†	Odds Ratio or Hazard Ratio (95% CI)	
	High-Flow Oxygen (N=106)	Standard Oxygen (N=94)	Noninvasive Ventilation (N=110)		Standard Oxygen vs. High-Flow Oxygen	Noninvasive Ventilation vs. High-Flow Oxygen
Death						
In ICU						
Unadjusted analysis				0.047	1.85 (0.84–4.09)	2.55 (1.21–5.35)
No. of patients	12	18	27			
% of patients (95% CI)	11 (6–19)	19 (12–28)	25 (17–33)			
Adjusted analysis**	—	—	—	—	2.55 (1.07–6.08)	2.60 (1.20–5.63)
At day 90						
Overall population						
Unadjusted analysis				0.02	2.01 (1.01–3.99)	2.50 (1.31–4.78)
No. of patients	13	22	31			
% of patients (95% CI)	12 (7–20)	23 (16–33)	28 (21–37)			
Adjusted analysis**	—	—	—	—	2.36 (1.18–4.70)	2.33 (1.22–4.47)

	Study Group			P Value
	High-Flow Oxygen (N=106)	Standard Oxygen (N=94)	Noninvasive Ventilation (N=110)	
Interval between enrollment and intubation — hr¶				
Overall population				0.27
Median	27	15	27	
Interquartile range	8–46	5–39	8–53	
Patients with $\text{PaO}_2:\text{FiO}_2 \leq 200$ mm Hg				0.32
Median	26	17	27	
Interquartile range	11–46	5–41	7–52	
Death				
Intubated patients				0.16
No. of patients/total. no.	12/40	20/44	27/55	
% of patients (95% CI)	30 (18–46)	45 (32–60)	49 (36–62)	

Ce que l'on peut dire

Conclusion : pas de différence du taux d'intubation mais

- Moins d'intubation qd P/F < 200
- Mortalité diminuée

- Patients sélectionnés : IRA sans hypercapnie et sans pathologie pulmonaire chronique
- Surmortalité dans groupe VNI (associé à HFNC)
 - Retard à l'intubation ?
 - ↗ VILI ($V_t = 9\text{ml/kg}$) ?

Byung Ju Kang
Younsuck Koh
Chae-Man Lim
Jin Won Huh
Seunghee Baek
Myongja Han
Hyun-Suk Seo
Hee Jung Suh
Ga Jin Seo
Eun Young Kim
Sang-Bum Hong

Failure of high-flow nasal cannula therapy may delay intubation and increase mortality

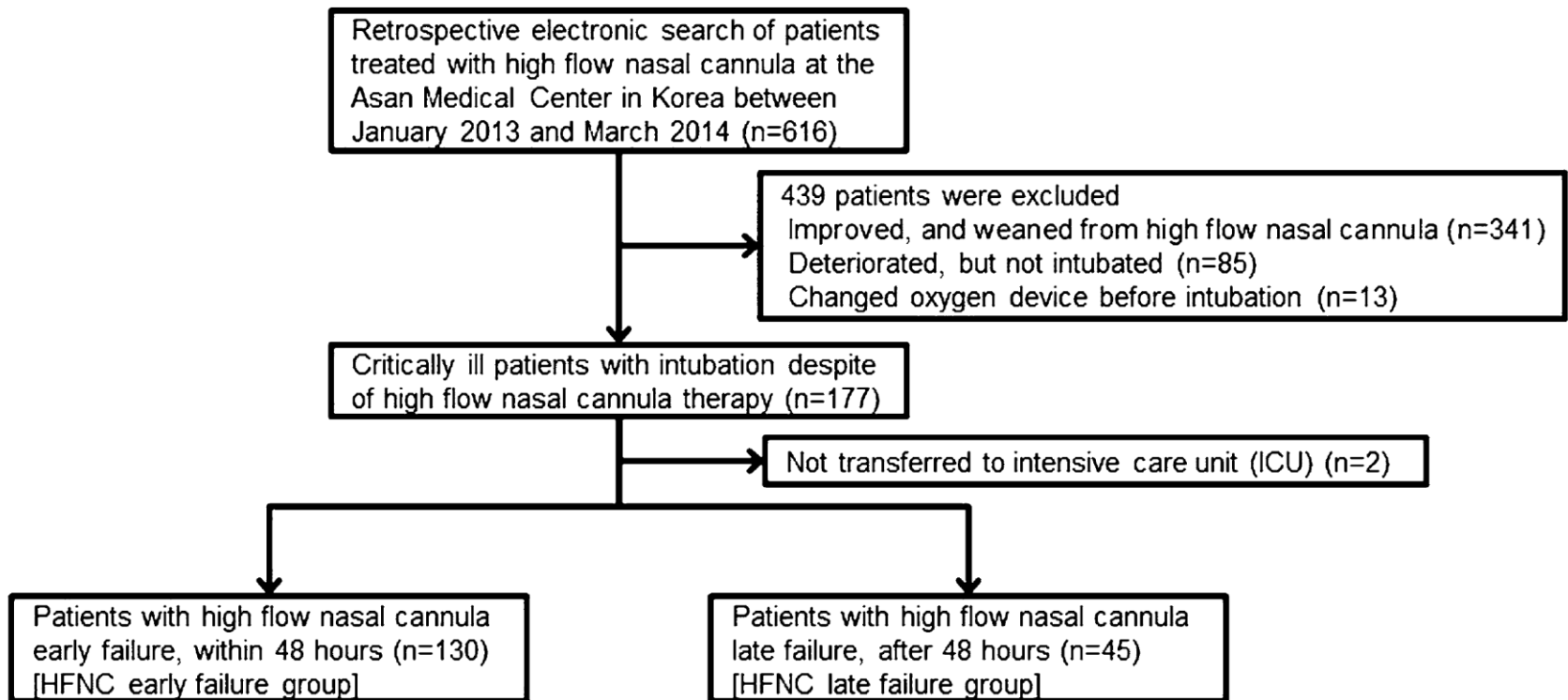
Primary objective

ICU mortality according to the intubation timing in respiratory failure patients who received HFNC that failed

Retrospective, single-center

Indications HFNC

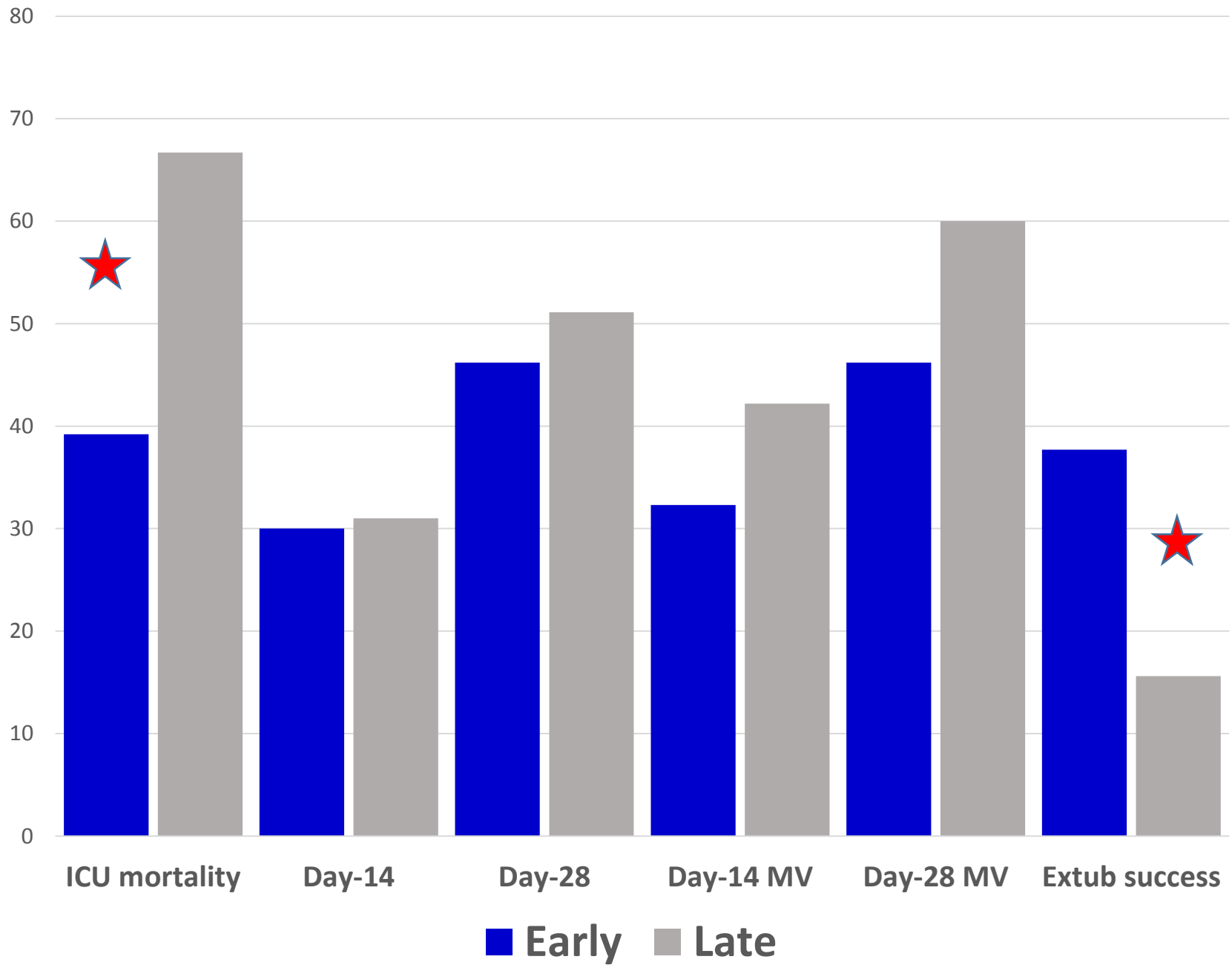
- O₂ standard > 9 lpm pour SpO₂ > 92%
- Signes cliniques de DR
 - RR > 24, mise en jeu muscles resp accessoires, balancement tho-abdo
- Patients à risque après extubation



Characteristics	Early HFNC failure group (<i>n</i> = 130)	Late HFNC failure group (<i>n</i> = 45)	<i>P</i> value ^a
Age, years ^b	66 (56–73)	68 (57–75.5)	0.354
Male sex, <i>n</i> (%)	85 (65.4)	35 (77.8)	0.123
Body mass index, kg/m ² ^b	22.2 (19.8–24.3)	21.1 (18.9–24.0)	0.326
HFNC treatment time, h ^b	10.1 (4.8–22.4)	126.1 (64.9–178.6)	<0.001
Underlying disease			
Diabetes mellitus, <i>n</i> (%)	44 (33.9)	7 (15.6)	0.020
Hypertension, <i>n</i> (%)	53 (40.8)	18 (40.0)	0.928
Solid malignancies, <i>n</i> (%)	34 (26.2)	9 (20.0)	0.409
Hematological malignancies, <i>n</i> (%)	40 (30.8)	15 (33.3)	0.749
Chronic kidney disease/dialysis, <i>n</i> (%)	18 (13.9)	4 (8.9)	0.387
Liver disease, <i>n</i> (%)	24 (18.5)	7 (15.6)	0.660
Use of immunosuppressive agents, <i>n</i> (%)	48 (36.9)	22 (48.9)	0.158
Coronary artery disease, <i>n</i> (%)	13 (10.0)	7 (15.6)	0.313
Heart failure, <i>n</i> (%)	22 (16.9)	8 (17.8)	0.896
Chronic obstructive pulmonary disease, <i>n</i> (%)	15 (11.5)	6 (13.3)	0.749
Tuberculosis-destroyed lung, <i>n</i> (%)	19 (14.6)	12 (26.7)	0.068
PaO ₂ /FiO ₂ before HFNC, mmHg ^b	158.6 (112.7–222.8) (<i>n</i> = 129)	180.0 (138.4–292.0) (<i>n</i> = 43)	0.061
PaO ₂ /FiO ₂ before intubation, mmHg ^b	90.6 (69.7–149.0)	86.3 (64.3–156.7)	0.594
APACHE II score ^b	25.0 (21.0–28.0)	25.0 (21.0–28.5)	0.832
SOFA score ^b	10.0 (7.0–13.0)	7.0 (5.0–11.0)	0.007

Etiologie DR

	Early HFNC failure group (<i>n</i> = 130)	Late HFNC failure group (<i>n</i> = 45)	<i>P</i> value
Etiology of respiratory failure before HFNC application			
Acute de novo respiratory failure, <i>n</i> (%)	43 (33.1)	15 (33.3)	0.975
Acute-on-chronic lung disease, <i>n</i> (%)	37 (28.5)	16 (35.6)	0.372
Cardiogenic pulmonary edema, <i>n</i> (%)	11 (8.5)	3 (6.7)	>0.999
Pulmonary edema due to renal failure, <i>n</i> (%)	5 (3.8)	1 (2.2)	>0.999
Septic shock other than respiratory infection, <i>n</i> (%)	12 (9.2)	3 (6.7)	0.763
After extubation, <i>n</i> (%)	22 (16.9)	7 (15.6)	0.832



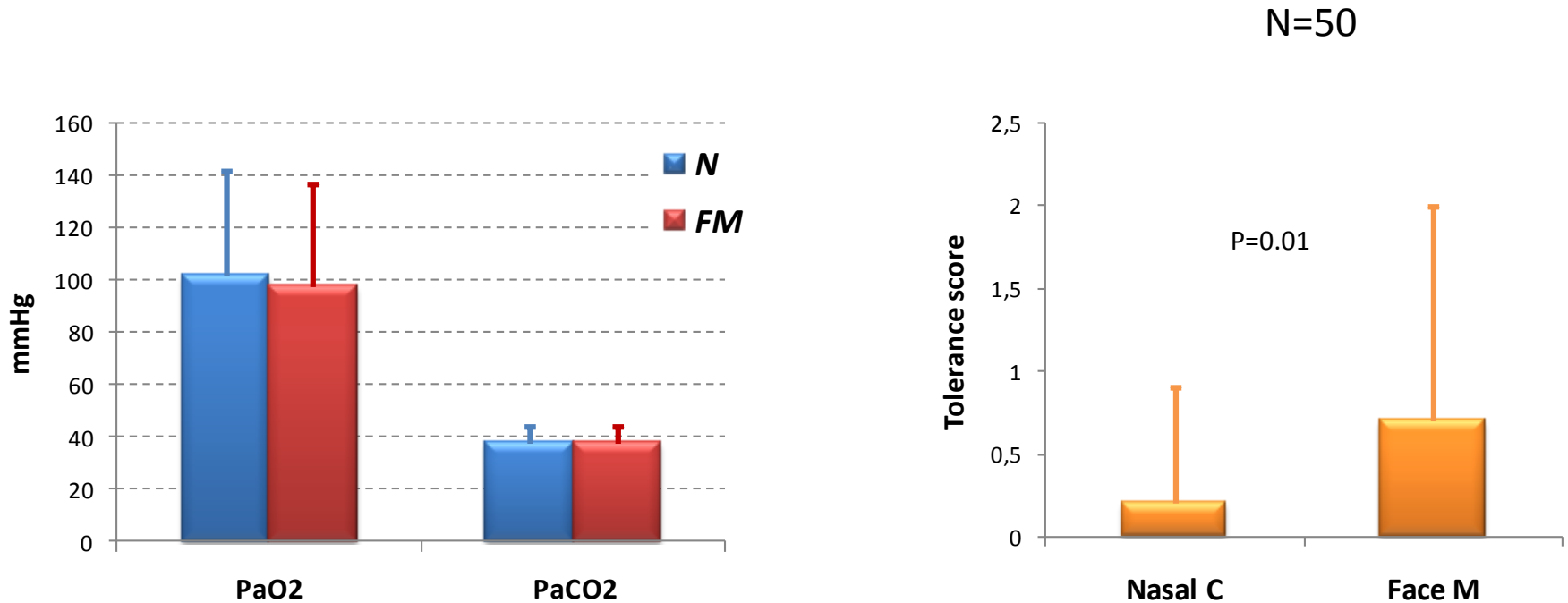
HFNC après extubation

D'un patient de réanimation

High-flow nasal oxygen vs high-flow face mask: A randomized crossover trial in extubated patients

Ravindranath Tiruvoipati*, David Lewis, Kavi Haji, John Botha

Department of Intensive Care Medicine, Frankston Hospital, Frankston, Victoria 3199, Australia



Tiruvoipati et al. J Crit Care 2010

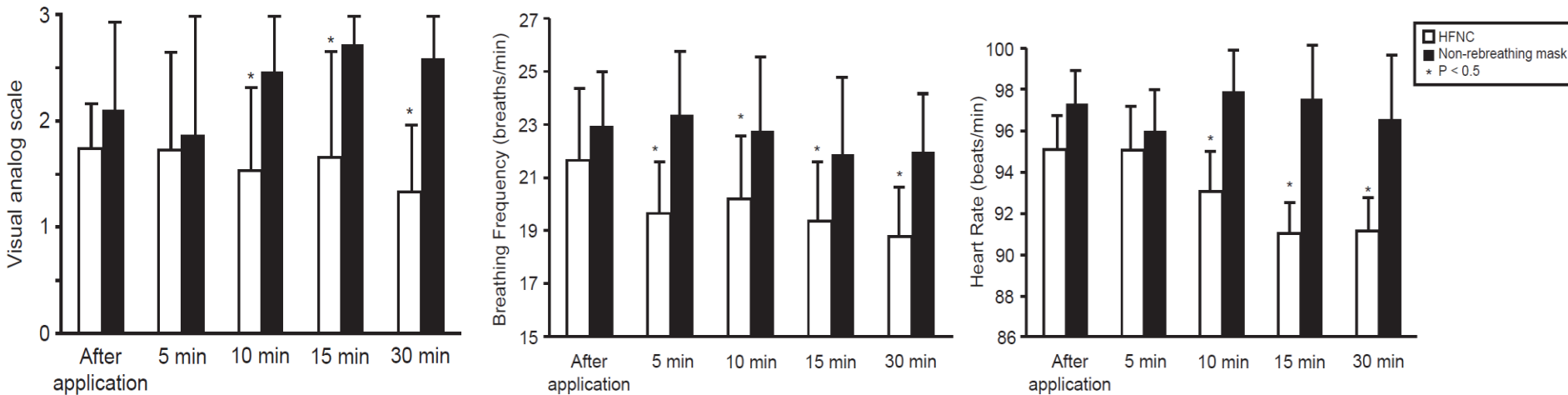
High-Flow Nasal Cannula Versus Conventional Oxygen Therapy After Endotracheal Extubation: A Randomized Crossover Physiologic Study

Nuttapol Rittayamai MD, Jamsak Tscheikuna MD, and Pitchayapa Rujiwit MD

Rittayamai *et al.* Respir Care 2014

17 pts after extubation

HFNC vs non-rebreathing face mask for 30'



Nasal High-Flow versus Venturi Mask Oxygen Therapy

after Extubation

Maggiore *et al.* AJRCCM 2014

Effects on Oxygenation, Comfort, and Clinical Outcome

Salvatore Maurizio Maggiore¹, Francesco Antonio Idone¹, Rosanna Vaschetto², Rossano Festa¹, Andrea Cataldo¹, Federica Antonicelli¹, Luca Montini¹, Andrea De Gaetano³, Paolo Navalesi^{4,5,6}, and Massimo Antonelli¹

- RCT
- Objectif I: P/F à 24h
- 2 centres
- Inclusion :
 - VM > 24h
 - Succès SBT (1 hour: PSV 6-8 cmH2O - PEEP 0, or T-piece)
 - PaO₂/FiO₂ ≤ 300 (fin SBT)
- Exclusion
 - VNI programmée
 - > 3 échecs SBT et PaCO₂ > 45 mm Hg et FR > 25 /min
- Objectif
 - SaO₂ 92-98% (88-95% qd hypercapnie)

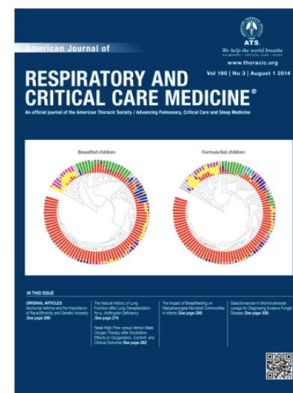
	Control Group (n = 52)	NHF (n = 53)	P Value
Age, yr	64 ± 17	65 ± 18	0.9
Male sex, n (%)	35 (67.3)	33 (62.3)	0.73
SAPS II	44 ± 16	43 ± 14	0.73
Type of admission			0.5
Medical, n (%)	31 (60)	35 (66)	
Surgical-trauma, n (%)	21 (40)	18 (34)	
Cause of acute respiratory failure			0.78
Pneumonia, n (%)	24 (46.2)	24 (45.3)	
Multiple trauma, n (%)	12 (23.1)	11 (20.8)	
Atelectasis, n (%)	5 (9.6)	4 (7.5)	
Shock, n (%)	3 (5.8)	5 (9.4)	
Cardiogenic pulmonary edema, n (%)	3 (5.8)	3 (5.7)	
Cardiac arrest, n (%)	2 (3.8)	3 (5.7)	
Other, n (%)*	3 (5.8)	3 (5.7)	
Length of mechanical ventilation before inclusion, d	5.2 ± 3.7	4.6 ± 4.1	0.43
Length of ICU stay before inclusion	5.6 ± 4.4	5.2 ± 4.4	0.67
PaO ₂ , mm Hg	93.4 ± 24.2	89.9 ± 19.5	0.41
PaCO ₂ , mm Hg	36 ± 7.1	34.7 ± 7.6	0.36
SaO ₂ , %	97.2 ± 2.6	96.9 ± 2.0	0.71
FiO ₂ , %	39 ± 7	38 ± 7	0.47

Nasal High-Flow versus Venturi Mask Oxygen Therapy after Extubation

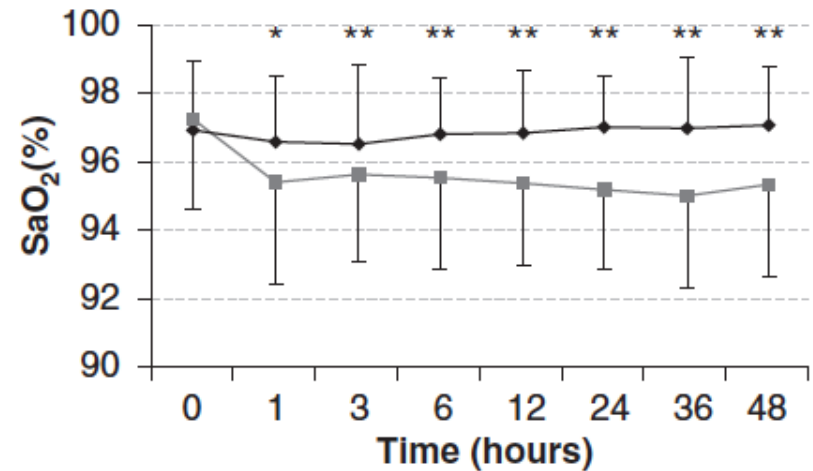
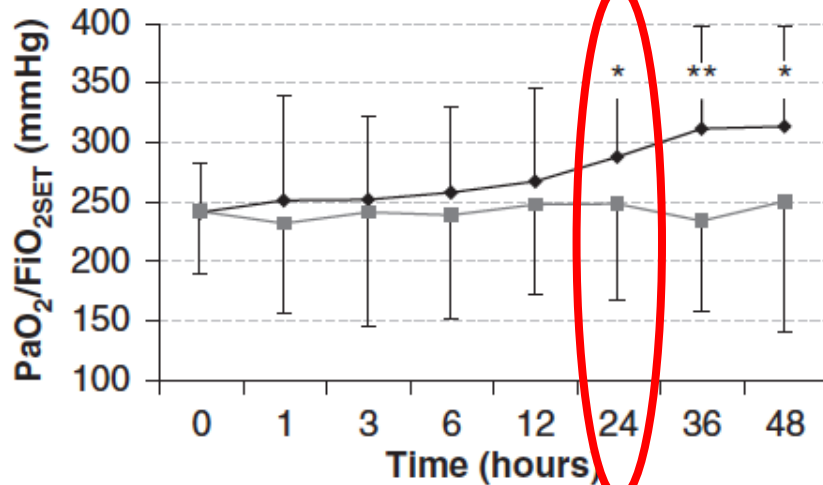
Effects on Oxygenation, Comfort, and Clinical Outcome

Salvatore Maurizio Maggiore¹, Francesco Antonio Idone¹, Rosanna Vaschetto², Rossano Festa¹, Andrea Cataldo¹, Federica Antonicelli¹, Luca Montini¹, Andrea De Gaetano³, Paolo Navalesi^{4,5,6}, and Massimo Antonelli¹

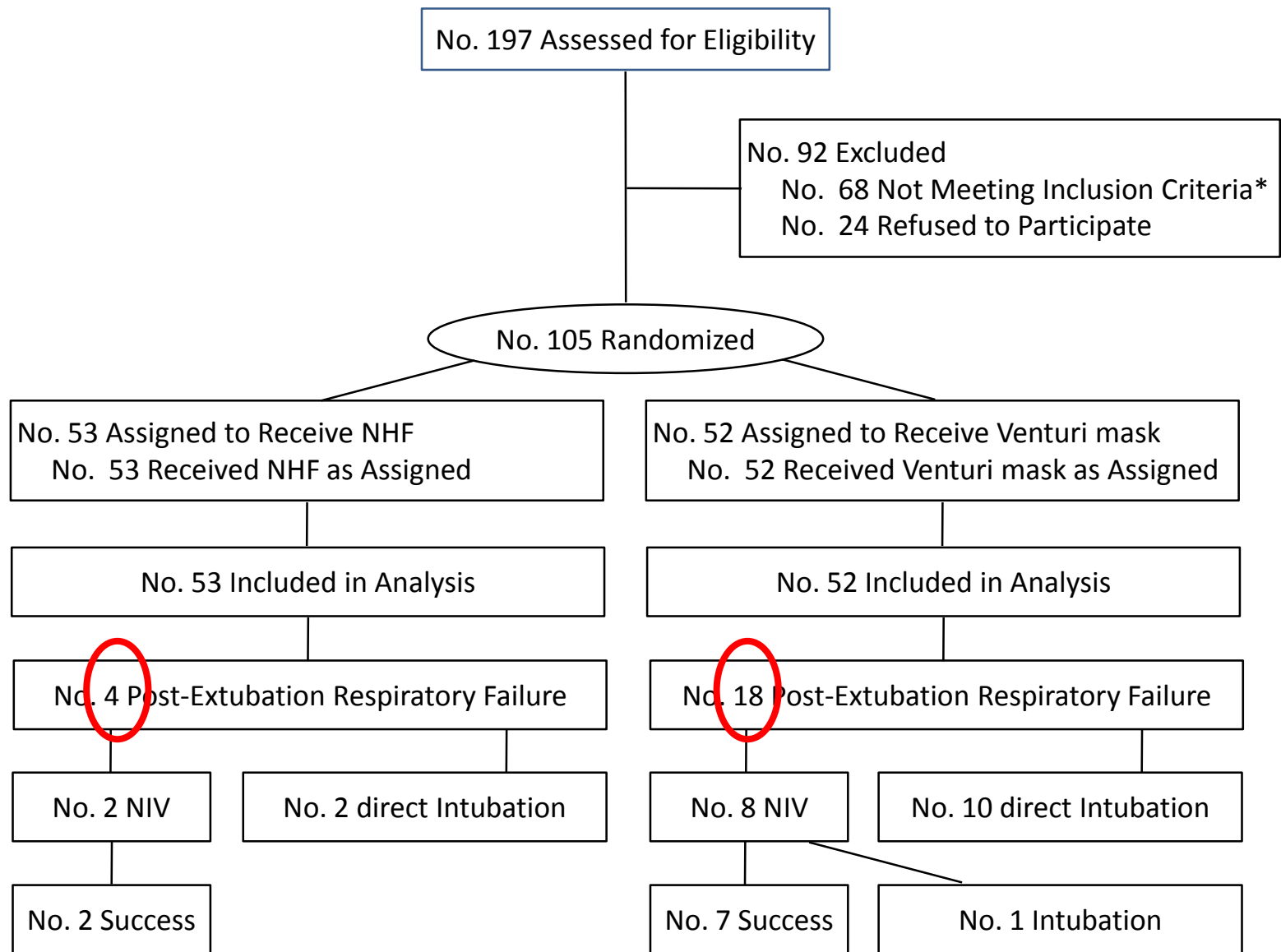
Am J Respir Crit Care Med 2014;190:282–288



Objectif I



◆ NHF ■ Venturi mask



	Control Group (n = 52)	NHF (n = 53)	P Value
Noninvasive ventilation, n (%)	8 (15.4)	2 (3.8)	0.042
Endotracheal intubation, n (%)	11 (21.2)	2 (3.8)	0.005
Cause of endotracheal intubation			
Hypercapnia with respiratory acidosis, n (%)	0	0	N/A
Changes in mental status, n (%)	1 (1.9)	1 (1.9)	0.989
Oxygen desaturation or hypoxia, n (%)	6 (11.5)	1 (1.9)	0.047
Unbearable dyspnea with respiratory muscle failure, n (%)	4 (7.7)	1 (1.9)	0.162
Persistent hypotension, n (%)	2 (3.8)	0	0.149
Inability to clear secretions, n (%)	6 (11.5)	1 (1.9)	0.047

LOS ICU: 11.7 ± 10.2 vs. 10.4 ± 8.5 d (HFNC vs. Venturi mask; P = 0.44)

ICU mortality: 11.3% vs. 9.6%; P = 0.77

The RINO Trial

(ReINtubation rate after Oxygen therapy)

- Multicenter, randomized, controlled, phase III, open trial (NCT02107183)
- 500 patients
- Nasal high-flow vs Venturi mask after extubation
- Study hypothesis: using Optiflow for delivering oxygen therapy after extubation may reduce the extubation failure rate and the need for reintubation as compared with the Venturi mask

HFNC après extubation

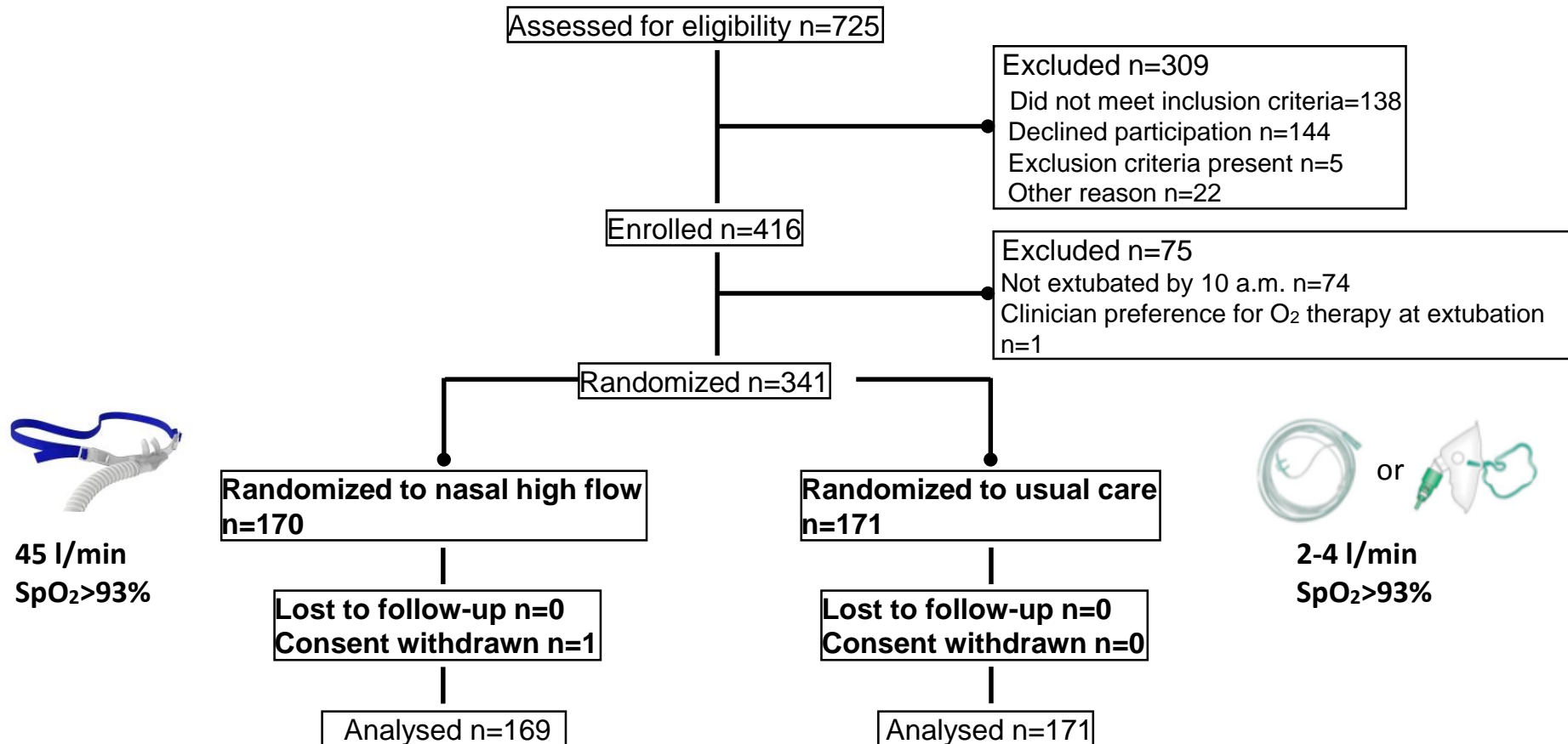
D'un opéré

Open-label, phase II study of routine high-flow nasal oxygen therapy in cardiac surgical patients

Parke *et al.* B J Anaesth 2013

R. Parke^{1,2*}, S. McGuinness¹, R. Dixon^{2,4} and A. Jull^{2,3}

A pragmatic, open-label RCT of patients following extubation after cardiac surgery

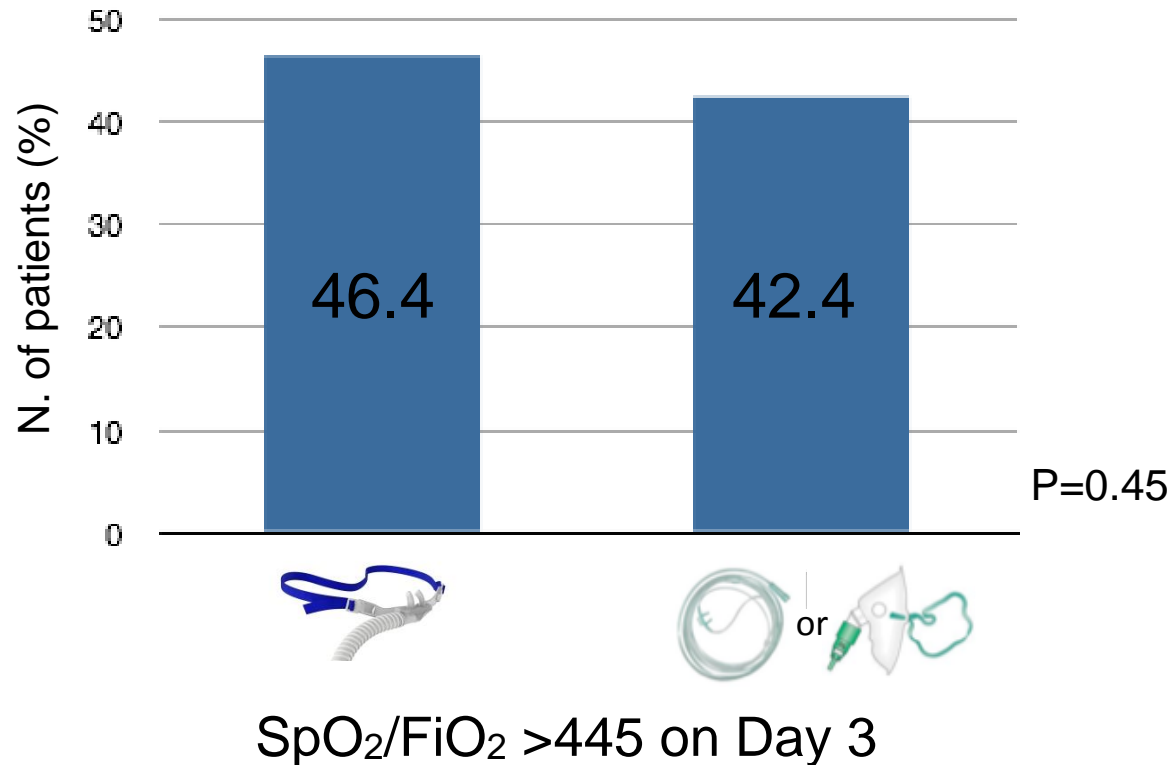


Open-label, phase II study of routine high-flow nasal oxygen therapy in cardiac surgical patients

Parke *et al.* B J Anaesth 2013

R. Parke^{1,2*}, S. McGuinness¹, R. Dixon^{2,4} and A. Jull^{2,3}

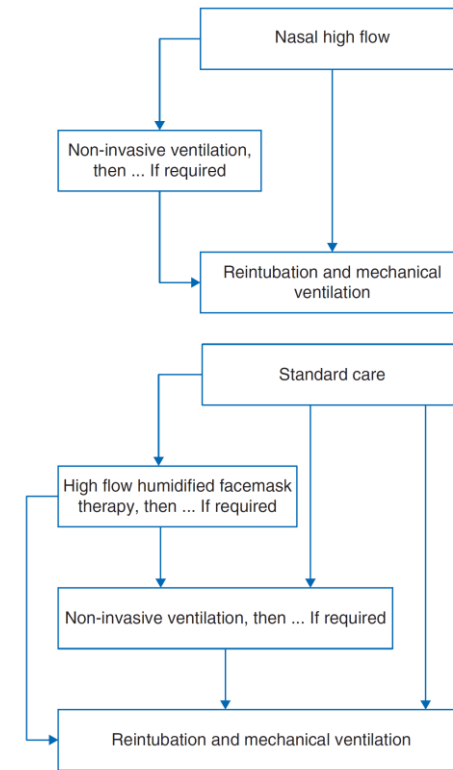
Primary outcome: N. of patients with SpO₂/FiO₂ >445 on Day 3



Open-label, phase II study of routine high-flow nasal oxygen therapy in cardiac surgical patients

R. Parke^{1,2*}, S. McGuinness¹, R. Dixon^{2,4} and A. Jull^{2,3}

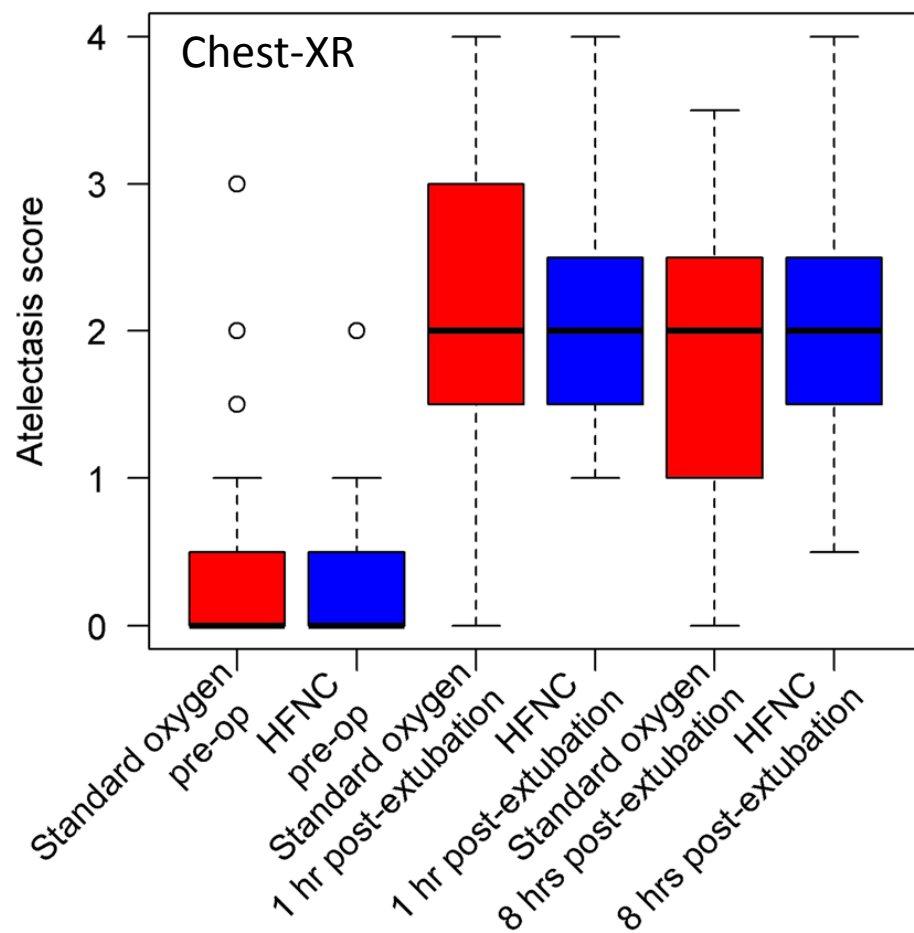
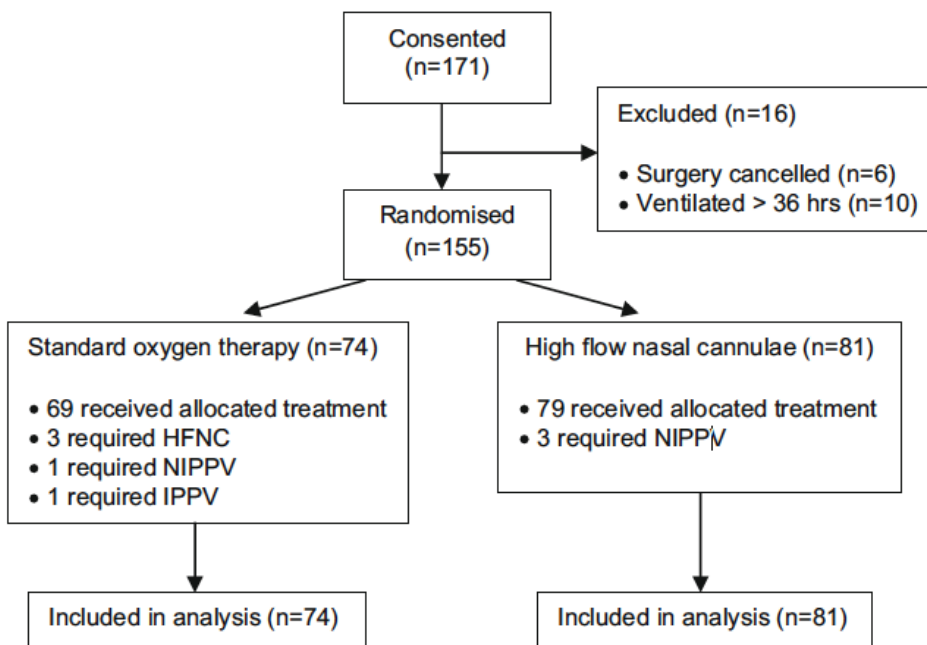
Parke *et al.* B J Anaesth 2013



	Usual care (N=171)	NHF (N=169)	OR (95%CI)	P value
Escalation therapy	77 (45%)	47 (28%)	0.47 (0.29-0.74)	0.001
N (% of those requiring escalation on that day)				
Up to 4 h after extubation	1 (33.3)	2 (66.7)	NR	NR
On Day 1 postoperative	35 (76.1)	11 (23.9)	0.3 (0.1-0.6)	<0.001
On Day 2 postoperative	35 (66)	18 (34)	0.5 (0.3-0.9)	0.01
On Day 3 postoperative	6 (28.6)	15 (71.4)	2.7 (1.0-7.1)	0.04
Reintroduction of oxygen therapy after Day2	54 (31.6)	29 (17.1)	0.45 (0.27-0.75)	0.002
NIV (CPAP or BiPAP)	5 (2.9%)	9 (5.3%)	1.87 (0.6-5.7)	0.27
Intubation	0	2	NR	NR

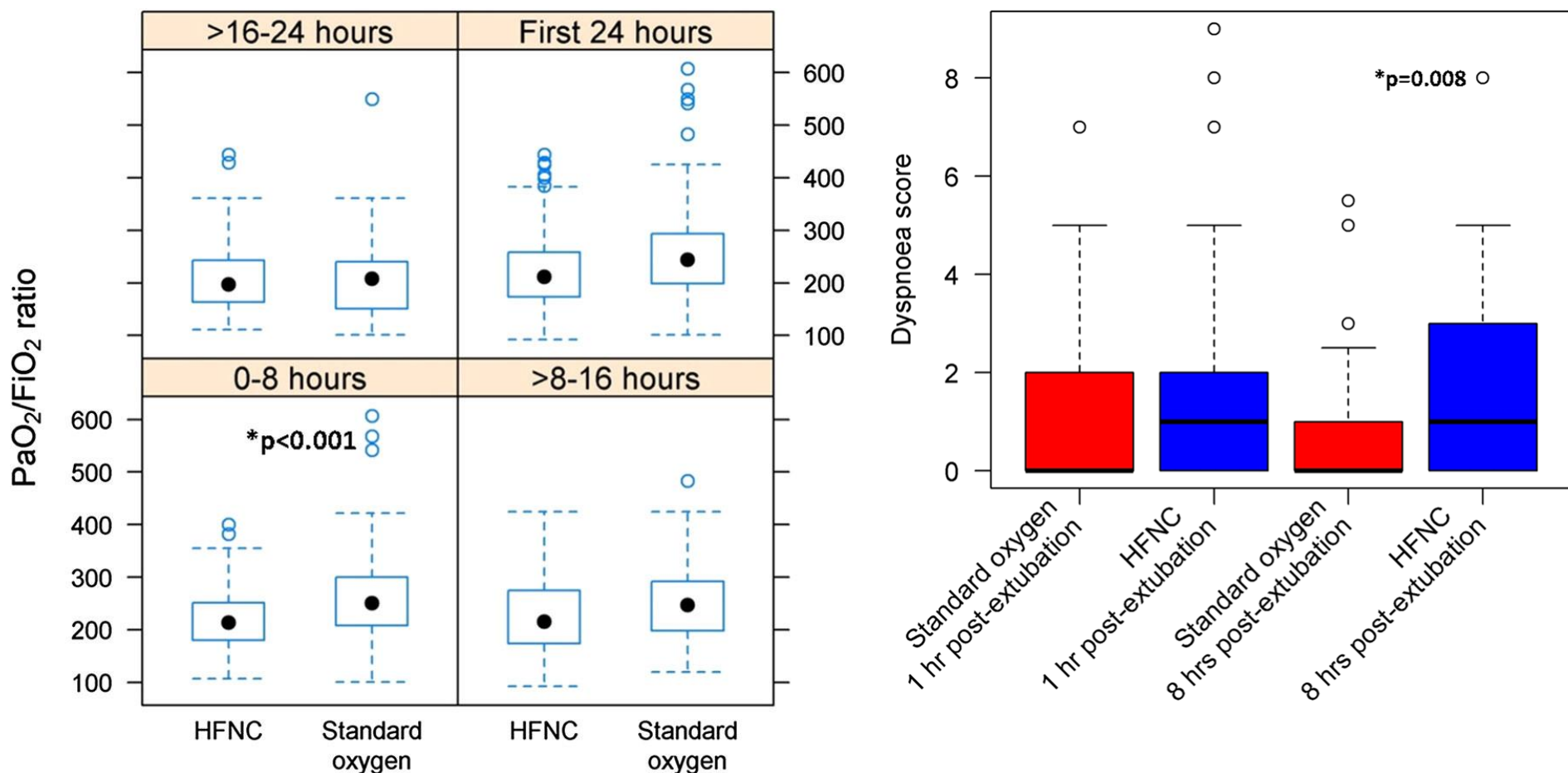
Amanda Corley
Taressa Bull
Amy J. Spooner
Adrian G. Barnett
John F. Fraser

Direct extubation onto high-flow nasal cannulae post-cardiac surgery versus standard treatment in patients with a BMI ≥ 30 : a randomised controlled trial



Amanda Corley
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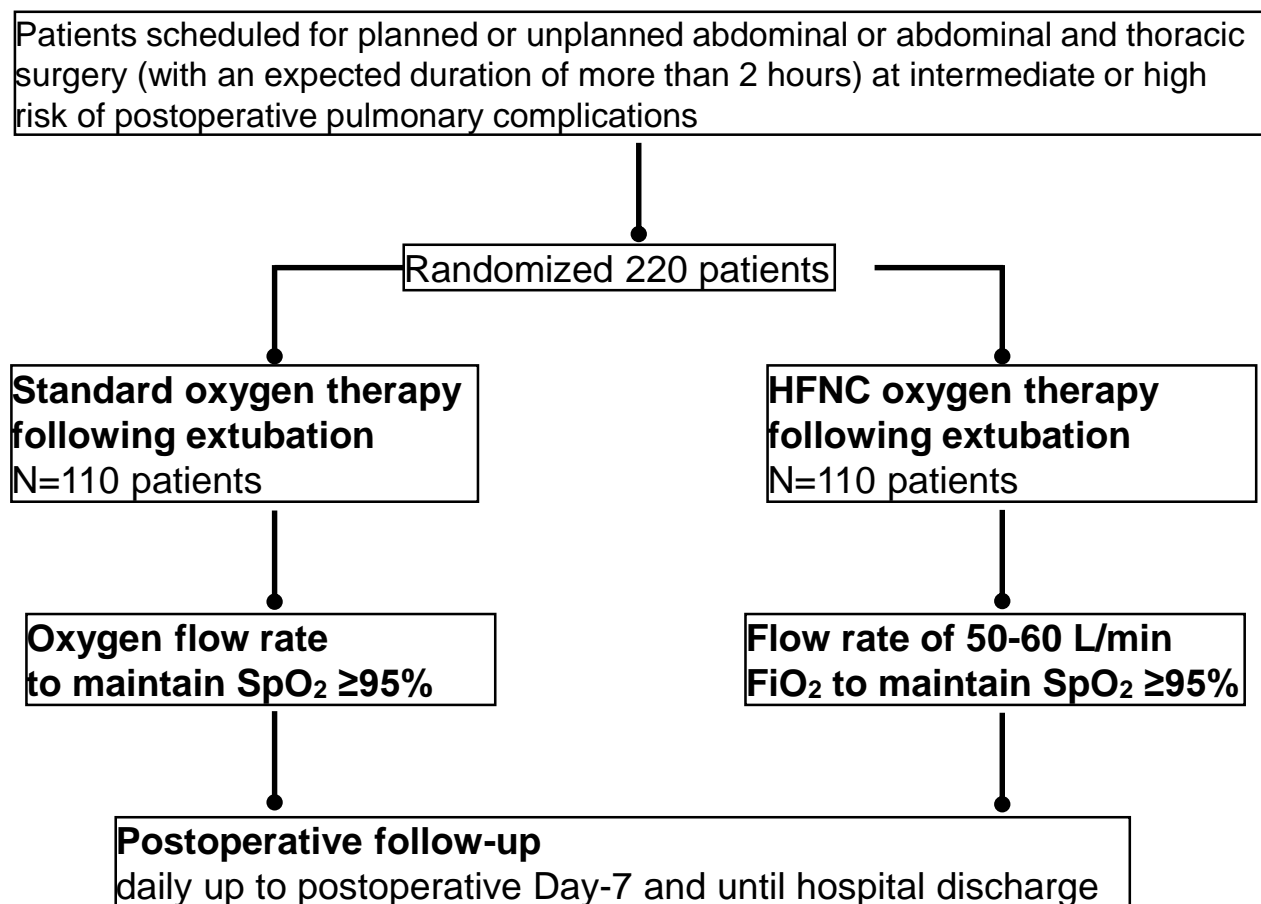
Direct extubation onto high-flow nasal cannulae post-cardiac surgery versus standard treatment in patients with a BMI ≥ 30 : a randomised controlled trial



STUDY PROTOCOL

Open Access

Optiflow® to Prevent **P**ost-**E**xtubation Hypoxemia after **R** **A**bdominal Surgery (**OPERA** Trial)

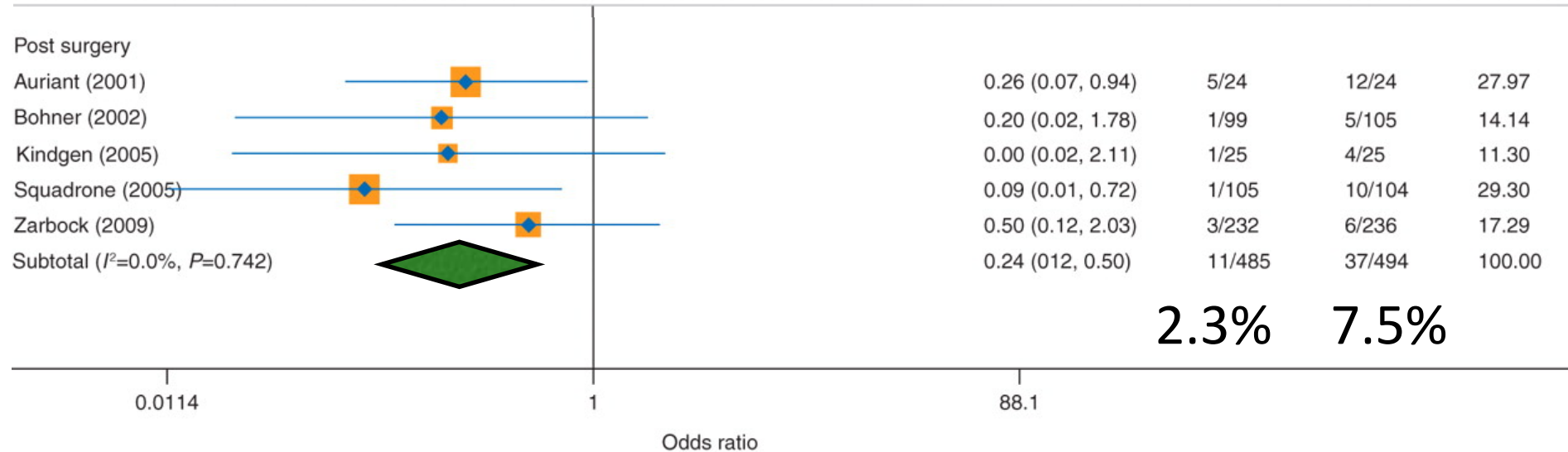


Non-invasive ventilation for weaning, avoiding reintubation after extubation and in the postoperative period: a meta-analysis

A. J. Glossop^{1*}, N. Shepherd², D. C. Bryden³ and G. H. Mills³

Reintubation

NIV after surgery



HFNC en post-opératoire

Original Investigation | CARING FOR THE CRITICALLY ILL PATIENT

High-Flow Nasal Oxygen vs Noninvasive Positive Airway Pressure in Hypoxemic Patients After Cardiothoracic Surgery A Randomized Clinical Trial

JAMA 2015;313(23):2331-9

François Stéphan, MD, PhD; Benoit Barrucand, MD; Pascal Petit, MD; Saida Rézaiguia-Delclaux, MD; Anne Médard, MD; Bertrand Delannoy, MD; Bernard Cosserant, MD; Guillaume Flicoteaux, MD; Audrey Imbert, MD; Catherine Pilorge, MD; Laurence Bérard, MD; for the BiPOP Study Group

Objectif: Déterminer que HFNC est non inférieur à la VNI pour la prévention des complications respiratoires dans les suites d'une chirurgie cardio-thoracique

High-Flow Nasal Oxygen vs Noninvasive Positive Airway Pressure in Hypoxemic Patients After Cardiothoracic Surgery A Randomized Clinical Trial

- Echec SBT ("à risque")
 - $SaO_2 < 90\%$ sous O_2 12 L pendant T-tube
 - Ou $PaO_2 < 75\text{mmHg}$ avec $FiO_2 > 50\%$ avec AI « faible »
- Succès SBT mais FdR IRA ("préventif")
 - body mass index > 30
 - FEVG $< 40\%$
 - Echec extubation antérieure
- Succès SBT mais échec extubation ("curatif")
 - P/F < 300
 - FR $> 25/\text{min}$ pendant au moins 2h
 - Mise en jeu muscles respiratoires accessoires ou ventilation paradoxale

Exclusion criteria were obstructive sleep apnea, tracheostomy, do-not-intubate status, delirium, nausea and vomiting, bradypnea, impaired consciousness, and hemodynamic instability.

Méthodes

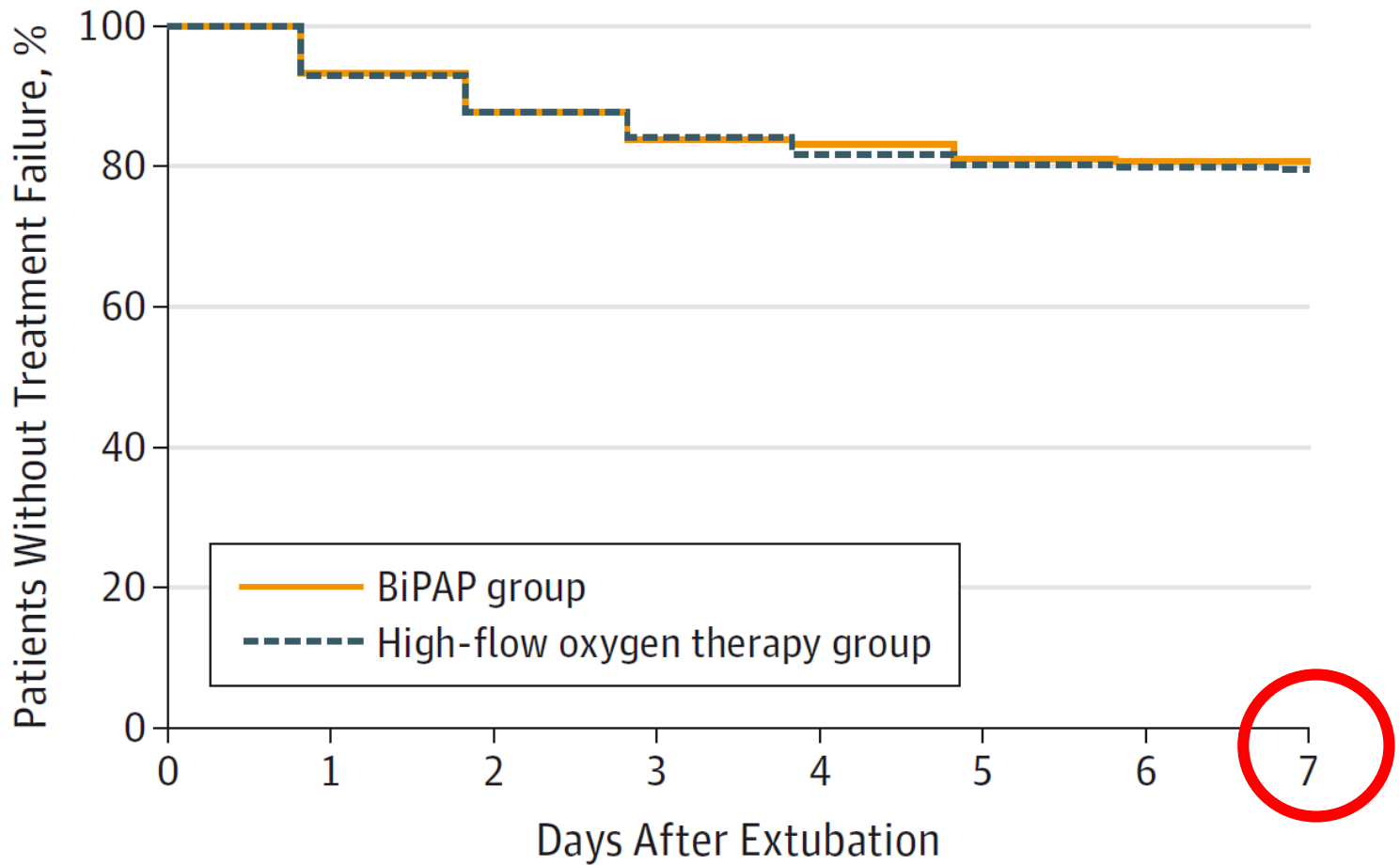
- Réglages initiaux
 - HFNC 50 lpm, FiO₂ 0,5 pour SaO₂ 92-98%
 - Bilevel positive airway pressure (BiPAP)
 - BiPap Vision (Respironics)
 - AI-PEEP (Drager Evita XL ou 4, Monnal T 75)
 - HME
 - AI 8 cmH₂O puis modifiée pour obtenir Vte 8 mL/kg et FR < 25/min
 - PEEP 4 cmH₂O; FiO₂ 0.5 ajustée pour SaO₂ 92-98%
 - 2 h puis environ 1 h toutes les 4h, ou davantage
 - Entre les sessions: O₂ lunettes, masque facial

Définitions

Stephan *et al.* JAMA 2015

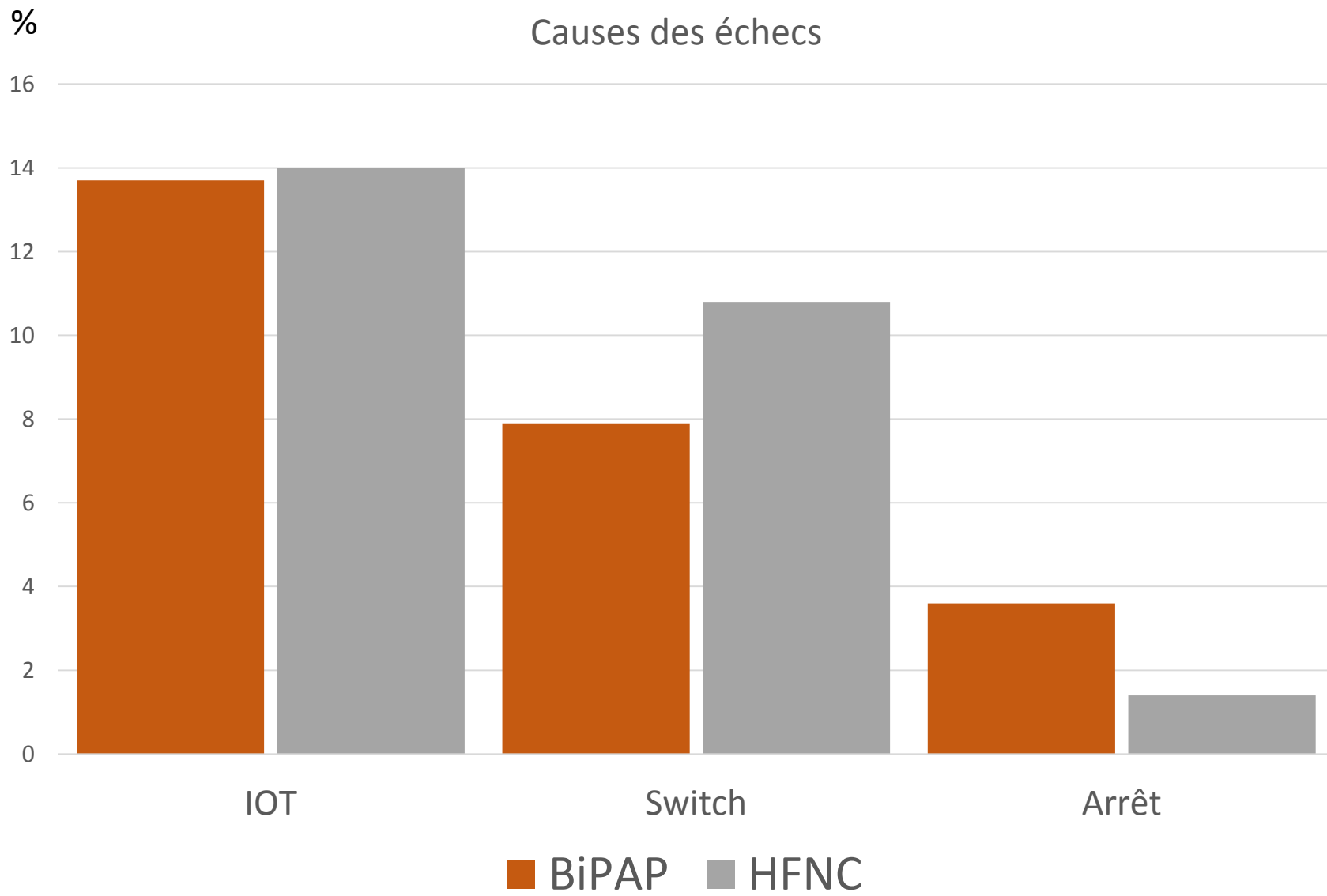
- Echec du traitement
 - Réintubation et VM
 - Switch pour un autre traitement
 - Arrêt prématuré
 - À la demande du patient
 - Pour motif médical
- Réintubation
 - Arrêt respiratoire
 - Pauses respiratoires avec alteration du niveau de vigilance
 - Instabilité hémodynamique
 - Sécrétions très abondantes (encombrement majeur)
 - Epuisement
 - SaO₂ < 88% sous FiO₂=1
 - Acidose respiratoire: pH < 7,30 et PaCO₂ > 50 mmHg

Characteristics	BiPAP (n = 416)	High-Flow Nasal Oxygen Therapy (n = 414)
Age, mean (95% CI), y	63.9 (62.6-65.2)	63.8 (62.5-65.2)
Men, No. (%)	278 (66.8)	273 (65.9)
Body mass index ^a		
Mean (95% CI)	28.2 (27.6-28.7)	28.3 (27.8-28.8)
>30, No. (%)	136 (32.7)	135 (32.6)
Smoking, No. (%) ^b		
Former	217 (52.2)	226 (54.6)
Current	69 (16.6)	83 (20.0)
SAPS II score at admission, mean (95% CI)	28.8 (27.7-30.0)	29.0 (27.8-30.1)
Surgical procedures, No. (%)		
Coronary artery bypass grafting	111 (26.7)	122 (29.5)
Valvular surgery	83 (20.0)	88 (21.3)
Combined cardiac surgery with coronary artery bypass grafting	27 (6.5)	26 (6.3)
Thoracic aorta	28 (6.7)	23 (5.6)
Pulmonary thromboendarterectomy	90 (21.6)	68 (16.4)
Lung resection	30 (7.2)	34 (8.2)
Heart, lung, and heart-lung transplantations	9 (2.2)	18 (4.3)
Others	38 (9.1)	35 (8.5)
Cardiopulmonary bypass, No. (%)	340 (81.7)	320 (77.2)
Duration of cardiopulmonary bypass, mean (95% CI), min	137 (129-146)	137 (128-146)
Time from surgery to randomization, median (IQR), d	1.0 (1.0-3.0)	1.0 (1.0-3.0)
Duration of mechanical ventilation at randomization, median (IQR), h	13.0 (6.0-27.5)	11.5 (5.0-25.4)



No. at risk

BiPAP	416	385	363	348	339	333	331	329
High-flow oxygen therapy	414	385	361	346	342	334	333	331



Pas d'amélioration du confort

Parameters	Mean (95% CI)							
	Baseline		1 Hour			6-12 Hours		
	BiPAP Group	HFNO Group	BiPAP Group	HFNO Group	P Value	BiPAP Group	HFNO Group	P Value
Dyspnea score, No./total (%) [95% CI]								
Improvement			266/404 (65.8) [61.0-70.7]	236/403 (58.6) [53.6-63.4]		229/379 (60.4) [55.3-65.4]	217/373 (58.2) [52.9-63.2]	
No improvement			120/404 (29.7) [25.3-34.4]	151/403 (37.5) [32.7-42.4]	.39	133/379 (35.1) [30.3-40.1]	139/373 (37.3) [32.4-43.4]	.99
Deterioration			18/404 (4.5) [2.7-7.0]	16/403 (4.0) [2.3-6.4]		17/379 (4.5) [2.6-6.1]	17/373 (4.6) [2.7-7.2]	
Comfort score, No. (%) [95% CI]								
Poor			51/397 (13.0) [9.7-16.5]	67/402 (16.7) [13.2-20.7]		67/376 (17.8) [14.1-22.1]	66/372 (17.7) [14.0-22.0]	
Acceptable			128/397 (32.2) [27.7-37.1]	101/402 (25.1) [21.0-29.7]	.32	110/376 (29.3) [24.7-34.1]	115/372 (31.0) [26.3-36.9]	.99
Good			218/397 (55.0) [49.9]	234/402 (58.2) [53.2-63.1]		199/376 (53.0) [47.7-58.1]	101/372 (51.0) [46.1-56.5]	

Events	Group		P Value
	BiPAP (n = 416)	HFNO (n = 414)	
Nosocomial pneumonia, No. (%) [95% CI]	90 (21.6) [17.8-25.9]	83 (20.0) [16.4-24.3]	.57
Pneumothorax, No. (%) [95% CI]	7 (1.7) [0.7-3.6]	8 (1.9) [0.9-3.9]	.86
Acute colonic pseudo-obstruction, No. (%) [95% CI]	8 (1.9) [0.9-3.9]	9 (2.2) [1.0-4.2]	.86
No. of days with respiratory support, median (IQR)	2 (1-3)	2 (1-3)	.59
Stay length, median (IQR), d			
ICU	6 (4-10)	6 (4-10)	.77
Hospital	14 (9-20)	13 (9-22)	.59

- Echec SBT (“à risque”)
 - SaO₂ < 90% sous O₂ 12 L pendant T-tube
 - Ou PaO₂ < 75mmHg avec FiO₂ > 50% avec AI « faible »
- Succès SBT mais FdR IRA (“préventif”)
 - body mass index > 30
 - FEVG < 40%
 - Echec extubation antérieure
- Succès SBT mais échec extubation (“curatif”)
 - P/F < 300
 - FR > 25/min pendant au moins 2h
 - Mise en jeu muscles respiratoires accessoires ou ventilation paradoxale

Table. Treatment Failure Rate According to the Eligibility Criteria

Eligibility Criteria	Treatment Failure, No./Total (%)		P Value ^a
	Bilevel Positive Airway Pressure	High-Flow Nasal Oxygen Therapy	
Failure of spontaneous breathing trial	7/25 (28.0)	11/27 (40.7)	.33
Successful spontaneous breathing trial in high-risk patients	19/151 (12.6)	8/139 (5.7)	.04
Failed extubation	65/240 (27.8)	68/248 (27.4)	.93

High-Flow Nasal Cannulae or Noninvasive Ventilation for Management of Postoperative Respiratory Failure

Lorenzo Del Sorbo, MD; Niall D. Ferguson, MD, MSc

How should clinicians apply the results of the trial by Stephan et al. in clinical practice? The answer is, as usual, it depends. In centers with considerable expertise with NIV and for patients who are hypercapnic, a reasonable approach would be to favor NIV in this setting. Alternatively, based on these data and others, for many postoperative patients HFNC appear to be a viable alternative that is better tolerated and may lead to non inferior clinical outcomes.

HFNC en pré-intubation

Pré- et per-intubation

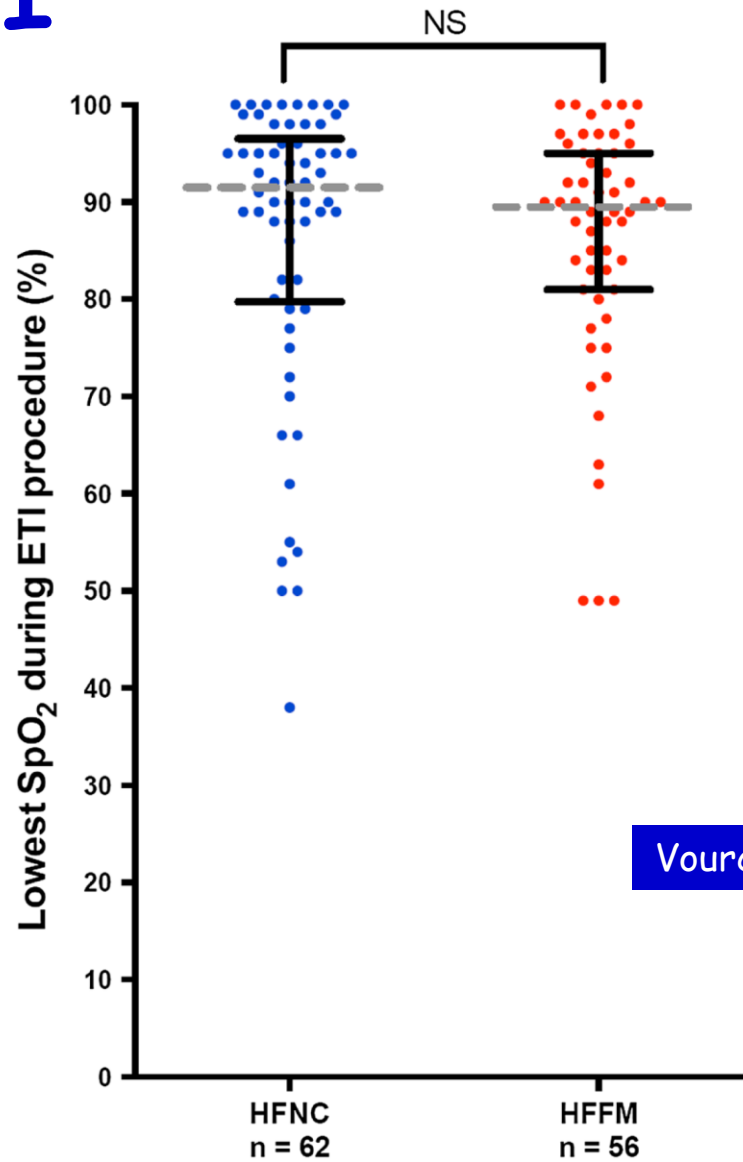
PREOXYFLOW

- Multicentrique
- IOT par juniors/seniors
- ARF
 - FR > 30 et FiO₂ ≥ 50 % pour SpO₂ ≥ 90 %
 - et PaO₂/FiO₂ ratio < 300 mmHg dans les 4h précédant inclusion
- HFNC 60 lpm FiO₂=1 pendant 4 minutes avant induction et pendant l'intubation vs. masque 15 lpm O₂ pendant 4 minutes avant induction
- Objectif I: SpO₂ la plus basse entre l'induction et la connexion au ventilateur

	High-flow nasal cannulae (<i>n</i> = 62)	High-flow face mask (<i>n</i> = 57)
Sex ratio M/F	39/23	39/18
Age, mean (SD), years	64.9 (14)	59.3 (14.5)
BMI, mean (SD) ^a	27.6 (5.8)	27.6 (7.3)
Medical patients, <i>n</i> (%)	54 (87.1)	50 (87.7)
SAPS II score, mean (SD) ^b	54.5 (20.2)	51.3 (16.5)
Comorbidities		
Chronic heart failure (NYHA III or IV), <i>n</i> (%)	4 (6.5)	3 (5.3)
Chronic respiratory failure, <i>n</i> (%)	5 (8)	5 (8.8)
Obstructive sleep apnoea syndrome, <i>n</i> (%)	1 (1.6)	6 (10.7)
COPD patients, <i>n</i> (%) ^c	7 (11.3)	6 (10.7)
Past upper airway tract cancer, <i>n</i> (%)	1 (1.6)	1 (1.8)
Diabetes requiring insulin therapy, <i>n</i> (%)	2 (3.2)	4 (7)
Vasopressor support at inclusion, <i>n</i> (%)	18 (29)	11 (19.3)
Glasgow Coma Score, mean (SD)	13.5 (2.8)	13.4 (2)
McCabe scale 2 or 3, <i>n</i> (%) ^d	30 (48.4)	23 (40.4)
Functional status KNAUS (class C or D), <i>n</i> (%) ^c	19 (30.6)	13 (22.8)
Respiratory failure aetiology, <i>n</i> (%)		
Pneumonia	25 (40.3)	29 (50.9)
Extra-respiratory ARDS	19 (30.6)	11 (19.3)
Cardiogenic pulmonary oedema	3 (4.8)	5 (8.7)
COPD exacerbation	4 (6.4)	0
Thoracic trauma	1 (1.6)	5 (8.7)
Other	10 (16.1)	7 (12.2)
Advanced oxygenation support in the last hour before inclusion, <i>n</i> (%)		
NIV	11 (17.8)	8 (14)
HFNC	10 (16.1)	3 (5.3)
Arterial blood gas oxygenation		
PaO ₂ /FiO ₂ , mean (SD), mm Hg	120.2 (55.7)	115.7 (63)
PaCO ₂ , mean (SD), mm Hg	39.8 (14.7)	37.4 (11)
SpO ₂ , mean (SD), %	94.6 (4.7)	93.4 (6)

	High-flow nasal cannulae (<i>n</i> = 62)	High-flow face mask (<i>n</i> = 57)
Airway description		
At least 2 difficult mask ventilation criteria ^a , <i>n</i> (%)	43 (69.4)	33 (57.9)
Medical history of difficult intubation, <i>n</i> (%)	0	2 (3.5)
Mouth opening less than 3 cm, <i>n/n</i> tot ^b (%)	4/52 (7.7)	8/48 (16.7)
Limitation of cervical mobility $\leq 35^\circ$, <i>n/n</i> total ^b (%)	2/44 (4.6)	8/42 (19)
Thyromental distance < 65 mm, <i>n/n</i> tot ^b (%)	4/51 (7.8)	10/40 (25)
Mallampati III or IV, <i>n/n</i> tot ^b (%)	10/39 (25.6)	12/32 (37.5)
Macocho score, <i>n</i> (%) ^c		
<3	25 (78.1)	11 (55)
≥ 3	7 (21.88)	9 (45)
First operator ^d		
Senior <i>n</i> (%)	11 (17.7)	16 (28)
Junior <i>n</i> (%)	51 (82.2)	41 (71.9)
Emergency intubation ^e		
Real emergency, <i>n</i> (%)	12 (19.3)	29 (50.8)
Relative emergency, <i>n</i> (%)	48 (77.4)	25 (43.8)
Deferred emergency, <i>n</i> (%)	2 (3.2)	3 (5.2)

Objectif I



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Median (IQR) 91.5 (80-96)

Mean (SD) 86.2 (15.2)

Median (IQR) 89.5 (81-95)

Mean (SD) 85.9 (12.9)

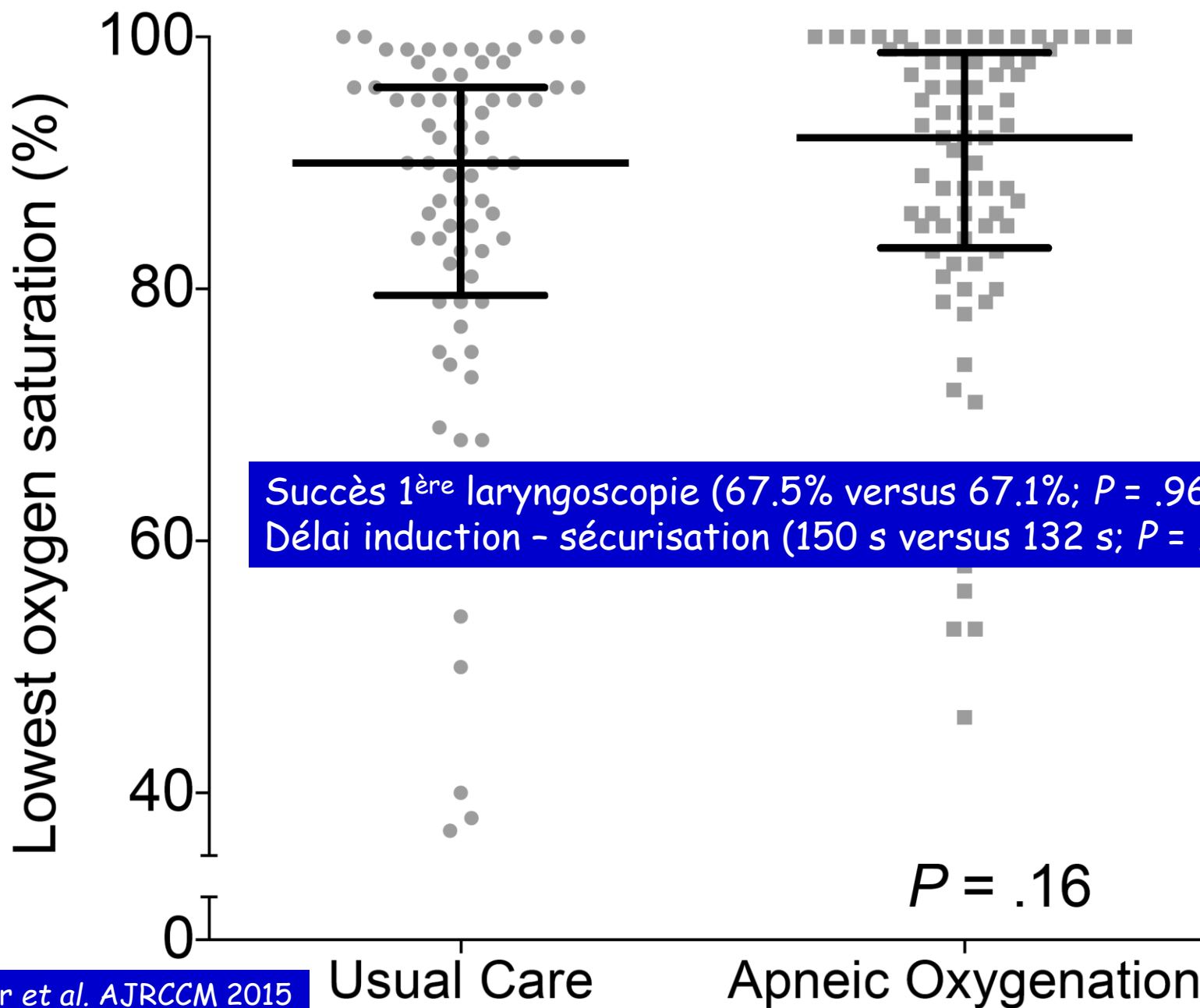
	High-flow (<i>n</i> = 62)	nasal cannulae	High-flow (<i>n</i> = 57)	face mask	<i>p</i> ^a
Primary outcome					
Lowest SpO ₂ during ETI procedure, median (IQR)	91.5 (80–96)		89.5 (81–95)		0.44
Secondary outcomes					
Preoxygenation					
Duration of preoxygenation, <i>n</i> (%)					
<4 min	4 (6.5)		3 (5.3)		
4 min or more	58 (93.5)		54 (94.7)		0.83
SpO ₂ at the beginning of preoxygenation, mean (SD)	95.4 (3.9)		94.1 (6.0)		0.15
SpO ₂ at the end of preoxygenation, mean (SD)	97.1 (3.8)		96.3 (4.4)		0.98
Failure to increase saturation to 90 % during preoxygenation, <i>n</i> (%)	4 (6.5)		2 (3.5)		0.49
Respiratory outcome					
PaO ₂ 1 h after ETI, median (IQR), mm Hg	98.2 (72–139.5)		89.5 (69.8–144)		0.94
Duration of mechanical ventilation, median (IQR), days	6 (4–14)		10 (5–17)		0.02
Ventilator-free days (day 28), median (IQR), days ^d	14 (0–22)		5 (0–16)		0.09
Ventilator-associated pneumonia, <i>n</i> (%)	6 (9.7)		8 (14)		0.62
Morbidity in ICU					
ICU length of stay, median (IQR), days	10 (6–16)		13 (7–24)		0.12
SOFA score, mean (SD)					
Day 1	8.8 (4.1)		9.2 (3.6)		0.57
Day 2 (<i>n</i> = 112)	8.5 (4)		9.3 (4.1)		0.65
Day 3 (<i>n</i> = 103)	7.1 (3.5)		8 (4.7)		0.63
Day 4 (<i>n</i> = 99)	6.1 (3.6)		7.3 (4.5)		0.23
Day 5 (<i>n</i> = 95)	5.6 (3.8)		6.8 (4.6)		0.21
Death in ICU, <i>n</i> (%)	21 (33.9)		23 (40.4)		0.46
Mortality at 28 days, <i>n</i> (%)	22 (35.4)		24 (42.1)		0.48

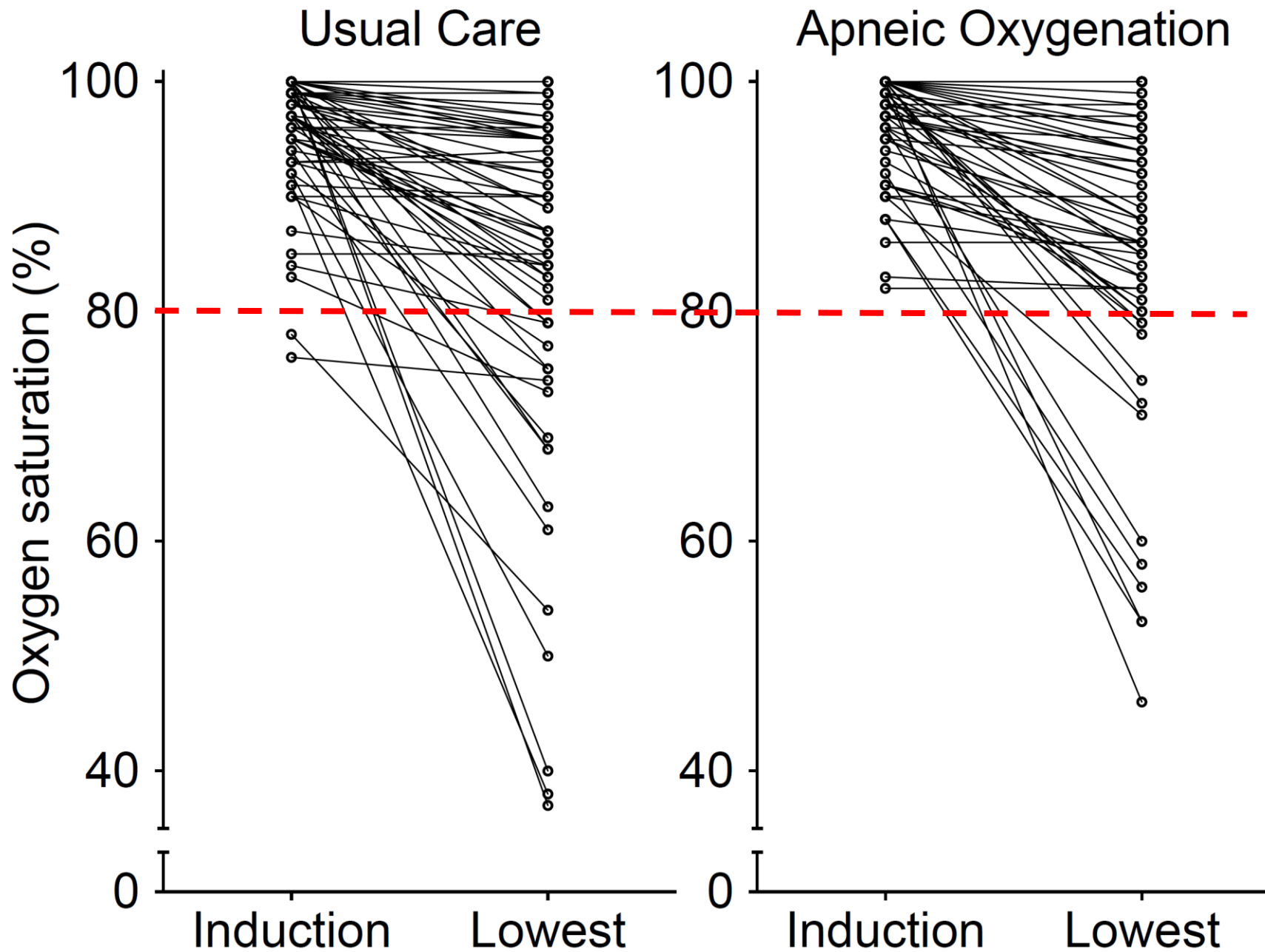
Per-intubation (apnée)

FELLOW

- Monocentrique
- IOT par fellows
- Compare efficacité HFNC 15 lpm $FiO_2=1$ vs. pas d' O_2 pendant premier essai de laryngoscopie
- Objectif I: SpO_2 la plus basse entre l'induction et la 2^{ème} minute après l'intubation

Semler et al. AJRCCM 2015





	Usual Care	Apneic Oxygenation	
Oxygenation Outcomes	(n = 73)	(n = 77)	P Value
Lowest oxygen saturation, median [IQR], %	90 [80-96]	92 [84-99]	.16
Lowest oxygen saturation < 90%, No. (%)	34 (47.2%)	34 (44.7%)	.87
Lowest oxygen saturation < 80%*, No. (%)	18 (25.0%)	12 (15.8%)	.22
Decrease in oxygen saturation, median [IQR], %	4.5 [1-14]	4.0 [0-12]	.60
Decrease in oxygen saturation > 3%, No. (%)	40 (55.6%)	41 (53.9%)	.87
Clinical Outcomes			
Duration of mechanical ventilation, median [IQR], days	3 [2-7]	3 [1-10]	.73
Intensive care unit length of stay, median [IQR], days	7 [3-10]	4 [2-9]	.24
Died within one hour of intubation, No. (%)	1 (2.8%)	0 (0.0%)	>.99
Died before hospital discharge, No. (%)	36 (49.3%)	27 (35.1%)	.10

Recommendations of the task force regarding the use of HFNC

Indication	Acute respiratory failure	In obese patients	Prior intubation	Following extubation	During bronchoscopy	For the delivery of aerosol therapy
Level of evidence	High To be confirmed	Low Additional trials are needed	Low to Moderate Additional trials are needed	Moderate Additional trials are needed	Moderate Additional trials are needed	Low to Moderate Additional trials are needed

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