

Sevrage respiratoire DU Kiné

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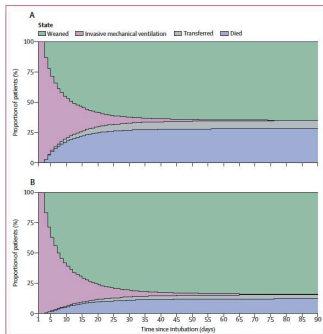


LES GRANDS PRINCIPES DU SEVRAGE

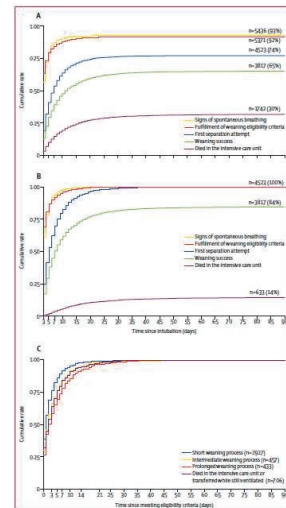
- Simple à Difficile
- Sédation
- Fonction Cardiaque
- Force musculaire

Weaning from mechanical ventilation in intensive care units across 50 countries (WEAN SAFE): a multicentre, prospective, observational cohort study

Tai Pham, Leo Heunks, Giacomo Bellani, Fabiana Madotto, Irene Aragao, Gaëtan Beduneau, Ewan C Gallagher, Giacomo Grasselli, Jon Henrik Laake, Jordi Mancebo, Oscar Penuelas, Lise Piquilloud, Antonio Pesenti, Hannah Wunsch, Frank van Haren, Laurent Brochard*, John G Laffey*, for the WEAN SAFE Investigators†



Lancet Respir Med 2023; 11: 465-76



Sevrage 3 situations

- 1) **Sevrage simple** (premier essai)
65%
- 2) **Sevrage prolongé** (plus d'un essai)
25%
- 3) **Sevrage (très) difficile**
10%

Sevrage 3 situations

	Funk	Penuelas	Sellares	Tonnellier
Simple	59 %	55 %	44 %	30 %
Prolongé	26 %	39 %	37 %	40 %
Difficile	14 %	6 %	18 %	30 %

Mécanismes du Sevrage non difficile

- Sevrage possible d'emblée:
 - ☒ screening quotidien
- Sevrage possible mais non identifié
 - ☒ Sur assistance
 - ☒ Trop de sédation
 - ☒ Pas de screening

Mécanismes du Sevrage difficile

- Pathologie pulmonaire/cardiaque sous jacente sévère
- Surcharge volémique
- Faiblesse musculaire respiratoire
- Métabolique/Nutrition/Anémie

Intensive Care Med 2007; 12: 1022-1026
DOI: 10.1185/0961321060132006
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TASK FORCE

Weaning from mechanical ventilation

J.-M. Bolea¹, J. Biot², A. Comros³, M. Herridge⁴, B. Marsh¹, C. Melot¹, R. Peari⁵, H. Silverman⁶, M. Stanchina⁷, A. Vieillard-Baron⁸, T. Welte⁹

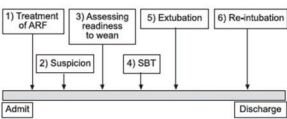


TABLE 3 Classification of patients according to the weaning process

Group/category	Definition
Simple weaning	Patients who proceed from initiation of weaning to successful extubation on the first attempt without difficulty
Difficult weaning	Patients who fail initial weaning and require up to three SBT or as long as 7 days from the first SBT to achieve successful weaning
Prolonged weaning	Patients who fail at least three weaning attempts or require >7 days of weaning after the first SBT

SBT: spontaneous breathing trial.



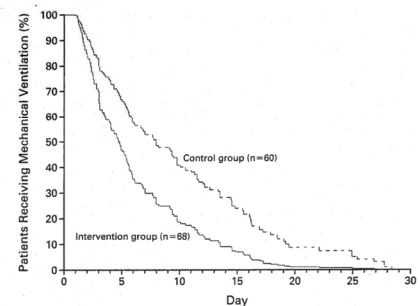
Sevrage du ventilateur

Sevrage de la sonde

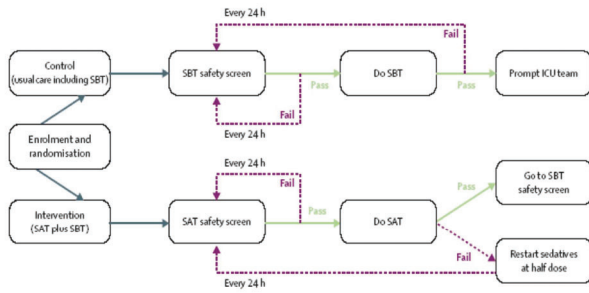


Protocole de sédation

Kress, JP, et al. N Engl J Med 2000; 342:1471-1477



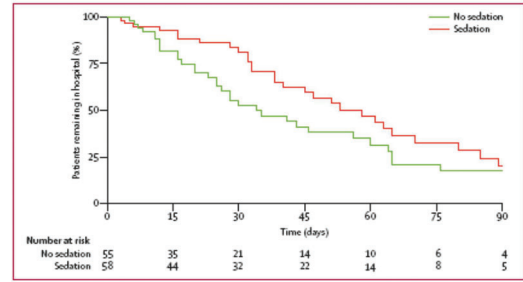
Efficacy and safety of a paired sedation and ventilator weaning protocol for mechanically ventilated patients in intensive care (Awakening and Breathing Controlled trial): a randomised controlled trial



Lancet 2008; 371: 126-34

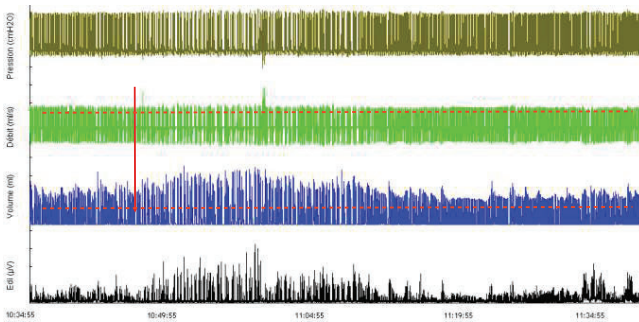
A protocol of no sedation for critically ill patients receiving mechanical ventilation: a randomised trial

Thomas Strøm, Torben Martinussen, Palle Toft



Lancet 2010

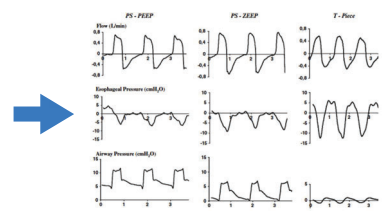
Test à l'anexate démontre l'inhibition de la sédation



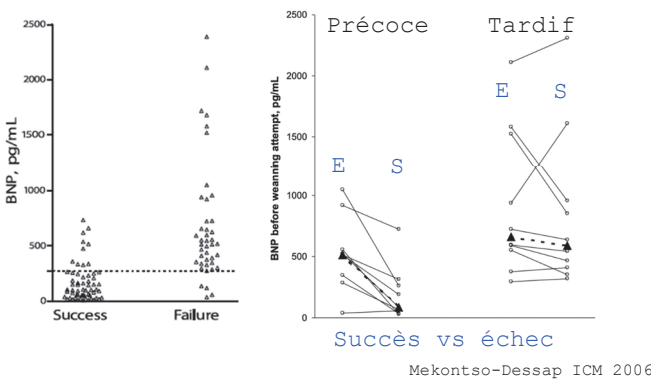
WEAVING FAILURE OF CARDIAC ORIGIN: RECENT ADVANCES

ORIGINAL

Physiological comparison of three spontaneous breathing trials in difficult-to-wean patients



Fonction cardiaque



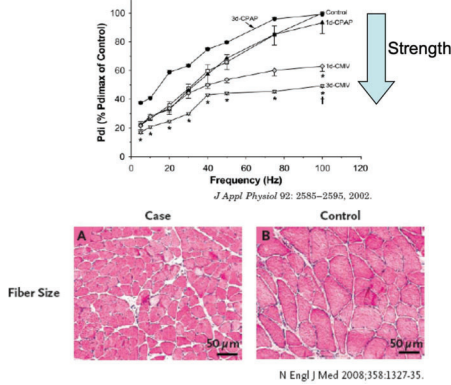
Mekontso-Dessap ICM 2006

Balance hydrique

Fluid balance and weaning outcomes

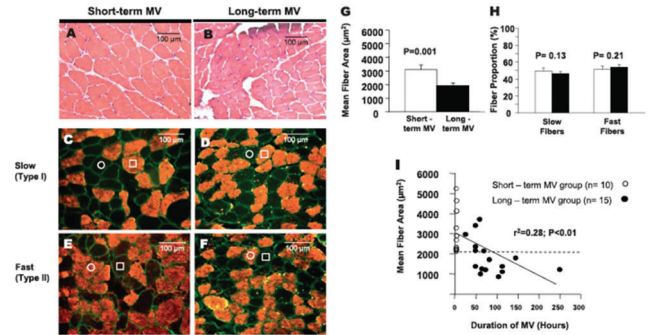
	Weaning success (n=39)	Weaning failure (n=48)	All patients (n=87)	p
Duration of ventilation (days)	2 (1 to 8)	3 (1 to 14)	3 (1 to 14)	0.03
APACHE II (day of BT)	17 (3 to 30)	15 (3 to 25)	16 (3 to 30)	0.2
Fluid balance 24 h (ml)	-625 (-4,380 to +3,274)	+242 (-3,923 to +4,272)	-91 (-4,380 to +4,272)	0.01
Cumulative fluid balance (ml)	-633 (-8,232 to +9,534)	+920 (-1,176 to +2,048)	-65 (-1,176 to +2,048)	0.06
Diuretics on day of trial	62%	51%	56%	0.2
Prealbumin (mg/dl)	14 (2 to 35)	12 (2 to 30)	13 (2 to 35)	0.8
fV ₁ (breaths min ⁻¹ l ⁻¹)	50 (13 to 260)	80 (17 to 300)	67 (13 to 300)	0.005
PaO ₂ /FIO ₂	240 (88 to 477)	244 (100 to 503)	240 (88 to 503)	0.9
Compliance (ml/cmH ₂ O)	44 (22 to 76)	39 (13 to 80)	43 (13 to 80)	0.2
Left ventricular dysfunction (%)	69	68	69	0.6

Faiblesse musculaire: Dysfonction diaphragmatique



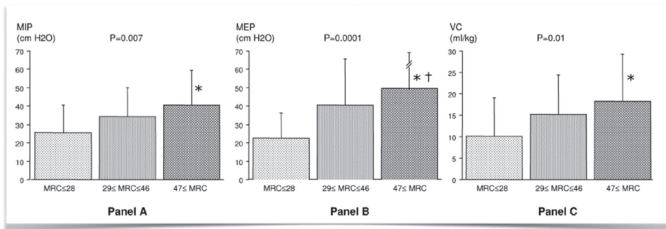
Rapidly Progressive Diaphragmatic Weakness and Injury during Mechanical Ventilation in Humans

Samir Jaber^{1,2,6}, Basil J. Petrof³, Boris Jung^{1,2}, Gérald Chanques^{1,2}, Jean-Philippe Berthet⁴, Christophe Rabuel⁵, Hassan Bouayrabine⁶, Patricia Courouble^{1,2}, Christelle Koechlin-Ramonato⁷, Mustapha Sebbane^{1,2}, Thomas Similowski⁸, Valérie Scheuermann⁹, Alexandre Mebazaa³, Xavier Capdevila^{1,2}, Dominique Mornet^{2,10}, Jacques Mercier^{2,10}, Alain Lacampagne², Alexandre Philipps², and Stefan Matecki^{2,10}



Respiratory weakness is associated with limb weakness and delayed weaning in critical illness*

Bernard De Jonghe, MD; Sylvie Bastuji-Garin, MD, PhD; Marie-Christine Durand, MD; Isabelle Malassin, MD; Pablo Rodrigues, MD; Charles Cerf, MD; Hervé Outin, MD; Tarek Sharshar, MD, PhD; for Groupe de Réflexion et d'Etude des Neuromyopathies En Réanimation



Crit Care Med 2007; 35:2007-2015

Table 6. Analysis of risk factors for low maximal inspiratory pressure (MIP) or maximal expiratory pressure (MEP)

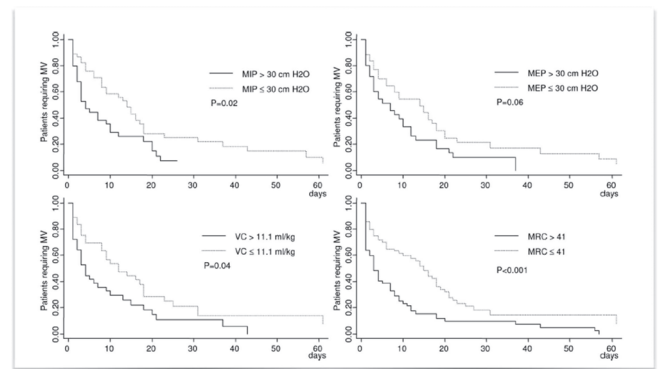
Univariate Analysis	MIP or MEP ≤ 30 cm H ₂ O (n = 53)	MIP and MEP > 30 cm H ₂ O (n = 26)	p Value
At ICU admission			
Age in years, median (IQR)	67 (53-78)	70 (54-76)	.9
Female sex, n (%)	22 (41.5)	7 (26.9)	.3
Admission SAPS II, median (IQR)	46 (36-58)	44 (38-50)	.4
COPD, n (%)	20 (37.7)	13 (50)	.3
Before awakening			
Days of MV, median (IQR)	10 (8-14)	10 (8-13)	.9
Days with ≥ 2 failed organs, median (IQR)	6 (4-10)	5 (0-9)	.2
Use of corticosteroids, n (%)	33 (62.3)	16 (61.5)	.9
Use of neuromuscular blockers, n (%) ^a	13 (24.5)	11 (42.3)	.1
Average daily morning BGL (mmol/L), median (IQR)	8.1 (7.0-9.3)	7.5 (6.8-8.6)	.1
Septic shock, n (%)	31 (58.5)	8 (30.8)	.02
Multivariate Logistic Regression Analysis of Low MIP or MEP ≤ 30 cm H₂O			
	OR	95% CI	p Value
Septic shock	3.17	1.17-8.58	.02

Crit Care Med 2007; 35:2007-2015

Table 5. Independent determinants of the risk of successful extubation delayed for ≥ 7 days after awakening

	Model with MIP			Model with MEP			Model with VC			Model with MRC					
	OR	95% CI	p Value	OR	95% CI	p Value	OR	95% CI	p Value	OR	95% CI	p Value			
MIP ≤ 30 cm H ₂ O	8.02	2.12-30.36	.002	MEP ≤ 30 cm H ₂ O	4.15	1.16-14.82	.03	VC ≤ 11.1 ml/kg	2.75	0.82-9.18	.1	MRC ≤ 41	3.03	1.23-7.43	.02
COPD	4.43	1.20-16.41	.03	COPD	4.56	1.24-16.75	.02	COPD	4.43	1.3-14.79	.02	COPD	2.74	1.10-6.85	.03
Cardiac insufficiency	4.56	1.25-19.71	.02	Cardiac insufficiency	3.79	1.07-13.39	.04	Cardiac insufficiency	3.24	0.92-11.38	.07	Cardiac insufficiency	2.14	0.82-5.61	.1

Crit Care Med 2007; 35:2007-2015

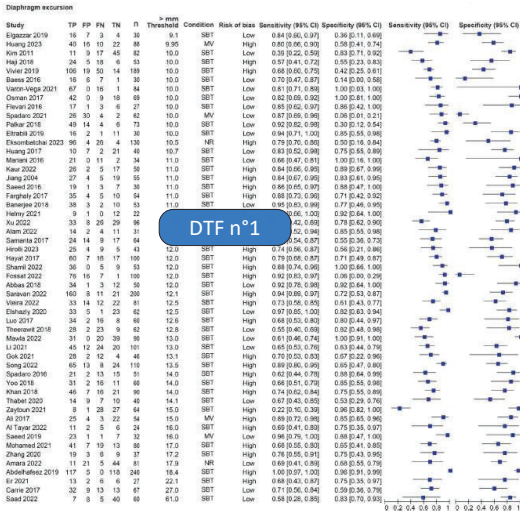
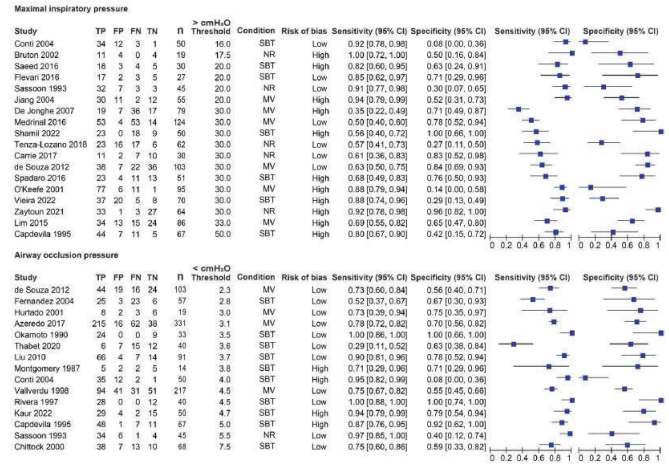
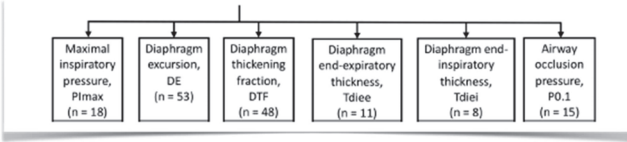


Crit Care Med 2007; 35:2007-2015

RESEARCH Open Access

Accuracy of respiratory muscle assessments to predict weaning outcomes: a systematic review and comparative meta-analysis

Diego Poddighe^{1,2†}, Marine Van Hollebeke^{1,2†}, Yasir Qaiser Choudhary¹, Débora Ribeiro Campos¹, Michele R. Schaeffer¹, Jan Y. Verbaek^{4,5}, Greet Hermans^{6,7}, Rik Gosselec^{1,2,7} and Daniel Langer^{1,2*}



Faiblesse musculaire

Review

J Appl Physiol 107: 962-970, 2009.
 First published April 30, 2009; doi:10.1152/jap.00165.2009.

HIGHLIGHTED TOPIC | The Respiratory Muscles in Chronic Obstructive Pulmonary Disease

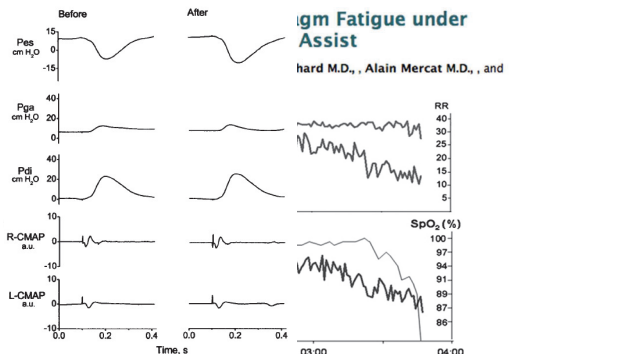
Role of the respiratory muscles in acute respiratory failure of COPD: lessons from weaning failure

Martin J. Tobin,¹ Franco Laghi,¹ and Laurent Brochard²

Is Weaning Failure Caused by Low-Frequency Fatigue of the Diaphragm? Voir de la fatigue ?

Franco Laghi, Steven E. Cattapan, Amal Iubran, Salim Parthasarathy, Paul Warshawsky, Yoon-Sub A. Chol, and Martin J. Tobin

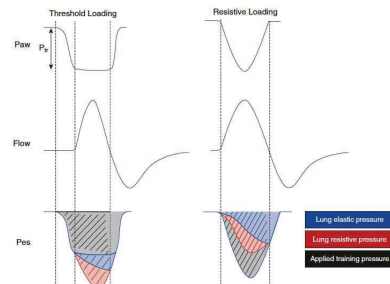
Am J Respir Crit Care Med Vol 167, pp 120-127, 2003



ORIGINAL RESEARCH

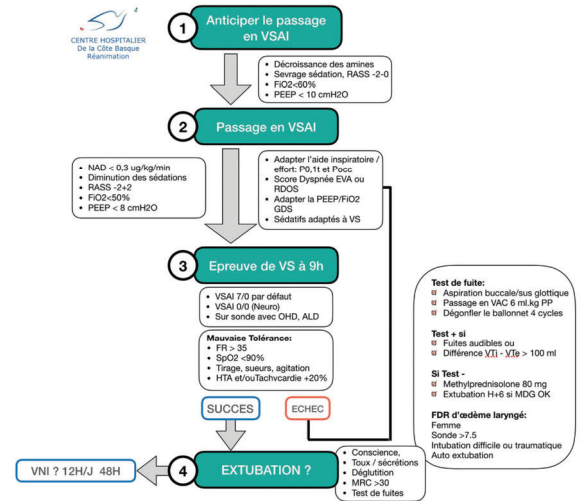
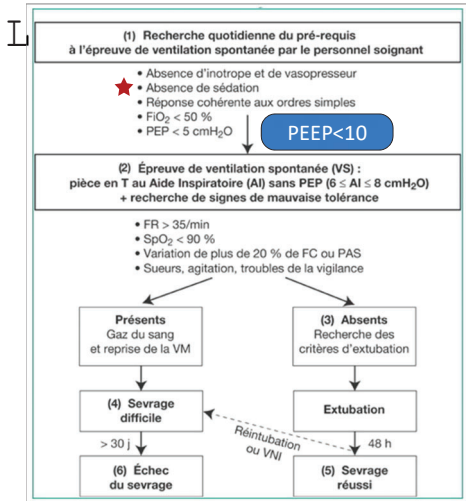
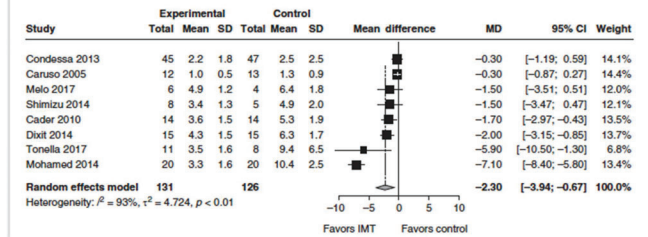
Inspiratory Muscle Rehabilitation in Critically Ill Adults: A Systematic Review and Meta-Analysis

Stefannie Vorona,¹ Umberto Sbatini,¹ Sulaiman Al-Maqbal,¹ Michele Berton,¹ Martin Dres^{2,3}, Bernie Bisset^{4,5}, Frank Van Haren^{6,7}, A. Daniel Martin⁸, Cristian Urrea¹, Debbie Brace¹, Matteo Parotto^{9,10,11}, Margaret S. Herridge^{12,13}, Neill K. J. Adhikari^{6,13,14}, Eddy Fan^{1,9,12,15}, Luana T. Melo¹⁶, W. Darlene Reid¹⁶, Laurent J. Brochard^{9,12}, Niall D. Ferguson^{1,9,12,14,15}, and Ewan C. Goligher^{9,15}



Outcome	Impact Effect (95% CI)	No. of Participants (RCTs)	Quality of the Evidence (GRADE)
Change in maximal inspiratory pressure from baseline after IMT	Mean difference in change 6 (5 to 8) cm H ₂ O higher in IMT group than in control group Pooled ratio of means for change in MIP relative to baseline MIP, 1.21 (1.16 to 1.25)	647 (15 RCTs)	⊙○○○ Very low ^{1,4}
Change in maximal inspiratory pressure from baseline after IMT (sensitivity analysis excluding studies at high risk of bias)	Mean difference 9 (7 to 12) cm H ₂ O higher in IMT group than in control group	175 (8 RCTs)	⊙⊙⊙⊙ High
Maximal inspiratory pressure after IMT	Mean difference 7 (6 to 8) cm H ₂ O higher in IMT group than in control group	575 (15 RCTs)	⊙○○○ Low ¹
Change in maximal expiratory pressure from baseline after IMT	Mean difference in change 9 (5 to 14) cm H ₂ O higher in IMT group than in control group Pooled ratio of means for change in MEP relative to baseline MEP, 1.39 (1.27 to 1.54)	153 (8 RCTs)	⊙⊙⊙○ Moderate ⁴
Change in maximal expiratory pressure from baseline after IMT (sensitivity analysis excluding studies at high risk of bias)	Mean difference in change 9 (5 to 14) cm H ₂ O higher in IMT group than in control group	106 (8 RCTs)	⊙⊙⊙⊙ High
Duration of ventilation	Pooled duration of ventilation was 4.1 (0.8 to 7.4) d shorter in IMT group than in control group	325 (8 RCTs)	⊙○○○ Very low ^{1,4,5}
Duration of ventilation (sensitivity analysis excluding studies at high risk of bias)	Pooled duration of ventilation was 4.6 (1.1 to 10.1) d shorter in IMT group than in control group	220 (8 RCTs)	⊙○○○ Low ^{1,5}
Duration of weaning from mechanical ventilation	Pooled duration of weaning from mechanical ventilation was 2.3 (0.7 to 3.9) d shorter in IMT group than in control group	257 (8 RCTs)	⊙○○○ Very low ^{1,5}
Duration of weaning (sensitivity analysis excluding studies at high risk of bias)	Pooled duration of weaning from mechanical ventilation was 3.2 (0.6 to 5.8) d shorter in IMT group than in control group	209 (8 RCTs)	⊙○○○ Low ^{1,5}
ICU length of stay	Length of stay in ICU was 3.1 (-1.0 to 7.1) d shorter in IMT group than in control group	28 (8 RCTs)	⊙○○○ Very low ^{1,4,5}
Mortality in ICU	Pooled relative risk of death in ICU was 0.67 (0.20 to 2.20) in IMT group compared with control group	197 (8 RCTs)	⊙○○○ Low ^{1,5}

The impact of inspiratory muscle training (IMT) on the duration of weaning from mechanical ventilation

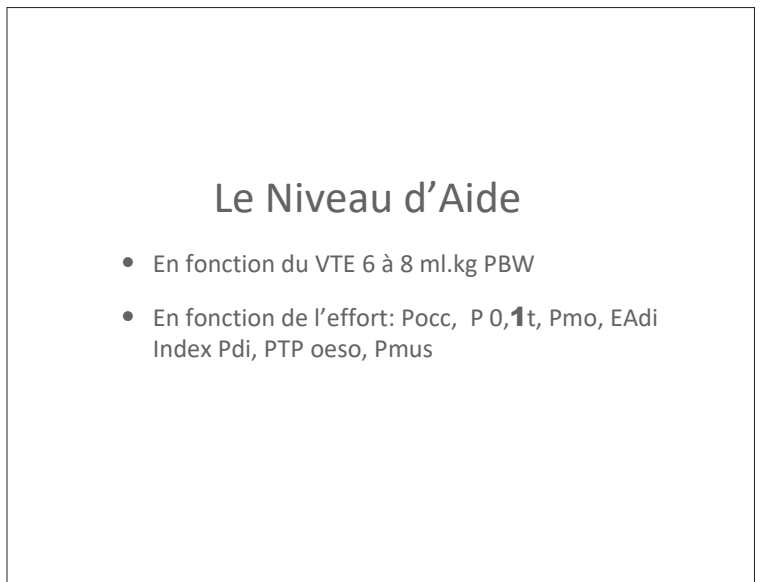
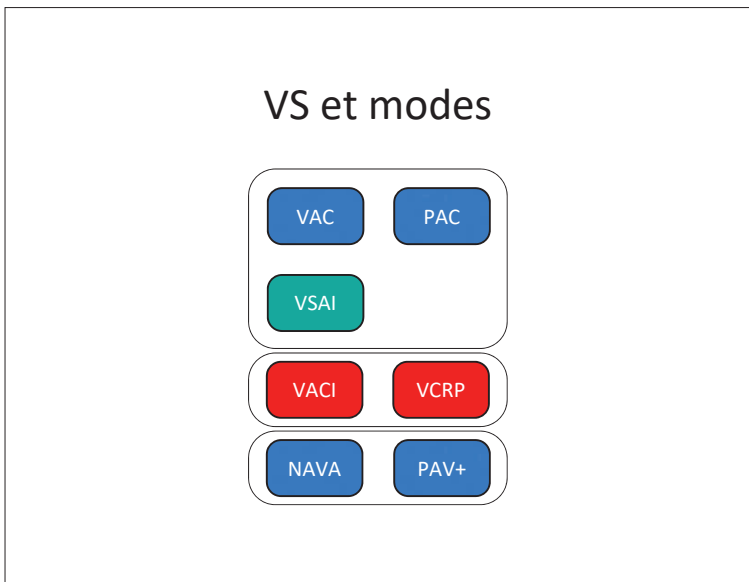
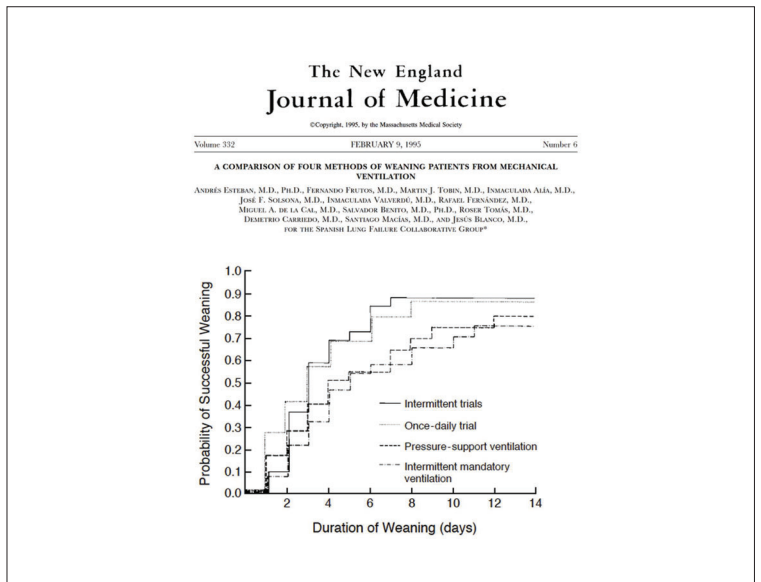
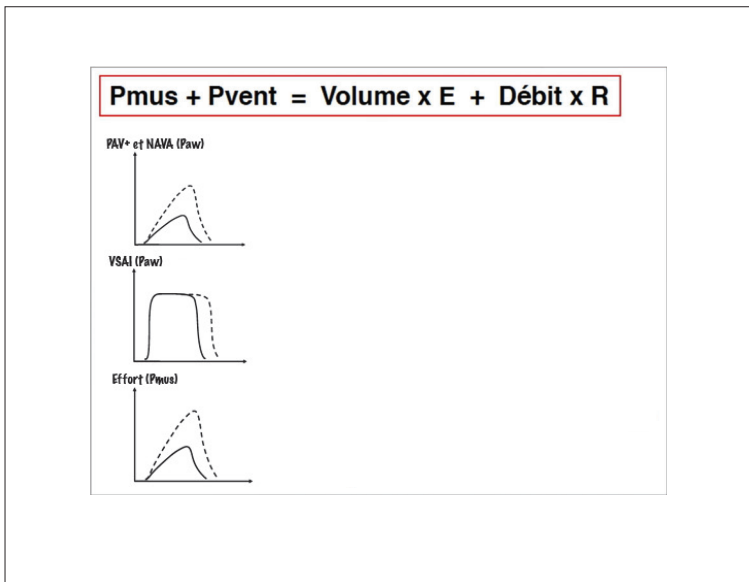
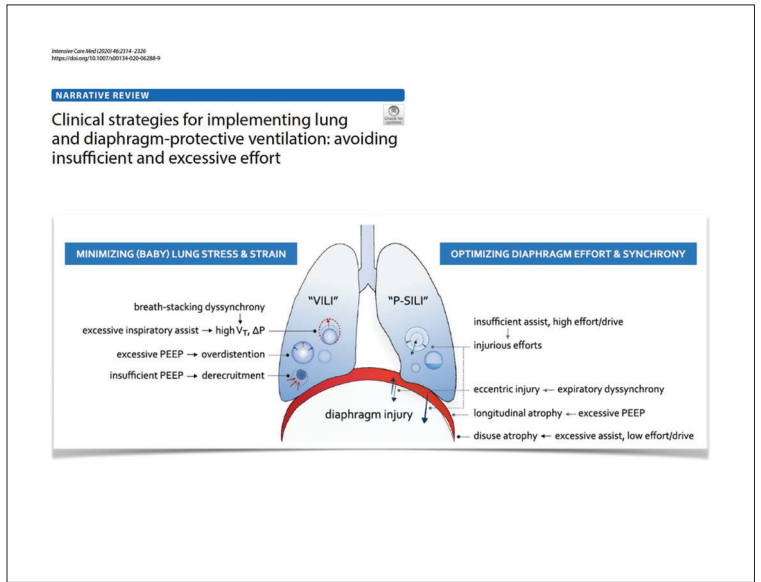
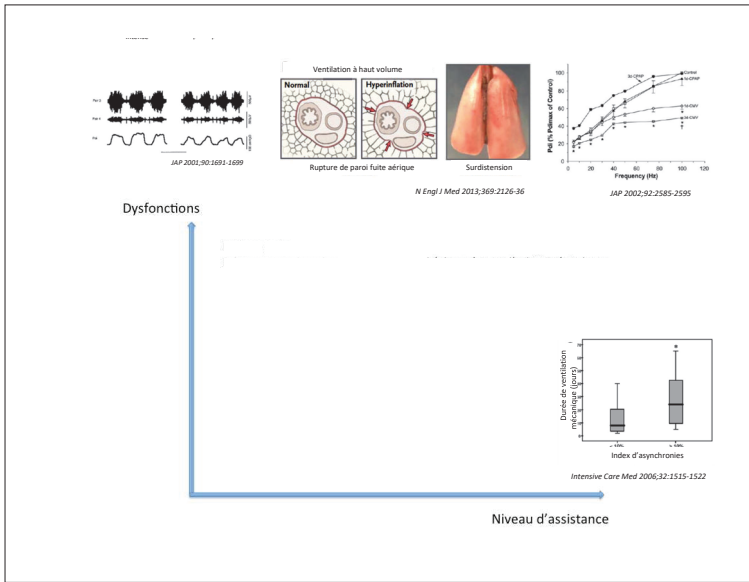


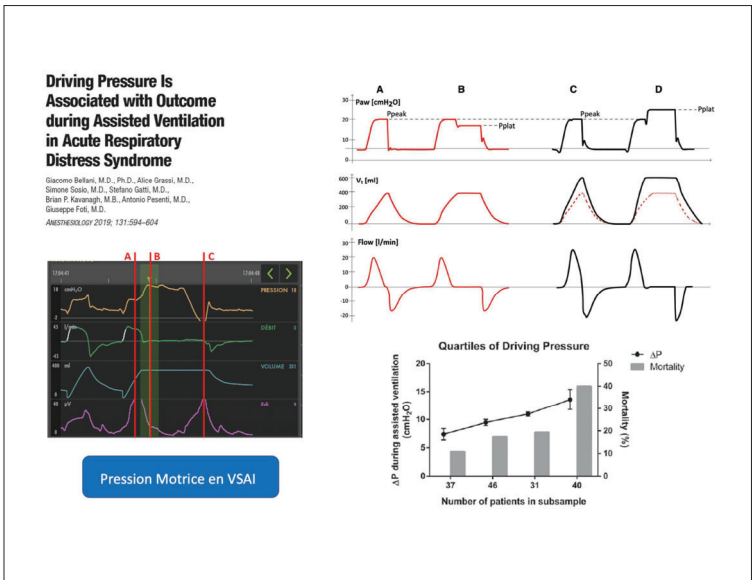
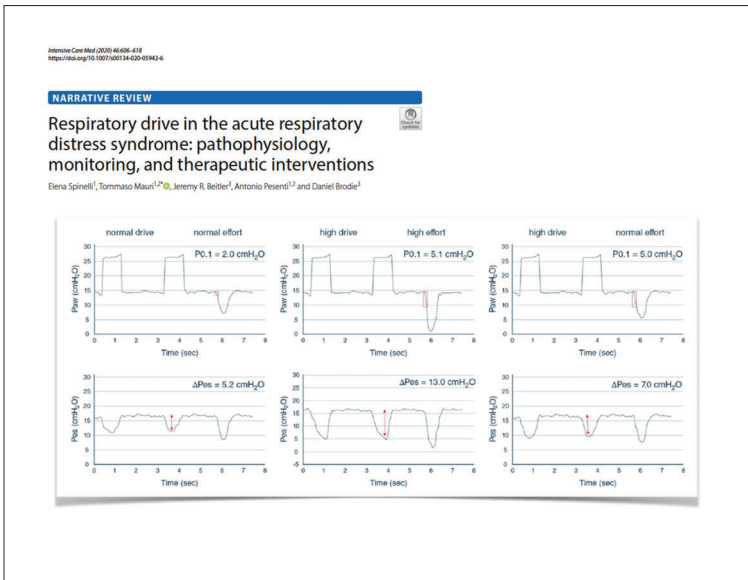
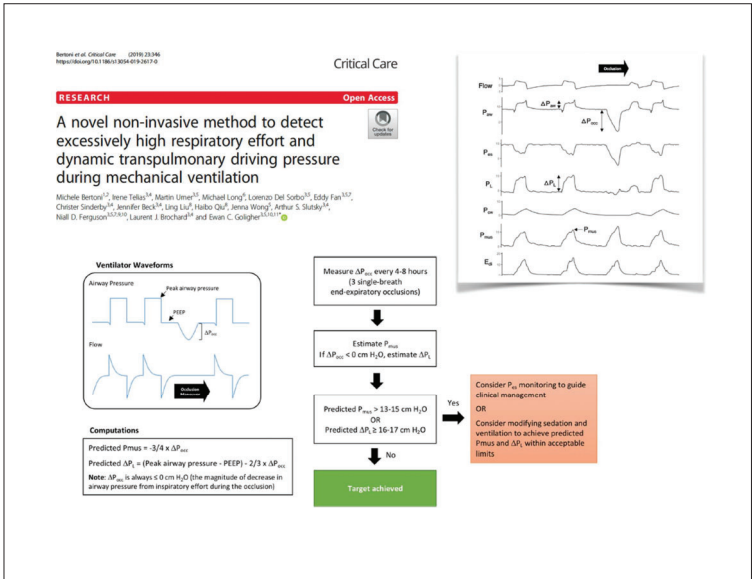
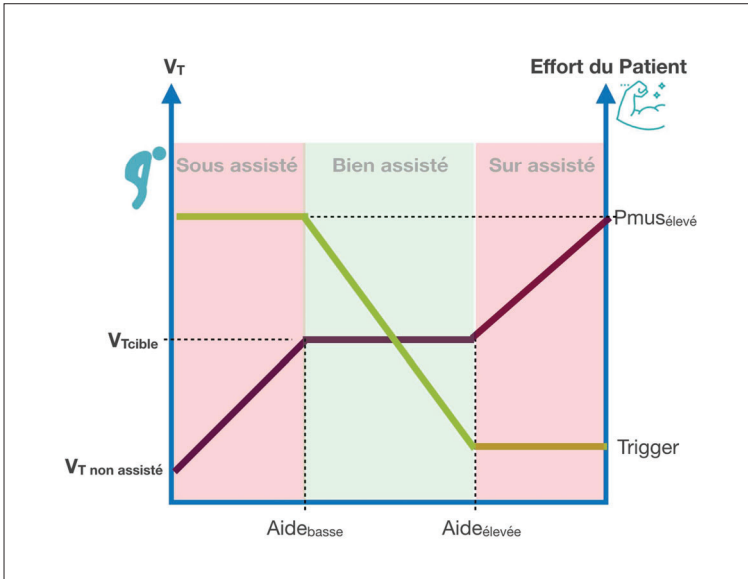
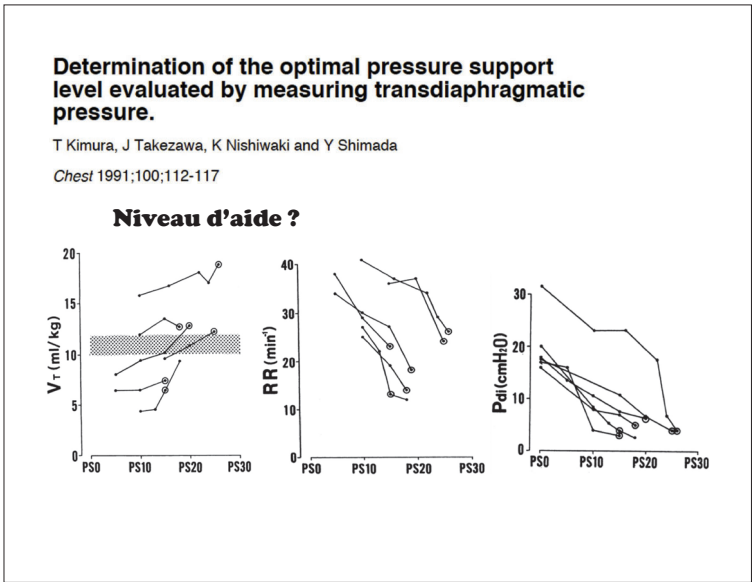
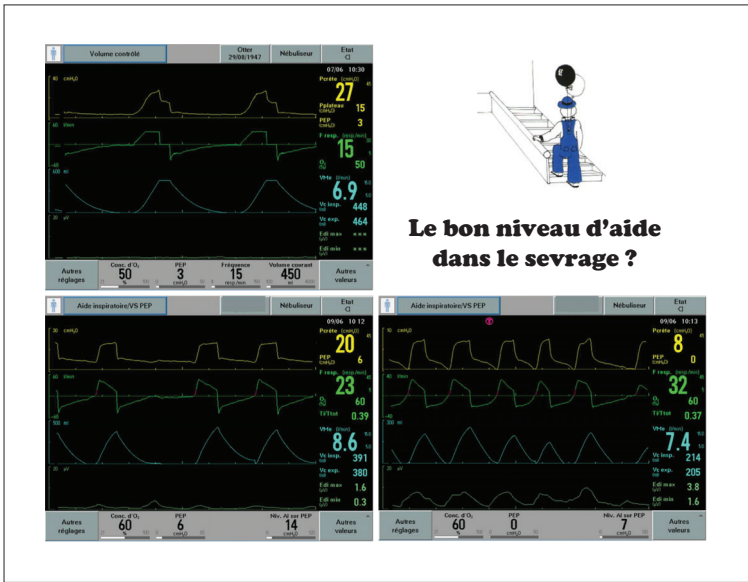
LA VENTILATION ASSISTEE LORS DU SEVRAGE

- Niveau d'aide
- Synchronisation

Problématiques du mode lors du sevrage

- Synchroniser le patient au respirateur/limiter les asynchronies
- Adapter l'aide inspiratoire aux besoins du patient
 - VSAI: aide fixe à adapter manuellement
- Intelligence artificielle en VSAI: Smartcare
- Neurally Adjusted Ventilatory Assist: NAVA
- Proportionnal Assist Ventilation: PAV+



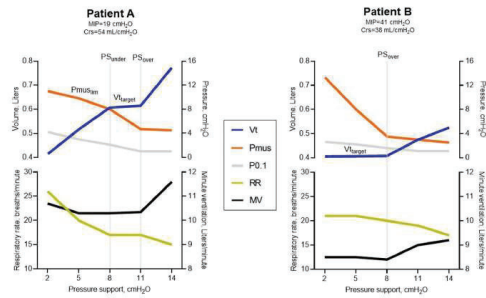


PERSPECTIVE

Open Access

Pressure support, patient effort and tidal volume: a conceptual model for a non linear interaction

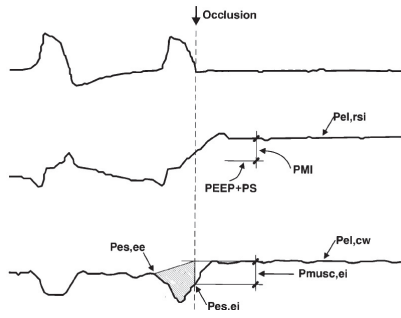
Mattia Docci^{1,2,3}, Giuseppe Foti^{1,4}, Laurent Brochard^{2,3} and Giacomo Bellani^{5,6*}



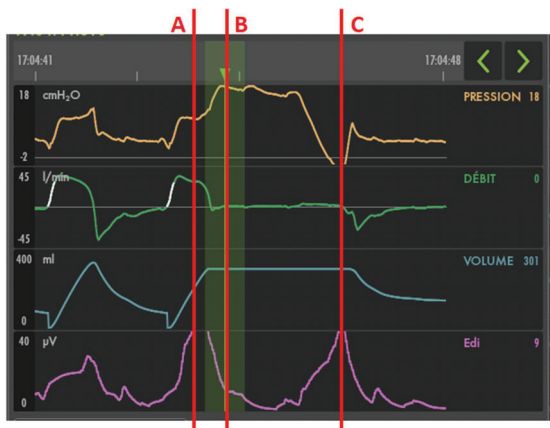
End-Inspiratory Airway Occlusion

A Method To Assess the Pressure Developed by Inspiratory Muscles in Patients with Acute Lung Injury Undergoing Pressure Support

GIUSEPPE FOTI, MAURIZIO CEREDA, GIULIANA BANFI, PAOLO PELOSI, ROBERTO FUMAGALLI and ANTONIO PESENTI



AM J RESPIR CRIT CARE MED 1997;156:1210-1216.



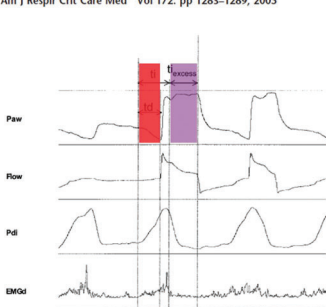
LA SYNCHRONIE PATIENT VENTILATEUR

La synchronie

Impact of Expiratory Trigger Setting on Delayed Cycling and Inspiratory Muscle Workload

Didier Tassaux, Marc Gallotier, Anne Battistini, and Philippe Jolliet

Am J Respir Crit Care Med Vol 172, pp 1283-1289, 2005

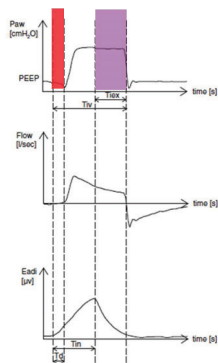


Intensive Care Med (2005) 10, 699-705

ORIGINAL

Neurally adjusted ventilatory assist improves patient-ventilator interaction

Low PEEP level
 Laurent Brochard
 Karim Brochard
 Anne Brochard
 Thierry Brochard
 Pierre-François Lemaire
 Philippe Jolliet
 Didier Tassaux



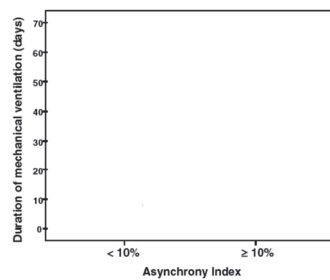
Problématiques des asynchronies?

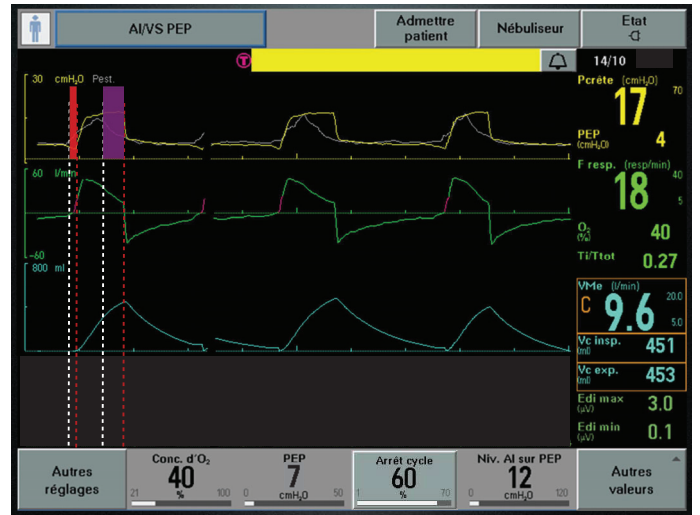
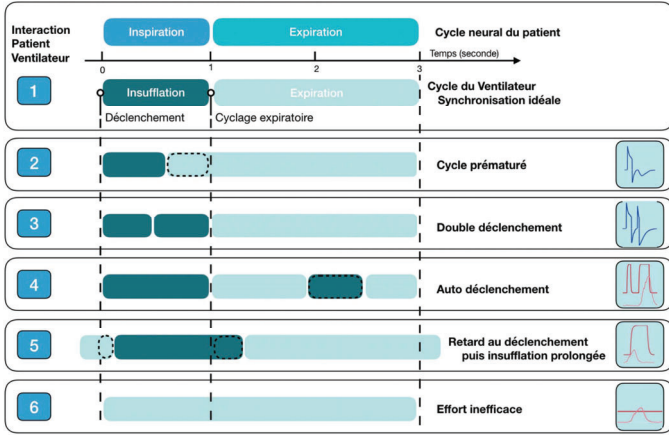
Intensive Care Med (2006) 11, 1515-1522
 DOI 10.1007/s00134-006-0301-8

ORIGINAL

Armand W. Thille
 Pablo Rodriguez
 Belen Cabello
 François Lellouche
 Laurent Brochard

Patient-ventilator asynchrony during assisted mechanical ventilation

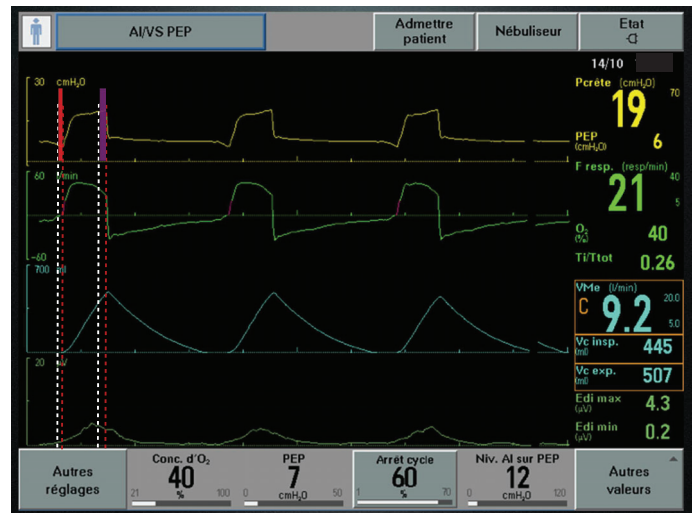
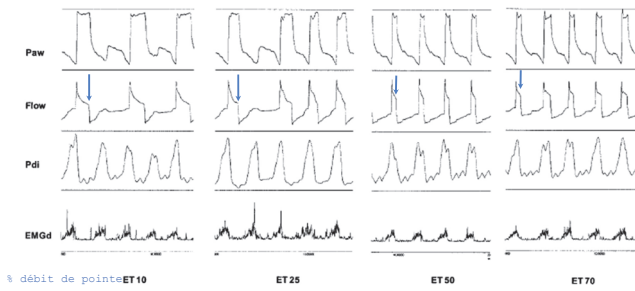




Impact of Expiratory Trigger Setting on Delayed Cycling and Inspiratory Muscle Workload

Didier Tassaux, Marc Gainnier, Anne Battisti, and Philippe Jolliet

Am J Respir Crit Care Med Vol 172. pp 1283–1289, 2005

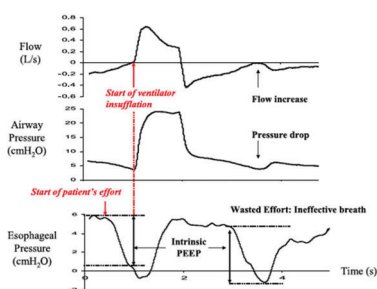


Intensive Care Med (2008) 34:1477–1486
DOI 10.1007/s00134-008-1121-9

ORIGINAL

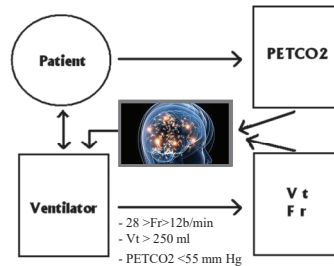
Arnaud W. Thille
Belen Cabello
Fabrice Galia
Aissam Lyazidi
Laurent Brochard

Reduction of patient-ventilator asynchrony by reducing tidal volume during pressure-support ventilation



Système Smartcare

- Système Néoganesh: règles, scénarios pour être dans une zone de confort: VTE FR EtCO₂ évaluation 2-3 min



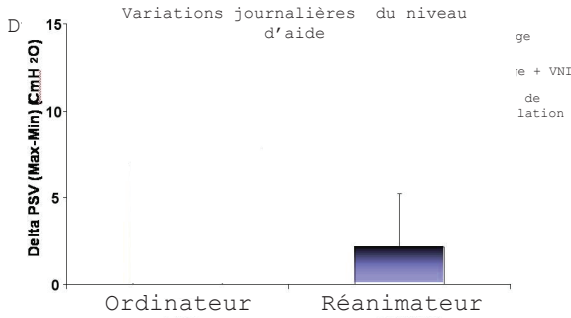
The knowledge base - 142 rules

SOME GENERAL RULES USED IN THE KNOWLEDGE-BASED SYSTEM

1. Never reduce the level of pressure support at night, except in case of excessive absolute ventilation.
2. Try to keep the patient within a zone of "comfort" with a respiratory rate between 12 and 28 breaths per minute, a tidal volume above a minimum threshold, and a PETCO₂ below a maximum threshold.
3. Do not let the tidal volume fall below 300 or 250 ml in patients whose body weight is above or below 55 kg, respectively.
4. Keep the respiratory rate between 12 and 28 breaths/min so that the patient is comfortable. In some patients, the upper limit can be moved up to 32 breaths/min.
5. Do not let the end-tidal CO₂ exceed 60 mm Hg in COPD patients and 50 mm Hg in patients with other disorders.
6. Decrease the level of pressure support by 2 cm H₂O when the patient has a stable ventilation within the comfort zone during at least 30 min with a level of pressure support less than 20 cm H₂O.
7. Decrease the level of pressure support by 4 cm H₂O when the patient has a stable ventilation within the comfort zone during at least 60 min with a level of pressure support of 20 cm H₂O or more.
8. Initiate the prewarning observation period when the level of pressure support is at the minimal value (5 or 9 cm H₂O) in intubated or in-tubated patients, respectively.
9. Consider that the patient is ready to be weaned after 1 or 2 h of stable ventilation at the minimal level of pressure support (1 h in those patients with a level of pressure support of 15 cm H₂O or less after 1 h of observation, 2 h in those with an initial level of pressure support greater than 15 cm H₂O).
10. Adjust the level of pressure support to the physiologic needs of the patient and evaluate every 2 min.
11. Consider that a patient requiring a PEEP level above 5 cm H₂O is not ready to be weaned.
12. The minimal level of pressure support is 40 cm H₂O.
13. In case of severe hypoventilation, switch to assist-control ventilation with preset parameters.

A Multicenter Randomized Trial of Computer-driven Protocolized Weaning from Mechanical Ventilation

François Lellouche, Jordi Mancebo, Philippe Joliet, Jean Roesler, Frédérique Schortgen, Michel Dojat, Bélen Cabello, Lila Bouadma, Pablo Rodriguez, Salvatore Maggiore, Marc Reynaert, Stefan Mersmann, and Laurent Brochard
Am J Respir Crit Care Med Vol 174. pp 894-900, 2006

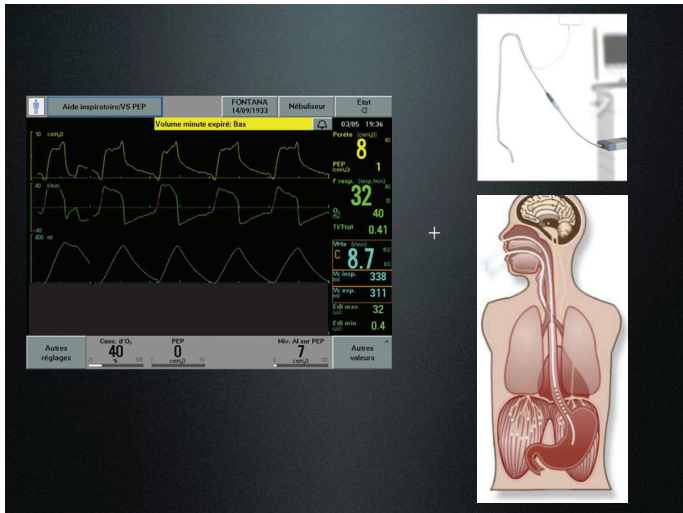
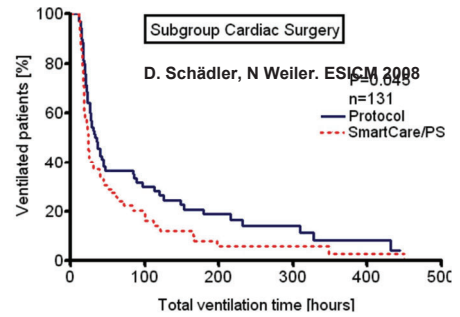


Intensive Care Med (2008) 34:1788-1795
DOI 10.1007/s00134-008-1179-4

ORIGINAL

Louise Rose
Jeffrey J. Presneill
Linda Johnston
John F. Cade

A randomised, controlled trial of conventional versus automated weaning from mechanical ventilation using SmartCare™/PS



SEPARATION DU VENTILATEUR

- INDEX PREDICTIFS
- TEST

The New England Journal of Medicine

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Volume 324 MAY 21, 1991 Number 21

A PROSPECTIVE STUDY OF INDEXES PREDICTING THE OUTCOME OF TRIALS OF WEANING FROM MECHANICAL VENTILATION

KARI L. YANO, M.D., AND MARTIN J. TORO, M.D.

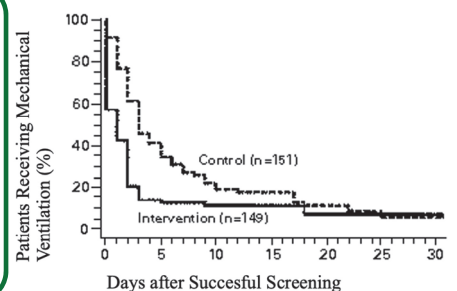
Premier Index FR/VT

Table 3. Effect of Duration of Mechanical Ventilation (<8 Days or ≥8 Days) on the Accuracy of Indexes in Predicting Weaning Outcome.*

INDEX	SENSITIVITY		SPECIFICITY		POSITIVE PREDICTIVE VALUE		NEGATIVE PREDICTIVE VALUE	
	<8 DAYS	≥8 DAYS	<8 DAYS	≥8 DAYS	<8 DAYS	≥8 DAYS	<8 DAYS	≥8 DAYS
Minute ventilation	0.79	0.75	0.75	0.08	0.65	0.35	0.40	0.33
Respiratory frequency	0.89	1.00	0.31	0.42	0.69	0.53	0.63	1.00
Tidal volume	1.00	0.88	0.50	0.58	0.78	0.58	1.00	0.88
Tidal volume/patient's weight	0.96	0.88	0.38	0.42	0.73	0.50	0.86	0.83
Maximal inspiratory pressure	1.00	1.00	0.00	0.25	0.64	0.47	1.00	1.00
Dynamic compliance	0.75	0.63	0.69	0.25	0.81	0.36	0.61	0.50
Static compliance	0.82	0.50	0.56	0.08	0.77	0.27	0.64	0.20
PaO ₂ /PaO ₂ ratio	0.79	0.88	0.38	0.17	0.69	0.41	0.50	0.67
Frequency/tidal volume ratio	1.00	0.88	0.63	0.67	0.82	0.64	1.00	0.89
CROP index	0.82	0.75	0.56	0.58	0.77	0.55	0.64	0.78

Identifying patients capable of breathing spontaneously and duration of mechanical ventilation

- Intervention Group
- 1) A daily screening of respiratory function (by the respiratory therapists of the unit)
 - PaO₂/FIO₂ > 200
 - PEEP < 5 cm H₂O
 - Adequate cough
 - f/Vt < 105 c/min
 - No vasopressor agents or sedatives
- 2) A 2-hour trial of spontaneous breathing
- 3) Notification of the physician of the successful results

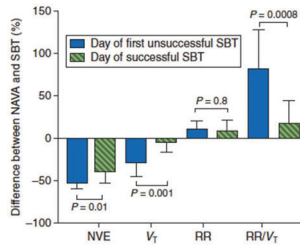


FEUX VERT

Ely E.W. et coll. N Engl J Med 1996; 335: 1864-9.

Neuro-ventilatory efficiency during weaning from mechanical ventilation using neurally adjusted ventilatory assist

H. Rozé^{1,2,3*}, B. Repusseau¹, V. Perrier¹, A. Germain¹, R. Séramondi¹, A. Dewitte^{1,3}, C. Fleureau¹ and A. Ouattara^{1,2}

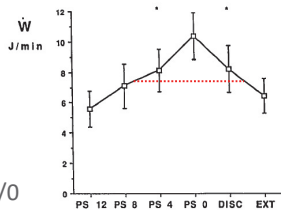


Augmentation de l'EAdi pour un même niveau de pression entre le début et la fin du sevrage



Inspiratory Pressure Support Compensates for the Additional Work of Breathing Caused by the Endotracheal Tube

Laurent Brochard, M.D.,¹ Fernando Riva, M.D.,¹ Hubert Lorino, Ph.D.,³ François Lemaire, M.D.,[§] Alain Harf, M.D., Ph.D.[¶]



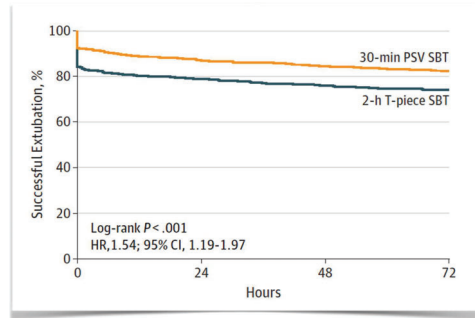
TEST VSAI 7/0

TABLE 1. Patient Characteristics

Patient	Age (yr)	Sex	Duration of mechanical ventilation (days)	Preop. Cause (%)	Preop. (%)	Diagnosis
1	81	F	5	104	0.50	CO2P, congestive heart failure
2	90	M	15	96	0.60	Refract. postoperative degenerative encephalopathy
3	80	M	3	149	0.40	CO2P, vascular surgery
4	76	M	3	132	0.55	CO2P, vascular surgery
5	78	F	24	111	0.59	CO2P, congestive heart failure
6	76	M	38	97	0.50	Chronic asthma, bacterial pneumonia
7	80	M	5	125	0.40	Abdominal surgery
8	61	F	14	105	0.50	Drug overdose
9	64	F	1	99	0.55	Cardiac surgery (myocardial)
10	18	M	3	99	0.55	Drug overdose
11	40	M	3	99	0.50	Severe meningitis

Effect of Pressure Support vs T-Piece Ventilation Strategies During Spontaneous Breathing Trials on Successful Extubation Among Patients Receiving Mechanical Ventilation: A Randomized Clinical Trial

Carlos Subiela, MD, Gonzalo Fernández, MD, PhD, Antonio Vilchez, MD, PhD, Raquel Rodríguez García, MD, Alejandro González Castro, MD, Carolina García, MD, Olga Rabal, MD, PhD, Luis Ventura, MD, Alexander López, MD, María Carmen de la Torre, MD, PhD, Elena Kovatch, MD, Valeria Arango, MD, Cecilia Hermosa, MD, Carmen Sánchez, MD, Ana Tello, MD, Eva Torres, MD, PhD, César Laborda, MD, Sara Gabalés, MD, Victoria Lacort, MD, María del Mar Fernández, MD, PhD, Anna Arnaiz, MD, PhD, Rafael Fernández, MD, PhD

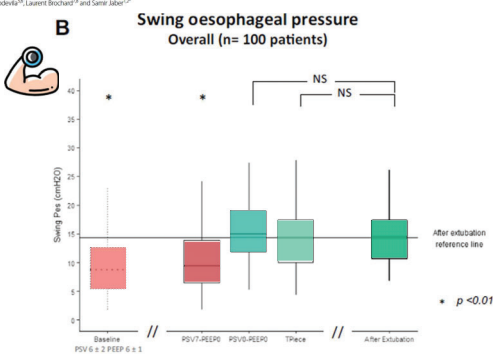


JAMA. 2019;321(22):2182-2182.

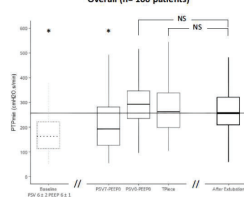
ORIGINAL
Spontaneous breathing trials should be adapted for each patient according to the critical illness. A new individualised approach: the GLOBAL WEAN study

Mathieu Capdeville^{1,2}, Yvan Assa^{1,3}, Clement Monot¹, Audrey De Jong^{1,4}, Aurélien Vignat¹, Julie Carli¹, Nicolas Molinari¹, Xavier Capdeville^{1,5}, Laurent Brochard¹ and Samir Jaber^{1,6}

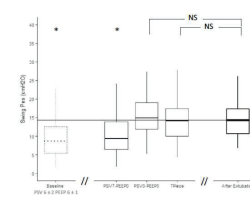
VSAI 7/0
Vs
VSAI 0/0



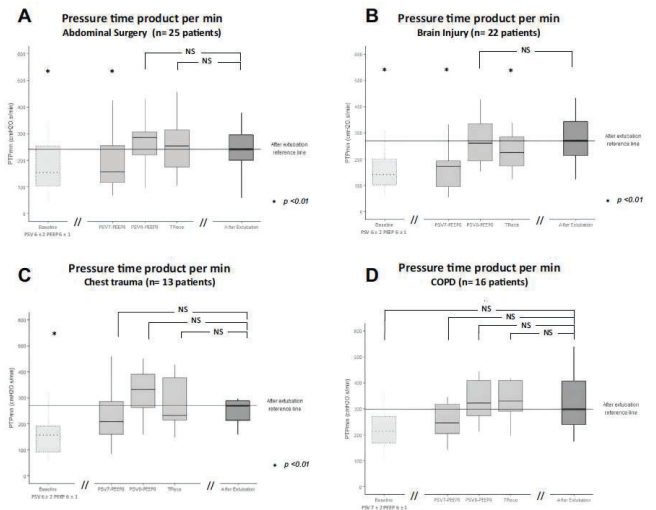
A Pressure time product per min Overall (n= 100 patients)



B Swing oesophageal pressure Overall (n= 100 patients)



Critical illness	SBT method	Significant difference in comparison to postextubation*
Overall (n=100)	PSV7-PEEP0	Yes
	PSV0-PEEP0	No
	TPiece	No
Abdominal surgery (n=25)	PSV7-PEEP0	Yes
	PSV0-PEEP0	No
Brain injury (n=22)	PSV7-PEEP0	Yes
	PSV0-PEEP0	No
Chest trauma (n=13)	PSV7-PEEP0	No
	PSV0-PEEP0	No
COPD (n=16)	PSV7-PEEP0	No
	PSV0-PEEP0	No
Miscellaneous (n=24)	PSV7-PEEP0	No
	PSV0-PEEP0	Yes
	TPiece	No

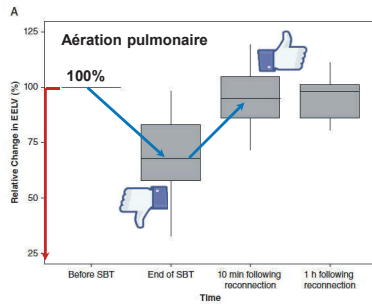


Critical Care Original Research

CHEST

Physiologic Effects of Reconnection to the Ventilator for 1 Hour Following a Successful Spontaneous Breathing Trial

Rémi Coudroy, PhD; Alice Lejars, MD; Maeva Rodriguez, MD; Jean-Pierre Frat, PhD; Christophe Rault, PhD; François Arrivé, MD; Sylvain Le Pape, PhD; and Arnaud W. Thille, PhD



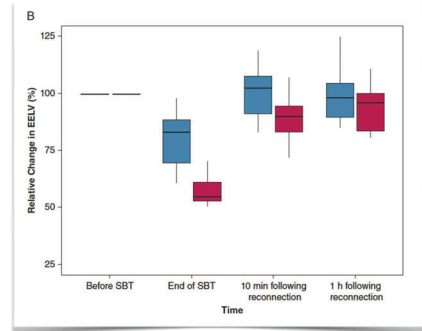
Critical Care Original Research

CHEST

Physiologic Effects of Reconnection to the Ventilator for 1 Hour Following a Successful Spontaneous Breathing Trial

Rémi Coudroy, PhD; Alice Lejars, MD; Maeva Rodriguez, MD; Jean-Pierre Frat, PhD; Christophe Rault, PhD; François Arrivé, MD; Sylvain Le Pape, PhD; and Arnaud W. Thille, PhD

■ VSAI 7/0
■ Tube en T



CHEST 2024;

The Lancet Respiratory Medicine
Volume 11, Issue 4, April 2021, Pages 319-328

Continued enteral nutrition until extubation compared with fasting before extubation in patients in the intensive care unit: an open-label, cluster-randomised, parallel-group, non-inferiority trial



A jeun 6 heures = Poursuite Nutrition jusqu'à extubation

Résultats essentiels

- Au total, 1150 patients ont été inclus en intention de traiter.
- L'interruption de la NE pré-extubation (n=513) n'a pas augmenté le risque de réintubation par rapport au groupe « poursuite NE » (n=617). En effet, dans la population en intention de traiter, 106 (17,2%) patients du groupe « poursuite NE » et 90 (17,5%) du groupe « arrêt NE » avaient un échec d'extubation dans les 7 jours suivant l'extubation (-0,4% ; IC95% -5,2 - 4,5).
- De plus, le délai entre test de ventilation spontanée / extubation et sortie de réanimation était raccourci dans le groupe « poursuite NE » ; hazard ratio à 1,39 (IC95%:1,13-1,71).

Noninvasive Positive-Pressure Ventilation for Postextubation Respiratory Distress: A Randomized Controlled Trial

JAMA
The Journal of the American Medical Association

Sean P. Keenan, MD, FRCPC, MSc
Caroline Powers, RRT
David C. McCormack, MD, FRCPC
Gary Block, MD, FRCPC

Outcomes	NPPV (n = 39)	Standard Therapy (n = 42)	P Value
Reintubation, No. (%)	28 (72)	29 (69)	.79
Pneumonia, No. (%)	16 (41)	17 (40)	.61
Duration of ventilation†			
Mean (SD)	8.4 (7.4)	17.5 (28.0)	.11
Median (range)	6.7 (0.5-28.6)	8.9 (2.0-146.7)	.12
ICU length of stay			
Mean (SD)	15.1 (10.9)	19.4 (25.0)	.32
Median (range)	11.9 (3.6-41.7)	10.8 (2.3-152.7)	.72
Hospital length of stay			
Mean (SD)	32.2 (25.4)	29.8 (28.4)	.69
Median (range)	19 (6-111)	22 (4-162)	.51
ICU survival, No. (%)	33 (85)	32 (76)	.34
Hospital survival, No. (%)	27 (69)	29 (69)	.99

JAMA. 2002;287:3238-3244

ORIGINAL ARTICLE

Noninvasive Positive-Pressure Ventilation for Respiratory Failure after Extubation

Andrés Esteban, M.D., Ph.D., Fernando Frutos-Vivar, M.D., Niall D. Ferguson, M.D., Yaseen Arabi, M.D., Carlos Apezteguia, M.D., Marco González, M.D., Scott K. Epstein, M.D., Nicholas S. Hill, M.D., Stefano Nava, M.D., Marco-Antonio Soares, M.D., Gabriel D'Empaire, M.D., Inmaculada Ala, M.D., and Antonio Anzueto, M.D.

CONCLUSIONS

Noninvasive positive-pressure ventilation does not prevent the need for reintubation or reduce mortality in unselected patients who have respiratory failure after extubation.

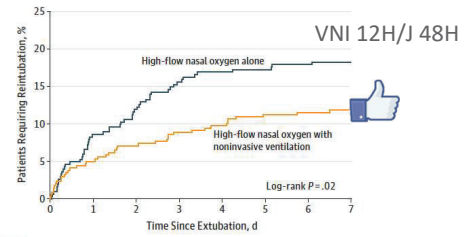
N Engl J Med 2004;350:2452-60.

Effect of Postextubation High-Flow Nasal Oxygen With Noninvasive Ventilation vs High-Flow Nasal Oxygen Alone on Reintubation Among Patients at High Risk of Extubation Failure A Randomized Clinical Trial



Sélection de patients à risque d'échec

Arnaud W. Thille, MD, PhD, Grégoire Muller, MD, Arnaud Gacouin, MD, Rémi Coudroy, MD, Maxime Decaville, MD, Romain Sonneville, MD, PhD, François Bellonci, MD, Christophe Gault, MD, Laurence Dangers, MD, Alexandre Laubette, MD, PhD, Séverin Cabasson, MD, Anahita Kouzli, MD, Emmanuel Vivier, MD, Anthony Le Meur, MD, Jean-Dominique Ricard, MD, PhD, Myriam Rozac, MD, Guillaume Barbier, MD, Christophe Lebet, MD, Stephan Ehrmann, MD, PhD, Caroline Sabatier, MD, Jeremy Bourouine, MD, Gaël Pradelle, MD, Pierre Bally, MD, Nicolas Terzi, MD, PhD, Jean-Delavrance, MD, PhD, Guillaume Laroche, MD, Pierre-Eric Darin, MD, Hodarou Ndaroumbe, MD, Aude Gibelin, MD, Laurence Zanes, MD, Nicolas Deye, MD, PhD, Alexandre Demour, MD, PhD, Adil Maamar, MD, Moez Ach-Nay, MD, René Robert, MD, PhD, Stéphanie Ragot, PharmD, PhD, Jean-Pierre Frate, MD, for the HIGH-WEAN Study Group and the REVA Research Network



No. at risk

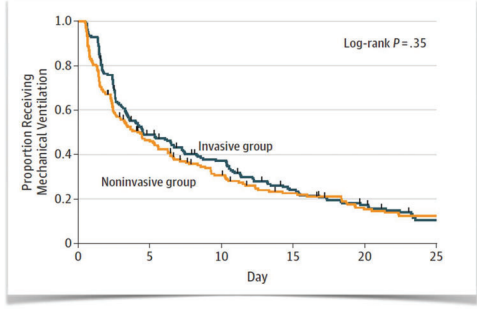
High-flow nasal oxygen	Alone	With
Alone	302	276
With	339	321
noninvasive ventilation	265	314
	253	308
	248	305
	246	294
	244	292
	243	291

Effect of Protocolized Weaning With Early Extubation to Noninvasive Ventilation vs Invasive Weaning on Time to Liberation From Mechanical Ventilation Among Patients With Respiratory Failure The Breathe Randomized Clinical Trial



Pas de précipitation

Geoff D. Perkins, MD, Dipesh Mishra, PhD, Simon Gates, PhD, Fang Gao, MD, Catherine Snelson, MB, Nicholas Hart, PhD, Luigi Camporota, PhD, James Varley, MB, Coralie Carls, MB, Ekanuraman Paramasivan, MB, Beverley Hodsdot, Daniel F. McArdle, MD, Timothy S. Walsh, MD through Blackboard, PhD, Louise Rowe, PhD, Sarah C. Lamb, DPM, Stavros Petros, PhD, Duncan Young, DM, Rajjit Lal, PhD, for the Breathe Collaborators



JAMA. 2018;320(18):1881-1888.

Conclusion

- Protocole de sevrage
- Screening quotidien des patients
- Adapter le niveau d'aide
- Améliorer l'interaction patient ventilateur
 - Synchronisation
 - Proportionnalité
 - Nouvel outil de monitoring au lit du patient: l'EAdi