Liberation from mechanical ventilation



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Weaning, liberation, discontinuing

- As soon as the <u>inciting factors</u> causing respiratory failure starts to improve, weaning may be initiated.
- Beginning with the day of intubation.
- All patients should be evaluated at least daily.
- The gradual process of transition from full ventilatory support to spontaneous breathing, including removal of endotracheal tube.
- Liberation or discontinuing ventilatory support. (overall process).
- Liberation from mechanical ventilation is a three-step process.

Outline



Classification of the patients



Five categories

- In the vast majority of ventilated patients, removal is quick and routine.
- Need a more systematic approach to discontinuing ventilatory support, normally about 15 - 20% of patients.
- Require <u>days to weeks</u> to wean from ventilatory support: < 5%
- Ventilator-dependent or "<u>unweanable</u>" patients: < 1%
- No chance for survival in whom the ventilator is discontinued while comfort measures are provided, terminal weaning or terminal extubation.

Evaluation of Systems to Determine Etiology of Respiratory Failure

- Neurologic factors
- Respiratory factors and ventilatory muscle function
- Metabolic factors
- Cardiovascular factors
- Psychological factors

Chest 120(6 Suppl):438S-444S, 2001

Evidence-Based Guidelines for Weaning and Discontinuing Ventilatory Support



Respir Care 40:244–259, 1995

Clinical criteria

- Improvement in the underlying cause of respiratory failure
- Adequate oxygenation
- Acid-base balance (Arterial pH > 7.25)
- Hemodynamic stability
- Ability to take spontaneous respiration



Weaning from mechanical ventilation: Readiness

Adequate oxygenation

- $P_aO_2/F_iO_2 \ge 150-200 \text{mHg}$
- SPO₂ 90% while receiving FiO₂ \leq 40% and PEEP \leq 5cmH₂O
- For chronic hypoxemia $\rightarrow P_aO_2/F_iO_2 \ge 120$ mHg
- Higher F_iO_2 (50%) may be acceptable in significant underlying lung disease.
- Higher PEEP (8 cmH₂O) may be acceptable to avoid atelectasis (obesity, abdominal distention, narrow ETT < 7mm)
- Normal Hb level, adequate cardiac output and tissue perfusion are assumed.



Egan's fundamentals of respiratory care 12th edition

Weaning from mechanical ventilation: Readiness t

PaO_2/F_iO_2 and SPO_2/F_iO_2

PaO_2/FiO_2	≥ 300	200	150	< 100
SpO ₂ /FiO ₂	315	235	190	150
FiO ₂	0.30	0.40	0.50	0.60
SpO ₂ (%)	> 94	94	95	< 90

Leasa et al. Crit Care (2021) 25:22

Hemodynamic stability

- Hemodynamic stable without myocardial ischemia
- Mean arterial pressure consistently > 60mmHg
- Systolic pressure > 90mmHg and < 180mmHg
- Use of vasopressors only in low and stable dose (dopamine < 5mcg/kg/min)



Weaning from mechanical ventilation: Readiness tes

Metabolic factors

- Nutrition should be adequate.
- 1.5-2.0 times resting energy expenditure.
- Protein intake should be 1-1.5g/kg per day.
- Avoid excessive carbohydrate \rightarrow CO₂ production.
- Avoid amino acid formulations (arginine/lysine) \rightarrow metabolic acidosis.

Metabolic factors

- Hypokalemia
- Hypocalcemia
- Hypophosphatemia
- Hypomagnesemia

Malnourished patients and those with chronic alcoholism

Hypothyroidism

Impaired respiratory muscle function.

Blunts the central response to hypercapnia and hypoxemia

Additional or optional requirement

- Hemoglobin level $\geq 7g/dL$
- Core temperature \leq 38.5C
- Mental status awake and alert or easily arousable $GCS \ge 8$
- RASS (Richmond agitation-sedation scale) -2 to +1
- As many as 47% of patients who spend >5 days in an ICU experience psychogenic disturbance

Outline



Requirements of an Ideal Weaning Index

- Assessment of the pathophysiological determinants of weaning outcome and psychological problems.
- <u>Accurately evaluate physiological function as it relates to the degree</u> of abnormality present.
- Ease of measurement and reproducible measurements.
- Minimum patient cooperation.
- High positive and negative predictive values.

Predictive factors of weaning from mechanical ventilation and extubation outcome: A systematic review



RSBI was the most frequently studied

RSBI was an important measurement tool in deciding whether to wean/extubate

A.R. Baptistella et al. / Journal of Critical Care 48 (2018) 56-62

RSBI (rapid shallow breathing index)

- RSBI < 105 (RR per minute \div V_T per liter) is a good <u>predictor for weaning</u>. 20 % may have false-positive results.
- RSBI of 80 is associated with 95% probability of successful liberation.

Chest. 1992;102:1829–1832

- RSBI around 50 \rightarrow successfully extubation.
- RSBI around 80 \rightarrow failed in the extubation.
- A discrepancy in the ideal RSBI score to predict weaning or extubation success.
- The most appropriate moment to measure RSBI ?

RSBI (rapid shallow breathing index) in predicting extubation outcome

- RSBI measured at <u>30 to 60 min</u> of SBT predicted the weaning outcome more effectively.
- RSBI at 120 min was significantly higher in patients with <u>extubation</u> failure and trial failure.
- When RSBI was <u>measured every 30 min</u> during 2 h of SBT.
 Initial RSBI was similar in extubation success and failure groups.
 RSBI remained unchanged or decreased in the success group.
 RSBI increased in the extubation failure group.

Weaning parameters in elderly patients

- An important difference between the age and weaning.
- With age split into quartiles (≤42, 43–54, 55–62, and 63+ years), the percentages of successful attempts decrease with increasing age (91%, 91%, 87%, and 84%, respectively)
- <u>Age > 65</u> is a <u>negative predictor</u> of weaning and extubation success.
- The longer the duration (in days) of IMV, the lower the chance of success in weaning and extubating.
- The <u>age + MV days</u> >100 or more predicted a poor outcome.

The maximum inspiratory pressure (MIP)

- Mechanical ventilation causes rapid diaphragmatic wasting, weakening the very muscle.
- (MIP) is a good parameter to <u>determine respiratory muscular capacity</u>, a predictive factor for weaning success.
- A successful weaning outcome was likely if MIP values < -30 cmH₂O and a weaning failure was likely if MIP < -20 cmH₂O
- Wide range of normal values can be closely related to the voluntary effort.

Measurement of drive to breathe P_{0.1}



Ultrasound (US) evaluation of the diaphragm

- A marker for diaphragmatic function and directly affect weaning and extubation outcomes
- The percent change in diaphragm thickness (T_{di}) ≥ 30% between endexpiration and end-inspiration (ΔT_{di}%), evaluated in the zone of apposition, has a sensitivity and specificity for extubation success of 88% and 71%
- $\Delta T_{di} \% > 20$ is a robust predictor of extubation success within 48 hr.
- ΔT_{di} % > 34.2 is a cutoff value associated with successful extubation.

Diaphragm Ultrasound in Weaning From Otheck for updates Mechanical Ventilation

Deepti Kilaru, MBBS; Nova Panebianco, MD, MPH; and Cameron Baston, MD, MSCE

- A potential contributing factor of weaning failure is diaphragm weakness or atrophy.
- Ventilator- induced diaphragm dysfunction (VIDD).
- Each day on the ventilator has been associated with <u>a reduction in</u> diaphragmatic thickness by an <u>average of 6%.</u>
- Mortality is higher in diaphragmatic dysfunction. (49% vs 7%)
- More commonly seen when mechanical ventilation is combined with sepsis.

Linear transducer placed longitudinally

B mode ultrasound image Musculoskeletal preset





Anterior axillary line Right 8th – 10th intercostal space

Cephalad

Diaphragm thickness Zone of apposition

Caudal

Diaphragm thickness

Diaphragm measured at the end of inspiration

Diaphragm measured at the end of expiration



Clinical application of diaphragm ultrasound

- Diaphragm thickening fraction (DTF) and diaphragm excursion are the variables that are assessed for predicting weaning.
- Thickening fraction =

[(End Inspiratory Diaphragm Thickness - End Expiratory Diaphragm Thickness) / End Expiratory Diaphragm Thickness] × 100.

- DTF >30% to 36% has been proposed as a <u>cutoff to predict successful</u> weaning from mechanical ventilation.
- Sensitivity range from 82% to 88% and specificity 71% to 88%
- DTF < 20% predictive of diaphragm paralysis.

Clinical application of diaphragm ultrasound

- DTF should be <u>obtained during a SBT</u> to help predict success in weaning.
- The performance characteristics of DTF are better at lower levels of pressure support (PS 10cmH₂O).
- More accurately demonstrates the expected post-extubation diaphragmatic workload.
- When compared directly with RSBI, DTF is more specific for predicting successful extubation.





Clinical application of diaphragm ultrasound

- Diaphragm excursion can be used to predict failure of extubation.
- An excursion > 10 to 14 mm has been shown to predict successful extubation with a sensitivity (78.9% to 87.5%) and specificity (70.8% to 71.5%)
- Even a completely paralyzed diaphragm will have an excursion proportional to lung compliance and driving pressure.
- Specific threshold during spontaneous breathing trials for all patients ?

The APACHE II and SOFA

- <u>APACHE II</u> scores were statistically higher in the weaning/extubation failure group.
- <u>May not always reflect the current state</u> of a patient, as the score is applied within 24 h of patient's admission to the ICU, and weaning can start several days later.
- The SOFA score is normally used continuously during patients' ICU stay, which accounts for the severity of a patient's illness <u>at the</u> moment of weaning or extubation.
- A more reliable predictor of weaning outcome.

The daily SOFA score



Nutrition and anemia

- Malnutrition \rightarrow impairment in respiratory function.
- Fatigue, decreased respiratory muscle strength and endurance, depletion of diaphragmatic muscle mass.
- Total protein, albumin and creatinine height index correlated with the weaning outcome.
- Anemia can exacerbate the insufficient global O₂ delivery (DO₂) and myocardial ischemia.
- Hb <10 g/dL more likely to have <u>unsuccessful extubations</u>.

Fluid and renal function

- Side effect of IMV is hypotension, caused by a reduction in venous return.
- <u>Positive fluid balance</u> in the 24 h prior to extubation can predict the extubation failure.
- From last 48 h, 72 h and even in accumulation since hospital admission is a significantly greater predictor of weaning failures.
- Parameters linked to renal function, such as <u>BUN</u>, <u>creatinine</u> and the patients' need for <u>hemodialysis</u>, can also predict the weaning and extubation outcome.

Respir Care 2014;59:1042-7

Respirology 2014;19:576-82

Outline


Methods of titrating ventilator support during weaning

- Simultaneous intermittent mandatory ventilation (SIMV)
- Pressure support ventilation (PSV)
- T-piece weaning (using a ventilator)

SIMV



Should not be used as a weaning mode.

Prolongs the weaning phase and total length of ventilatory support.

The least effective method.

Respir Care 47:1007-1017, 2002

Closed–loop control modes for ventilator discontinuation

- Automated tube compensation (ATC)
- Volume-target pressure support ventilation
- Automode and variable pressure support/variable pressure control
- Mandatory minute ventilation
- Adaptive support ventilation
- Artificial intelligence system

Pilbeam's Mechanical Ventilation Physiological and Clinical Applications, 6e 6th Edition

Potential advantages of ATC

- Support or overcome the WOB imposed by the artificial airway.
- Improve patient-ventilator synchrony through variable compensation of inspiratory flow based on patient demand.
- Unload the inspiratory muscles and increase alveolar ventilation without adverse cardiopulmonary side effects.
- <u>Reduce the risk of air trapping</u> caused by expiratory resistance from the endotracheal tube.
- Facilitate accurate prediction of readiness for extubation.



Control of the expiratory valve

Spontaneous breathing trial (SBT)

- Perhaps the best way to determine readiness to wean is a carefully supervised SBT.
- Consider ready for weaning is tolerate an SBT for 30-120min.
- 77-85% of patients who pass an SBT can be successfully weaned and extubated.
- Simple T-piece (using a ventilator)
- CPAP (5cmH₂O)
- Low level of pressure support (5-8cmH₂O)

Liberation From Mechanical Ventilation in Critically Ill Adults



Executive Summary of an Official American College of Chest Physicians/American Thoracic Society Clinical Practice Guideline

- For acutely hospitalized patients ventilated >24 h.
- We suggest that the initial SBT be conducted with inspiratory pressure augmentation (PS 5-8 cm H₂O) rather than without (T-piece or CPAP)

JAMA | Original Investigation | CARING FOR THE CRITICALLY ILL PATIENT

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

- 1153 patients, 18 ICUs in Spain.
- A SBT consisting of 30 minutes of PSV (PS 8cmH₂O), compared with 2 hours of T-piece ventilation.
- PSV led to significantly higher rates of successful extubation.
- These findings support the use of a shorter, less demanding ventilation strategy for spontaneous breathing trials.

PSV: higher successful extubation

Figure 2. Probability of Successful Extubation After First SBT in Each Group



	No./Total		
	30-min PSV SBT	2-h T-Piece SBT	Risk Ratio (95% CI)
ge, y			
≤70	306/370	289/385	1.10 (1.02-1.19)
>70	167/205	139/193	1.13 (1.01-1.26)
ength of mechanical ventilation, d			
≤4	263/304	249/326	1.13 (1.05-1.22)
>4	210/271	179/252	1.09 (0.99-1.21)
PACHE II score			
≤20	329/410	296/404	1.10 (1.02-1.18)
>20	133/164	132/174	1.07 (0.96-1.20)
)PD			
No	387/465	351/460	1.09 (1.02-1.16)
Yes	86/110	77/118	1.20 (1.02-1.41)
atient admission type			
Medical	329/404	283/396	1.14 (1.05-1.23)
Nonrespiratory	182/215	154/206	1.13 (1.03-1.25)
Respiratory	147/189	129/190	1.15 (1.01-1.30)
Surgical	114/140	114/142	1.10 (0.91-1.14)
Planned	33/35	23/29	1.19 (0.97-1.46)
Emergency	81/105	91/113	0.96 (0.83-1.10)
Trauma	30/31	31/40	1.25 (1.04-1.49)

Figure 3. Unadjusted Risk Ratios for Successful Extubation After First SBT in Predefined Subgroups

JAMA. 2019;321(22):2175-2182

Risk Ratio (95% CI)

2

0.6

Favors

PSV SBT

PSV

Comparison of T-piece and pressure support ventilation as spontaneous breathing trials in critically ill patients: a systematic review and meta-analysis



Yuting Li, Hongxiang Li and Dong Zhang*

- 10 RCT, 3165 patients
- T-piece and PSV as SBTs are considered to have <u>comparable predictive</u> power of successful extubation in critically ill patients.
- No significant difference in the rate of reintubation, ICU and hospital length of stay, and ICU and hospital mortality



	T-Piec	ce	PS			Odds Ratio	Odds Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95% CI	M-H, Fixed, 95% Cl
Chittawatanarat 2018	38	260	26	260	16.0%	1.54 [0.91, 2.62]	
Esteban 1997	36	246	38	238	23.8%	0.90 [0.55, 1.48]	
Koh 2000	4	22	4	20	2.5%	0.89 [0.19, 4.15]	
Santos Pellegrini 2018	35	99	40	91	19.4%	0.70 [0.39, 1.25]	Rate of
Subira 2019	58	578	59	575	38.3%	0.98 [0.67, 1.43]	reintubation
Total (95% CI)		1205		1184	100.0%	0.99 [0.78, 1.26]	
Total events	171		167				
Heterogeneity: Chi ² = 4.2	20, df = 4 (P = 0.3	8); l² = 5°	%			
Test for overall effect: Z	= 0.06 (P =	= 0.95)	-				0.01 0.1 1 10 100 Favours T-Piece Favours PS







	т	-Piece			PS			Mean Difference		Mean Differ	ence	
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Fixed, 95% Cl		IV, Fixed, 9	5% CI	
Chittawatanarat 2018	17.3333	10.4366	260	18.5	11.9276	260	51.0%	-1.17 [-3.09, 0.76]				
Santos Pellegrini 2018	32.5	31.5973	99	32	33.1449	91	2.2%	0.50 [-8.73, 9.73]				
Subira 2019	26	17.8362	578	26.3333	18.5796	575	42.8%	-0.33 [-2.44, 1.77]				
Teixeira 2015	25.1	16.5	66	27.6	19.8	46	3.9%	-2.50 [-9.47, 4.47]		-		
Total (95% CI)			1003			972	100.0%	-0.82 [-2.20, 0.55]				
Heterogeneity: Chi ² = 0.6	3, df = 3 (P = 0.89);	l² = 0%)					⊢ -100	-50 0		I 100
Test for overall effect: Z =	= 1.17 (P =	= 0.24)			E	osp	otal		-100	Favours T-Piece Fa		100
Fig. 10 Forest plot for hospital length of stay						LC	S					

Clinical Signs and Symptoms Indicating Problems during a SBT

- Respiratory rate > 30 to 35 /min (clinicians also should watch for 10 / min or RR < 8 /min).
- Tidal volume (V_r) \downarrow < 250 to 300mL
- Deterioration of ABG values and SpO₂
- BP changing significantly, as demonstrated by
 - SBP \downarrow 20 mmHg or \uparrow 30 mmHg
 - SBP >180 mmHg or a change of 10 mmHg diastolic
- Heart rate ↑ > 20% or > 140 / min.
- Sudden onset of frequent PVC > 4 to 6/min.

Patient evaluation during SBT

- Irregular ventilatory pattern.
- Palpable scalene muscle during inspiration.
- Palpable abdominal muscle tensing during expiration.
- Inability to alter the ventilatory pattern on command.
- 1 or 2 signs \rightarrow usually need continue support.
- \geq 3 signs \rightarrow poor prognosis for ventilator removal

Hess DR. Mechanical ventilation of the adult patient

Rule of thumb in initial approach to liberating

- SBT is the best approach.
- Ideally applied with <u>zero PS and zero PEEP</u> for 30-120 min via the mechanical ventilator. (for monitoring)
- <u>Single daily trial</u> may be preferred.
- Patients who have been received MV for >72 hr.
- The most common method is SBT interspersed with continued ventilatory support.
- Patient should not be overstressed, exhaustion \rightarrow delay weaning.
- It will take >24 h to completely recover from fatique.

AMERICAN THORACIC SOCIETY DOCUMENTS

An Official American Thoracic Society/American College of Chest Physicians Clinical Practice Guideline: Liberation from Mechanical Ventilation in Critically III Adults

Rehabilitation Protocols, Ventilator Liberation Protocols, and Cuff Leak Tests

- For acutely hospitalized adults who have been mechanically ventilated for more than 24 hours.
- Suggest protocolized rehabilitation directed toward early mobilization (conditional recommendation, low certainty in the evidence).
- A ventilator liberation protocol is suggested.

Daily screening liberation assessment

- Global variation exists in the use of protocols.
- Written directives to screen for readiness 5-83%
- Frequency of daily screening varied widely.
- Liberation protocols may not be beneficial in all settings.
- Should be tailored to the patients population.
- Not superior to usual care in highly staffed, closed ICU in an academic hospital.

AJRCCM. 2004;169(6):673

JAMA 2021 Mar 23; 325(12): 1173–1184

Recommendations in SBT

- An SBT should be performed every 24 hours.
- Avoid pushing patients to the point of exhaustion during the weaning process because this ultimately can delay liberation.
- It is important that clinicians wait >24 hours before attempting another SBT in patients for whom it fails.
- Frequent SBTs over a single day are not helpful and can lead to serious consequences.
- Even twice-daily SBTs offer no advantage over testing once a day.

Outline



Weaning and extubation

- > 10% of extubations fail, and IMV has to be reinitiated.
- The pathogenesis of weaning and extubation failure may differ significantly.
- "weaning success" when a patient successfully passes an SBT.
- "extubation success" when a patient is extubated after the SBT and is <u>not</u> reintubated during the next 48 h.

Eur Respir J 2007;29:1033-1056

If failed

- 7 times more likely to die.
- 31 times more likely to spend \geq 14 days in ICU.
- <u>6 times more likely to need transfer to long term or rehabilitation facility.</u>

CHEST 1997; 112:186-92

Incidence of failed extubation and mortality

CHEST /112 / 1 1JULY, 1997

Study (Year)	Patient Type	No. of Patients	% Reintubated	% Mortality
Sahn et al (1973) ⁵	MICU/SICU	100	17.0	NR
Hilberman et al $(1976)^8$	Cardiac surgery	124	17.7	NR
Tahvanainen et al (1983) ⁶	MICU	47	19.0	22.2
DeHaven et al (1986) ¹⁰	SICU/trauma	48	6.3	NR
Demling et al (1988) ¹	General SICU	400	5.5	40.0
0	Burn/trauma unit	300	3.3	10.0
Krieger et al (1989) ⁷	MICU/SICU	269	10.4	NR
Sassoon et al (1993)11	MICU	40	12.5	NR
Mohsenifar et al (1993) ¹²	RICU	29	14.3	NR
Lee et al (1994) ³	MICU	52	17.0	-33.3
Brochard et al (1994)9	MICU/SICU	109	11.0	NR
Torres et al $(1995)^2$	MICU/SICU	170	23.5	35.0
Esteban et al (1995) ⁴	MICU/SICU	530	15.7	NR
Current study	MICU	289	14.5	42.5
*MICU=medical ICU; SICU=s	urgical ICU; RICU=respiratory	ICU; NR=not reported.	3.3-23.5%	22.2-42.5%

Table 5—Studies Reporting the Incidence of Failed Extubation*

Incidence of failed extubation and mortality

AJRCCM Vol 187, Iss. 12, pp 1294–1302, Jun 15, 2013

TABLE 1. RATES OF PLANNED EXTUBATION FAILURE AND MORTALITY

Study (Reference)	Number of Extubations	Rate of Extubation Failure [% (<i>n</i>)]		ICU Mortality in Reintubated Patients [% (<i>n</i>)]	ICU Mortality in Nonreintubated Patients (%)
Esteban <i>et al.,</i> 1997 (1)	397	19 (74)	['	27 (20)	3
Esteban <i>et al.</i> , 1999 (2)	453	13 (61)	1 '	33 (20)	5
Epstein <i>et al.</i> , 1997 (4)	287	14 (40)	1 '	43 (17)	12
Vallverdu et al., 1998 (3)	148	15.5 (23)	1 '	35 (8)	5.6
Thille <i>et al.</i> , 2011 (6)	168	15 (26)	1 '	50 (13)	5
Frutos-Vivar <i>et al.</i> , 2011 (14)	1,152	16 (180)	1 '	28 (50)	7
Funk <i>et al.</i> , 2009 (38)	257	10 (26)	1 '	Not available	Not available
Tonnelier <i>et al.</i> , 2011 (39)	115	10 (12)	1 '	Not available	Not available
Sellares et al., 2011 (34)	181	20 (36)	1 '	Not available	Not available
Peñuelas et al., 2011 (40)	2,714	10 (278)	 ′	26 (72)	5
		10-19%		26-50%	

Risk factors for post-extubation stridor

- Prolonged intubation (variably defined as ≥36 hours to ≥6 days)
- Age > 80 years
- <u>A large ETT</u>

(>8 mm in men, >7 mm in women)

- A ratio of ETT to laryngeal diameter greater than 45% on CT
- A small ratio of patient height (mm) to ETT diameter (mm)

- An elevated APACHE II score
- A GCS score <8
- Traumatic intubation
- Female gender
- A history of <u>asthma</u>
- Excessive tube mobility due to insufficient fixation
- Insufficient or lack of sedation
- Aspiration



Extubation management in the adult intensive care

Cuff leak test

- Qualitative: stethoscope
- Quantitative assessment assessment: Cuff leak volumes
- Cuff leak volumes ≥ 110 mL or >24% of the delivered tidal volume is considered a normal cuff leak test.

Extubation management in the adult intensive care u

- <u>Simultaneous assessment of both cough and cuff leak may improve</u> prediction of post extubation stridor.
- The absence of both an audible cough and a cuff leak indicates the patient is 10 times more likely to develop post extubation stridor.

J Crit Care. 2004;19(1):23

AMERICAN THORACIC SOCIETY DOCUMENTS

An Official American Thoracic Society/American College of Chest Physicians Clinical Practice Guideline: Liberation from Mechanical Ventilation in Critically III Adults

Rehabilitation Protocols, Ventilator Liberation Protocols, and Cuff Leak Tests

- Performing a <u>cuff leak test</u> in mechanically ventilated adults who meet extubation criteria and are deemed <u>high risk</u> for post extubation stridor.
- For adults who have <u>failed a cuff leak test</u> but are otherwise ready for extubation, we suggest administering <u>systemic steroids</u> for ≥4 hours before extubation.





Akira Kuriyama, MD, MPH; Noriyuki Umakoshi, MD; and Rao Sun, MD, PhD

CrossMark

- Prophylactic Corticosteroids for Prevention of Postextubation Stridor and Reintubation in Adults A Systematic Review and Meta-analysis
- Administration of prophylactic corticosteroids before elective extubation was associated with significant reductions in the incidence of postextubation airway events and reintubation (57% reduction), with few adverse events.
- It is reasonable to select patients at high risk for airway obstruction who may benefit from prophylactic corticosteroids.

Relative risk of post extubation airway events

Trial (Year)	Favors Corticosteroids	Favors Control	RR (95% CI)	Corticosteroids event/total	Control event/total	Weight
Participants selected	with cuff-leak test					
Cheng (2006)			0.31 (0.14-0.69)	8/85	13/43	9.61
Lee (2007)			0.36 (0.13-1.05)	4/40	11/40	7.60
Baloch (2010)			0.40 (0.17-0.94)	6/46	15/46	9.16
Cheng (2011)			0.40 (0.17-0.94)	6/38	20/33	9.22
Yu (2014)			0.33 (0.17-0.67)	11/109	16/53	10.55
Lin (2016)			0.28 (0.12-0.65)	7/83	13/43	9.27
Subtotal ($I^2 = 0.0\%$,	P = .99)		0.34 (0.24-0.48)			
Unselected participar	nts					
Unselected participar						
Gaussorgues (1987)			2.00 (0.37-10.74)		2/138	
Gaussorgues (1987) Darmon (1992)		→	0.67 (0.32-1.40)	11/327	17/337	10.11
Gaussorgues (1987) Darmon (1992) Ho (1996)		→ 	0.67 (0.32-1.40) 0.68 (0.29-1.61)	11/327 7/39	17/337 10/38	10.11 9.14
Gaussorgues (1987) Darmon (1992) Ho (1996) Shih (2007)		> 	0.67 (0.32-1.40) 0.68 (0.29-1.61) 1.22 (0.56-2.69)	11/327 7/39 9/49	17/337 10/38 11/49	9.14 9.73
Gaussorgues (1987) Darmon (1992) Ho (1996)		 	0.67 (0.32-1.40) 0.68 (0.29-1.61)	11/327 7/39 9/49	17/337 10/38	10.11 9.14
Gaussorgues (1987) Darmon (1992) Ho (1996) Shih (2007)		 	0.67 (0.32-1.40) 0.68 (0.29-1.61) 1.22 (0.56-2.69)	11/327 7/39 9/49 11/355	17/337 10/38 11/49	10.11 9.14 9.73

Relative risk of reintubation

Trial (Year)	Favors Corticosteroids	Favors Control	RR (95% CI)	Corticosteroids event/total	Control event/total	Weight
Participants Selecte	ed with cuff-leak test					
Cheng (2006)		-	0.32 (0.11-0.91)	5/85	8/43	18.87
Lee (2007)			0.50 (0.05-5.30)	1/40	2/40	4.59
Baloch (2010)		-	0.22 (0.05-0.97)	2/46	9/46	10.82
Cheng (2011)			0.26 (0.08-0.87)	3/38	10/33	15.31
Yu (2014)		• · · · · · · · · · · · · · · · · · · ·	0.97 (0.18-5.14)	4/109	2/53	8.74
Lin (2016)			0.52 (0.08-3.55)	2/83	2/43	6.72
Subtotal (/2 = 0.0%	P = -80		0.35 (0.20-0.64)			
Unselected Particip Gaussorgues (1987		* *	5.00 (0.24-103.20)		0/138	2.85
		→	5.00 (0.24-103.20)	2/138	0/138	2.85
Darmon (1992)		+	0.41 (0.08-2.11)		5/337	9.06
Ho (1996)			0.33 (0.01-7.74)		1/38	2.61
Shih (2007)			1.25 (0.36-4.38)	5/49	4/49	14.30
François (2007)	<		0.07 (0.01-0.52)	1/355	14/343	6.13
Subtotal (<i>I</i> ² = 49.6)	%, <i>P</i> = .09)		0.53 (0.15-1.89)			
Total (<i>I</i> ² = 10.8%, <i>P</i> =	= .34)		0.42 (0.25-0.71)	27/1,309	57/1,163	100.00
	0.01	1 10	Note	: Box size is prop	ortional to stu	dy weight

Risk factors for reintubation

- A weak cough (cough peak expiratory flow rate ≤60 L/min)
- Frequent suctioning (every 1-2 hr, sputum volume >2.5 mL/hour)
- Glasgow Coma Score < 8
- A positive fluid balance during the 24 hours preceding extubation.
- Pneumonia as the reason for the initial intubation.
- Patients who are ≥65 years old with severe chronic cardiac or respiratory disease.
- A reduced or absent cuff leak, those with altered mental status.



Extubation management in the adult intensive ca

Reintubation

- Ranges from 4-33%
- <u>10-19%</u> may be clinically acceptable.
- Nosocomial pneumonia ¹ 8 folds
- Mortality 1 6-12 folds
- Up to 80% of intentionally self extubate do not require reintubation.

Liberation From Mechanical Ventilation in Critically Ill Adults



Executive Summary of an Official American College of Chest Physicians/American Thoracic Society Clinical Practice Guideline

- For patients at <u>high risk for extubation failure</u> who have been receiving mechanical ventilation for > 24 h and who have passed an SBT, we recommend extubation to preventive NIV
- Include those patients with <u>hypercapnia</u>, <u>COPD</u>, <u>congestive heart</u> failure, or other serious comorbidities.

(Strong Recommendation, Moderate Quality Evidence)

Official ERS/ATS clinical practice guidelines: noninvasive ventilation for acute respiratory failure

Bram Rochwerg 1, Laurent Brochard2,3, Mark W. Elliott4, Dean Hess5, Nicholas S. Hill6, Stefano Nava7 and Paolo Navalesi8 (members of the steering committee); Massimo Antonelli9, Jan Brozek1, Giorgio Conti9, Miquel Ferrer10, Kalpalatha Guntupalli11, Samir Jaber12, Sean Keenan13,14, Jordi Mancebo15, Sangeeta Mehta16 and Suhail Raoof17,18 (members of the task force)

TASK FORCE REPORT ERS/ATS GUIDELINES

Eur Respir J 2017; 50: 1602426

Should NIV be used to prevent respiratory failure post-extubation?

- NIV be used to prevent post-extubation respiratory failure in high-risk patients post-extubation. (Conditional recommendation, low certainty of evidence.)
- Patients at risk: <u>age >65years</u>, <u>underlying cardiac or respiratory disease</u>.
- NIV <u>should not</u> be used to prevent post-extubation respiratory failure in <u>non-high-risk patients</u>. (Conditional recommendation, very low certainty of evidence.)

Mortality

		NIV	,	Contr	ol		Risk Ratio	Risk Ratio
	Study or Subgroup	Events	Total	Events	Total	Weight	IV, Fixed, 95% CI	IV, Fixed, 95% CI
	1.2.1 Unselected Pati	ients						
	Su 2012	3	202	2	204		1.51 [0.26, 8.97]	
	Subtotal (95% CI)		202		204	14.7%	1.51 [0.26, 8.97]	
	Total events	3		2				
	Heterogeneity: Not ap	•					At r	isk patients
	Test for overall effect	Z = 0.46	$5 (\mathbf{P} = 0)$.65)				
	1.2.2 At Risk Patients	s						
	Ferrer 2006	2	79	12	83	21.6%	0.18 [0.04, 0.76]	
	Ferrer 2009	3	54	4	52	22.1%	0.72 [0.17, 3.07]	
	Nava 2005	3	48	9	49	29.9%	0.34 [0.10, 1.18]	
	Ornico 2013	1	20	7	18		0.13 [0.02, 0.95]	
	Subtotal (95% CI)		201		202	85.3%	0.31 [0.15, 0.64]	
	Total events	9		32				NUN/
	Heterogeneity: Chi ² =				$l^2 = 0\%$			NIV
	Test for overall effect	Z = 3.15	0 (P = 0)	.002)				
	Total (95% CI)		403		406	100.0%	0.39 [0.20, 0.76]	•
	Total events	12		34				
	Heterogeneity: Chi ² =	5.31, df	= 4 (P	= 0.26);	$l^2 = 25$	%		0.01 0.1 1 10 100
	Test for overall effect:							Favours [NIV] Favours [control]
	Test for subgroup diff	ferences:	Chi ² =	2.65, df	= 1 (P	= 0.10),	$^{2} = 62.3\%$	
							Eur Respir J 2017; 50: 1602426	

Re-intubation

	NIV	,	Contr	ol		Risk Ratio	Risk Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95% Cl	M-H, Fixed, 95% Cl
1.1.1 Unselected Pa	tients						
Jiang 1999	13	47	7	46	9.4%	1.82 [0.80, 4.14]	+
Su 2012	21	202	16	204	21.2%	1.33 [0.71, 2.47]	
Subtotal (95% CI)		249		250	30.7%	1.48 [0.90, 2.42]	►
Total events	34		23				
Heterogeneity: Chi ²				$l^2 = 0\%$		A	t risk patients
Test for overall effec	t: $Z = 1.54$	(P = 0)	0.12)				
1.1.2 At Risk Patien	ts						
Ferrer 2006	9	79	18	83	23.4%	0.53 [0.25, 1.10]	_ _
Ferrer 2009	6	54	10	52	13.6%	0.58 [0.23, 1.48]	
Kihlnani 2011	3	20	5	20	6.7%	0.60 [0.17, 2.18]	
Nava 2005	4	48	12	49	15.8%	0.34 [0.12, 0.98]	
Ornico 2013	1	20	7	18	9.8%	0.13 [0.02, 0.95]	
Subtotal (95% CI)		221		222	69.3%	0.44 [0.28, 0.70]	◆
Total events	23		52				
Heterogeneity: Chi ²	= 2.43, df	= 4 (P	= 0.66);	$l^2 = 0\%$			NIV
Test for overall effec	t: $Z = 3.50$	O(P = 0)	.0005)				
Total (95% CI)		470		472	100.0%	0.76 [0.55, 1.05]	
Total events	57		75				•
Heterogeneity: Chi ²		f = 6 (f		$ 1^2 = 5$	7%		
Test for overall effect				,			
Test for subgroup d				f = 1 (F)	e = 0.000	5), $l^2 = 91.9\%$	Favours [NIV] Favours [control]
							Eur Respir J 2017; 50: 1602426
Should NIV be used in the treatment of respiratory failure that develops post-extubation?

• NIV should not be used in the treatment of <u>patients with established</u> <u>post-extubation respiratory failure</u>. (Conditional recommendation, low certainty of evidence.)

Mortality and re-intubation



Eur Respir J 2017; 50: 1602426

Should NIV be used to facilitate weaning patients from invasive mechanical ventilation?

 NIV be used to facilitate weaning from mechanical ventilation in patients with hypercapnic respiratory failure.

(Conditional recommendation, moderate certainty of evidence)

• No recommendation for hypoxemic patients.

Mortality

	NIV		Contr			Risk Ratio	Risk Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95% CI	M-H, Fixed, 95% Cl
1.1.1 COPD							
Chen 2001	0	12	3	12	2.9%	0.14 [0.01, 2.50]	<
CRGNMV 2005	1	47	7	43	6.1%	0.13 [0.02, 1.02]	
Nava 1998	2	25	7	25	5.9%	0.29 [0.07, 1.24]	
Prasad 2009	5	15	9	15	7.5%	0.56 [0.24, 1.27]	
Rabie Agmy 2004	1	19	2	18	1.7%	0.47 [0.05, 4.78]	
Rabie Agmy 2012	7	134	26	130	22.1%	0.26 [0.12, 0.58]	
Wang 2004	1	14	2	14	1.7%	0.50 [0.05, 4.90]	· · · · · · · · · · · · · · · · · · ·
Zheng 2005	3	17	3	16	2.6%	0.94 [0.22, 4.00]	
Zou 2006	3	38	11	38	9.2%	0.27 [0.08, 0.90]	
Subtotal (95% CI)		321		311	59.8%	0.33 [0.21, 0.50]	•
Total events	23		70				
Heterogeneity: Chi ² =				$I^{2} = 0\%$			NIV
Test for overall effect:	Z = 5.10) (P < 0	.00001)				
1.1.2 Mixed Etiology							
Carron 2014	3	32	6	32	5.0%	0.50 [0.14, 1.83]	
Ferrer 2003	6	21	12	22	9.8%	0.52 [0.24, 1.14]	
Girault 1999	0	17	2	16	2.2%	0.19 [0.01, 3.66]	· · · ·
Girault 2011	16	69	9	69	7.5%	1.78 [0.84, 3.75]	
Hill 2000	1	12		0			
1111 2000	-	12	1	9	1.0%	0.75 [0.05, 10.44]	
Tawfeek 2012	2	21	1 6	21	1.0%	0.75 [0.05, 10.44] 0.33 [0.08, 1.47]	
	2 9						
Tawfeek 2012 Trevisan 2008 Vaschetto 2012	2	21 28 10	6	21 37 10	5.0% 7.2% 2.5%	0.33 [0.08, 1.47] 1.19 [0.56, 2.53] 0.67 [0.14, 3.17]	
Tawfeek 2012 Trevisan 2008	2 9	21 28	6 10	21 37	5.0% 7.2%	0.33 [0.08, 1.47] 1.19 [0.56, 2.53]	
Tawfeek 2012 Trevisan 2008 Vaschetto 2012 Subtotal (95% CI) Total events	2 9 2 39	21 28 10 210	6 10 3 49	21 37 10 216	5.0% 7.2% 2.5% 40.2%	0.33 [0.08, 1.47] 1.19 [0.56, 2.53] 0.67 [0.14, 3.17]	
Tawfeek 2012 Trevisan 2008 Vaschetto 2012 Subtotal (95% Cl) Total events Heterogeneity: Chi ² =	2 9 2 39 9.29, df	21 28 10 210 = 7 (P	6 10 3 49 = 0.23);	21 37 10 216	5.0% 7.2% 2.5% 40.2%	0.33 [0.08, 1.47] 1.19 [0.56, 2.53] 0.67 [0.14, 3.17]	
Tawfeek 2012 Trevisan 2008 Vaschetto 2012 Subtotal (95% CI) Total events	2 9 2 39 9.29, df	21 28 10 210 = 7 (P	6 10 3 49 = 0.23);	21 37 10 216	5.0% 7.2% 2.5% 40.2%	0.33 [0.08, 1.47] 1.19 [0.56, 2.53] 0.67 [0.14, 3.17]	
Tawfeek 2012 Trevisan 2008 Vaschetto 2012 Subtotal (95% Cl) Total events Heterogeneity: Chi ² =	2 9 2 39 9.29, df	21 28 10 210 = 7 (P	6 10 3 49 = 0.23);	21 37 10 216 I ² = 25	5.0% 7.2% 2.5% 40.2%	0.33 [0.08, 1.47] 1.19 [0.56, 2.53] 0.67 [0.14, 3.17]	
Tawfeek 2012 Trevisan 2008 Vaschetto 2012 Subtotal (95% Cl) Total events Heterogeneity: Chi ² = Test for overall effect:	2 9 2 39 9.29, df	21 28 10 210 = 7 (P = 0) 3 (P = 0)	6 10 3 49 = 0.23);	21 37 10 216 I ² = 25	5.0% 7.2% 2.5% 40.2%	0.33 [0.08, 1.47] 1.19 [0.56, 2.53] 0.67 [0.14, 3.17] 0.85 [0.59, 1.22]	
Tawfeek 2012 Trevisan 2008 Vaschetto 2012 Subtotal (95% CI) Total events Heterogeneity: Chi ² = Test for overall effect: Total (95% CI) Total events	2 9 2 9.29, df Z = 0.88	21 28 10 210 = 7 (P = 0) 531	6 10 3 49 = 0.23); 0.38) 119	21 37 10 216 I ² = 25 527	5.0% 7.2% 2.5% 40.2% %	0.33 [0.08, 1.47] 1.19 [0.56, 2.53] 0.67 [0.14, 3.17] 0.85 [0.59, 1.22]	
Tawfeek 2012 Trevisan 2008 Vaschetto 2012 Subtotal (95% Cl) Total events Heterogeneity: Chi ² = Test for overall effect: Total (95% Cl)	2 9 2 9.29, df Z = 0.88 62 23.53, d	21 28 10 210 = 7 (P = 0) 531 f = 16	6 10 3 49 = 0.23); 0.38) 119 (P = 0.10	21 37 10 216 I ² = 25 527	5.0% 7.2% 2.5% 40.2% %	0.33 [0.08, 1.47] 1.19 [0.56, 2.53] 0.67 [0.14, 3.17] 0.85 [0.59, 1.22]	0.01 0.1 1 10 100 Favours [NIV] Favours [control

COPD

Weaning failure

		NIN	/	Contr	rol		Risk Ratio	Risk Ratio
	Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95% CI	M–H, Fixed, 95% Cl
	1.2.1 COPD							
	Nava 1998	3	25	8	25	6.7%	0.38 [0.11, 1.25]	
	Rabie Agmy 2004	4	19	6	18	5.1%	0.63 [0.21, 1.88]	
COPD	Rabie Agmy 2012	28	134	52	130	43.9%	0.52 [0.35, 0.77]	-
	Subtotal (95% CI)		178		173	55.7%	0.51 [0.36, 0.73]	◆
	Total events	35		66				
	Heterogeneity: Chi ² =	NIV						
	Test for overall effect	: Z = 3.7	O(P = 0)	0.0002)				
	1.2.2 Mixed Etiology							
	Carron 2014	5	32	11	32	9.2%	0.45 [0.18, 1.16]	
	Girault 1999	4	17	4	16	3.4%	0.94 [0.28, 3.14]	
	Girault 2011	23	69	22	69	18.3%	1.05 [0.65, 1.69]	
	Hill 2000	4	12	1	9	1.0%	3.00 [0.40, 22.47]	
	Tawfeek 2012	3	21	10	21	8.3%	0.30 [0.10, 0.94]	
	Vaschetto 2012	1	10	5	10	4.2%	0.20 [0.03, 1.42]	
	Subtotal (95% CI)		161		157	44.3%	0.74 [0.52, 1.05]	•
	Total events	40		53				
	Heterogeneity: $Chi^2 = 9.17$, $df = 5$ (P = 0.10); $I^2 = 45\%$ Test for overall effect: Z = 1.68 (P = 0.09)							
	Total (95% CI)		339		330	100.0%	0.61 [0.48, 0.79]	•
	Total events	75		119				
	Heterogeneity: Chi ² =	12.07, d	f = 8 (I	P = 0.15)	; I ² = 3	4%		0.01 0.1 10 100
	Test for overall effect	Z = 3.8	5 (P = 0)	0.0001)				0.01 0.1 I 10 100 Favours [NIV] Favours [control]
	Test for subgroup dif	ferences:	Chi ² =	2.00, df	= 1 (P	= 0.16), I	² = 50.0%	Favours [iviv] Favours [control]

SYSTEMATIC REVIEW

Non-invasive ventilation as a strategy for weaning from invasive mechanical ventilation: a systematic review and Bayesian meta-analysis

CrossMark

University of Warwick, Coventry, UK

Joyce Yeung, Keith Couper, Elizabeth G. Ryan, Simon Gates, Nick Hart and Gavin D. Perkins

- The use of NIV in weaning from mechanical ventilation decreases hospital mortality, the incidence of VAP and ICU stay.
- NIV as a weaning strategy appears to be most beneficial in COPD.

Intensive Care Med (2018) 44:2192–2204 https://doi.org/10.1007/s00134-018-5434-z

Hospital mortality rate for NIV and invasive weaning

COPD	n/N (Intervention vs Control)	OR < 1	Posterior Mean Odds Ratio (95% HDI)	Observed Odds Ratio (95% CI)
Sirault et al. (1999) Chen et al. (2001) Vang et al. (2004) Vang et al. (2005) Cheng et al. (2005) Cou et al. (2006) Sirault et al. (2011) Mohamed and Ibrahim (2012) Rong (2012)	0 / 17 vs 2 / 16 0 / 12 vs 3 / 12 1 / 14 vs 2 / 14 1 / 47 vs 7 / 43 3 / 17 vs 3 / 16 3 / 38 vs 11 / 38 16 / 69 vs 9 / 69 1 / 15 vs 3 / 15 2 / 33 vs 8 / 31		0.58 (0.05, 1.28) 0.55 (0.03, 1.19) 0.66 (0.09, 1.41) 0.46 (0.06, 0.93) 0.77 (0.14, 1.59) 0.45 (0.09, 0.85) 1.30 (0.46, 2.50) 0.59 (0.07, 1.23) 0.46 (0.08, 0.92)	0.17 (0.01, 3.73) 0.11 (0.00, 2.36) 0.56 (0.06, 4.84) 0.16 (0.03, 0.96) 0.93 (0.18, 4.88) 0.24 (0.06, 0.86) 1.96 (0.82, 4.73) 0.37 (0.05, 2.89) 0.22 (0.05, 0.99)
Anshra et al. (2014) Pooled COPD	2 / 25 vs 8 / 25 29 / 287 vs 56 / 279		0.43 (0.03, 0.02) 0.43 (0.13, 0.81)	0.22 (0.05, 1.02)
Mixed population Trevisan et al. (2008) Charra et al. (2009) Vaschetto et al. (2012) Carron et al. (2014) Wang et al. (2014) Perkins et al. (2018) Pooled mixed population	9 / 28 vs 10 / 37 0 / 12 vs 0 / 12 2 / 10 vs 3 / 10 3 / 32 vs 6 / 32 2 / 26 vs 5 / 27 35 / 182 vs 36 / 182 51 / 290 vs 60 / 300		0.95 (0.32, 1.85) 0.74 (0.05, 1.71) 0.69 (0.11, 1.44) 0.60 (0.13, 1.16) 0.59 (0.11, 1.16) 0.89 (0.50, 1.35) 0.88 (0.25, 1.48)	1.28 (0.45, 3.65) 1.00 (0.02, 54.47 0.63 (0.09, 4.20) 0.48 (0.12, 1.96) 0.42 (0.08, 2.07) 0.97 (0.58, 1.62)
Overall pooled	80 / 577 vs 116 / 579	0.02 0.05 0.1 0.2 0.5 Odds Ratio	0.58 (0.29, 0.89)	

Fig. 3 Forest plot comparing hospital mortality rates for NIV and invasive weaning, by patient population (COPD vs. mixed ICU population). The estimated odds ratio (OR) from the posterior distribution for each study is shown as a circle, with 95% HDI represented by horizontal lines. The observed OR are given by crosses. The pooled OR estimates (and 95% HDI) are also displayed as the last row for each patient population, and the overall pooled estimate for all studies is displayed as the last row. An OR < 1 means that the intervention is superior

High flow nasal cannula (HFNC)

- Some high risk patients should receive HFNC for 24-48 hr.
- Patients with severe hypoxemic respiratory failure or high O₂ requirement.
- Provide small amount of PEEP.
- HFNC establishes 1cmH₂O for 10L/min flow delivered.



Egan's fundamentals of respiratory care 12th edition

Extubation management in the adult intensive ca

JAMA | Original Investigation | CARING FOR THE CRITICALLY ILL PATIENT

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

- In mechanically ventilated patients at high risk of extubation failure.
- The use of HFNC with NIV immediately after extubation significantly decreased the risk of reintubation compared with HFNC alone.

Table 2. Primary, Secondary, and Explore	ratory Outcomes
--	-----------------

	No. (%) High-Flow Nasal Oxygen Alone (n = 302)	High-Flow Nasal Oxygen With NIV (n = 339)	Absolute Difference, % (95% CI)	P Value
Primary Outcome				
Reintubation at day 7	55 (18)	40 (12)	-6.4 (-12.0 to -0.9)	.02
Secondary Outcomes				
Postextubation respiratory failure at day 7	88 (29)	70 (21)	-8.5 (-15.2 to -1.8)	.01
Reintubation				
At 48 h	36 (12)	24 (7)	-4.8 (-9.6 to -0.3)	.04
At 72 h	47 (16)	30 (9)	-6.7 (-11.9 to -1.7)	.009
Up until ICU discharge	59 (20)	41 (12)	−7.4 (−13.2 to −1.8)	.009
Length of stay, median (IQR), days				
In ICU	11 (7 to 19)	12 (7 to 19)	0.5 (-1.6 to 2.6)	.55
In hospital	23 (15 to 39)	25 (15 to 42)	2.3 (-1.4 to 6.1)	.31
Mortality				
In ICU	26 (9)	21 (6)	-2.4 (-6.7 to 1.7)	.25
In hospital	46 (15)	54 (16)	0.7 (-5.0 to 6.3)	.80
At day 28	33 (11)	39 (12)	0.6 (-4.4 to 5.5)	.82
At day 90	65 (21)	62 (18)	-3.2 (-9.5 to 2.9)	.30
Exploratory Outcomes				
Patients meeting reintubation criteria during ICU stay	65 (22)	49 (14)	-7.1 (-13.1 to -1.1)	.02
Mortality or reintubation in ICU	64 (21)	51 (15)	-6.2 (-12.2 to -0.2)	.04
Mortality of reintubated patients	21/59 (36)	11/41 (27)	-8.8 (-25.7 to 9.9)	.35

Reintubation rate

Figure 2. Kaplan-Meier Analysis of Time From Extubation to Reintubation for the Overall Study Population



The median observation time was 7 days (interquartile range, 7-7) in both treatment groups.

JAMA. 2019;322(15):1465-1475

Figure 3. Kaplan-Meier Analysis of Time From Extubation to Reintubation According to Predefined Strata



Results in hypercapnic patients with arterial partial pressure of carbon dioxide (Paco₂) greater than 45 mm Hg (A) and in nonhypercapnic patients with Paco₂ of 45 mm Hg or less (B) are shown. The median observation time was 7 days (interquartile range, 7-7) in both treatment groups.

JAMA. 2019;322(15):1465-1475

The morbid obese patients

- BMI > 30-35
- CPAP should be used during SBT and post extubation.
- The larger the patient the greater the likelihood of atelectasis.
- Majority of these patients have sleep apnea
- High level of PEEP ($\geq 10 \text{cmH}_2\text{O}$) is required to stabilize the lung.





Weaning patients with obesity from ventilatory support

Robert M. Kacmarek^{a,b,c}, Hatus V. Wanderley^{a,b,c}, Jesús Villar^{d,e} and Lorenzo Berra^{a,c}

- <u>Obesity</u> greatly alters the respiratory system mechanics causing atelectasis and prolonged duration of mechanical ventilation.
- WOB is markedly increased because of their negative transpulmonary pressure.
- Patients should immediately be transitioned to Mask CPAP ≥10 cmH2O or equal to the applied PEEP during weaning or to their ordered CPAP setting for sleep apnea.



Comparison of the FRC

Lung recruitability



To maintain same FRC, the airway pressure in the subject with obesity needs to increase of about 20 cmH₂O above atmospheric pressure.

When lungs are recruitable, pressure reached during exhalation shows higher volumes than during inspiration, advocating for decremental PEEP trial rather than incremental PEEP trial.

PEEP for failed and success SBT



Take home message

- Discontinuing mechanical ventilation is a three-step process that consists of readiness testing, weaning and extubation.
- Verification the problems leading to mechanical ventilation have been resolved is the first step in successfully liberating a patient from ventilatory support.
- Evaluation of appropriate criteria and the use of <u>therapist-driven</u> protocols or nurse-directed protocols can facilitate the process.
- RSBI is the preferred indicator.

Take home message

- SBT and PSV result in faster discontinuation.
- Diaphragm ultrasound may be useful in identifying readiness.
- Carefully monitoring during weaning.
- Review the common causes of weaning failure.
- Assess for the ability to maintain and protect airway and the presence of upper airway edema before extubation.
- CPAP should be applied during SBT in morbid obese patients.

Thank you for your listening