

Dünya genelinde yoğun bakıma yatışların önemli bir bölümü (%20-40)* LUNG-SAFE

Değişken solunum dinamikleri → P-SILI artış

Hem AHRF hem ARDS açısından aynı spektrumun parçasıdır.

RESEARCH

Open Access

Causes and characteristics of death in patients with acute hypoxemic respiratory failure and acute respiratory distress syndrome: a retrospective cohort study

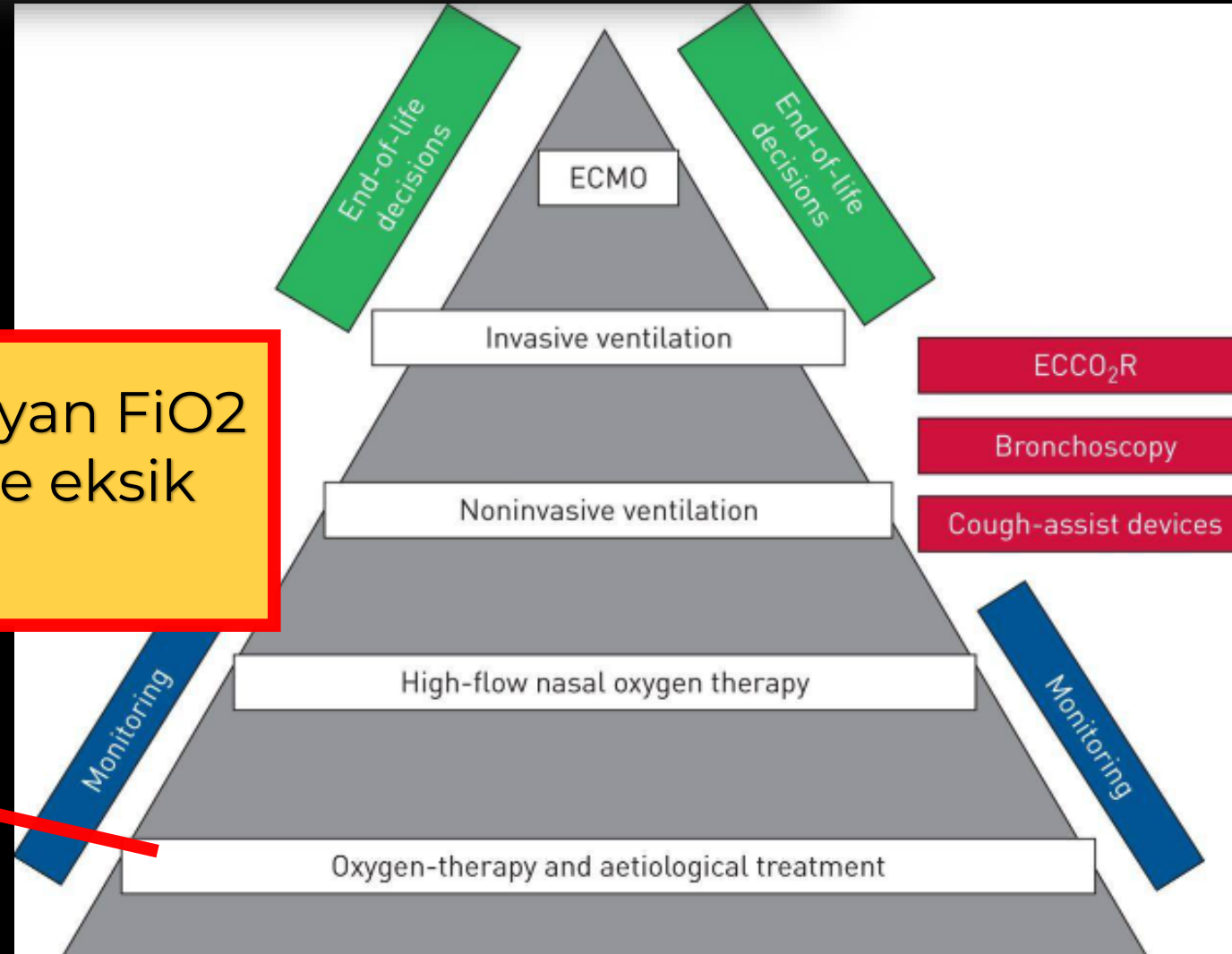


**Sepsis (%26),
Pulmoner disfonksiyon (%22)
Nörolojik disfonksiyon (%19)**

ARDS ve non ARDS arasında fark yok

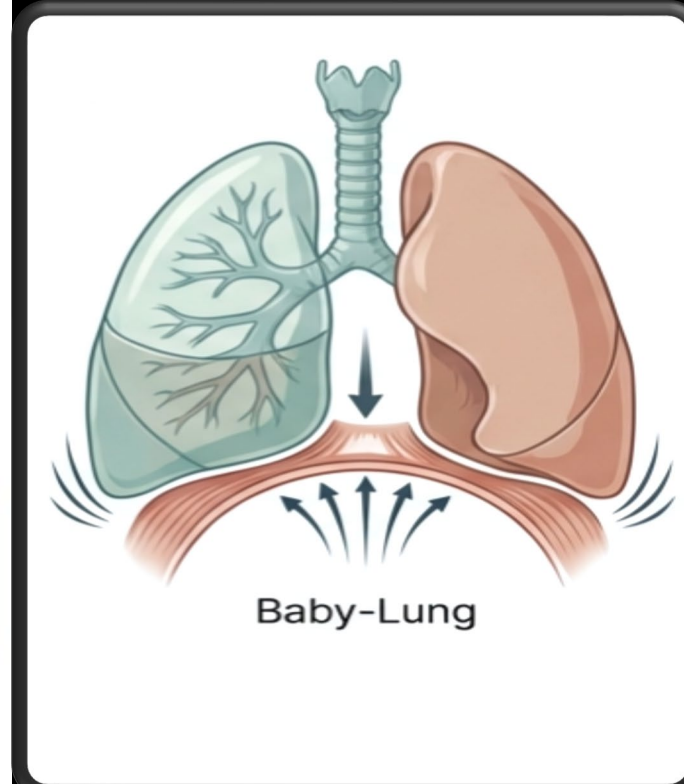
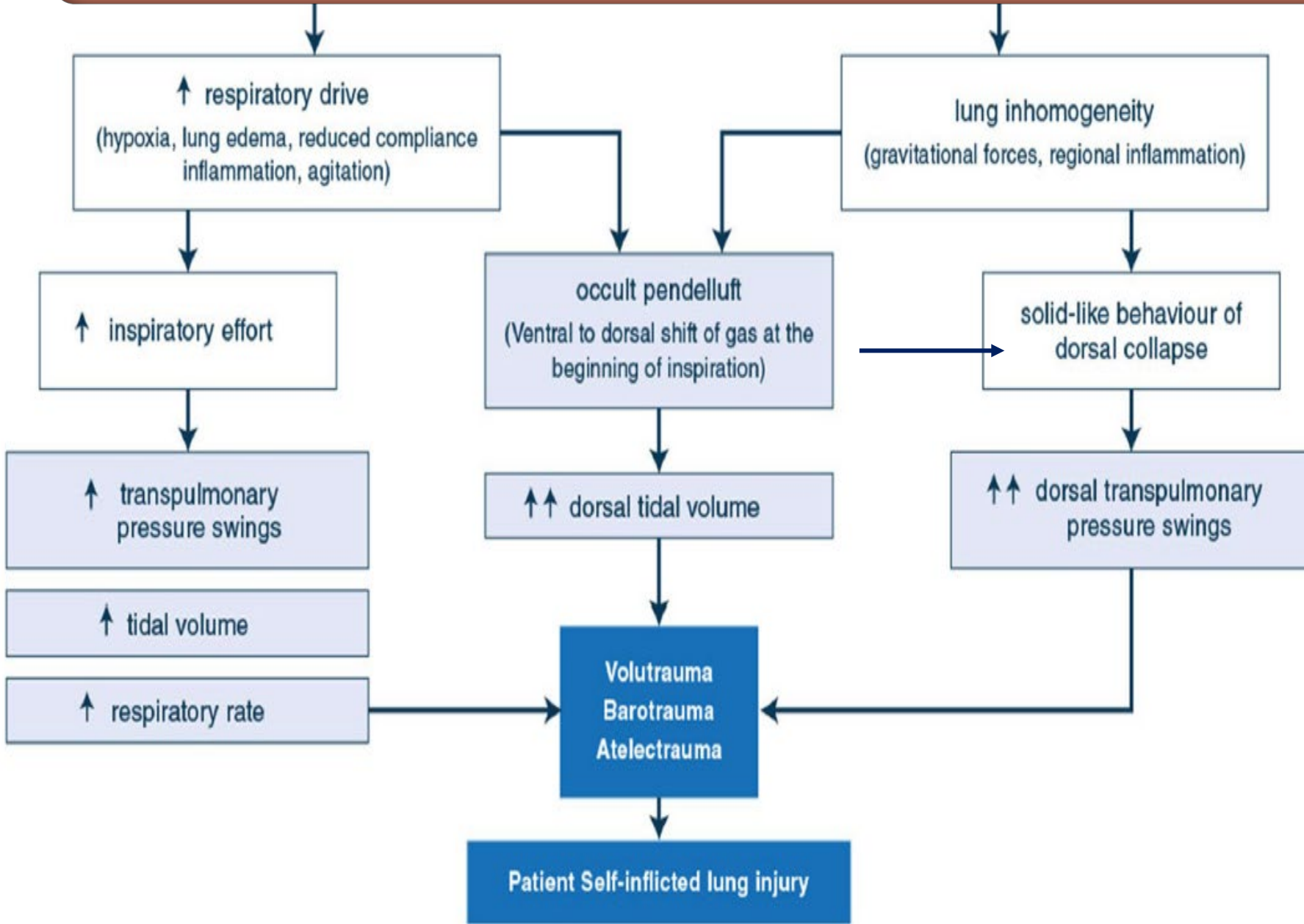
Highlights in acute respiratory failure

Raffaele Scala¹ and Leo Heunks²

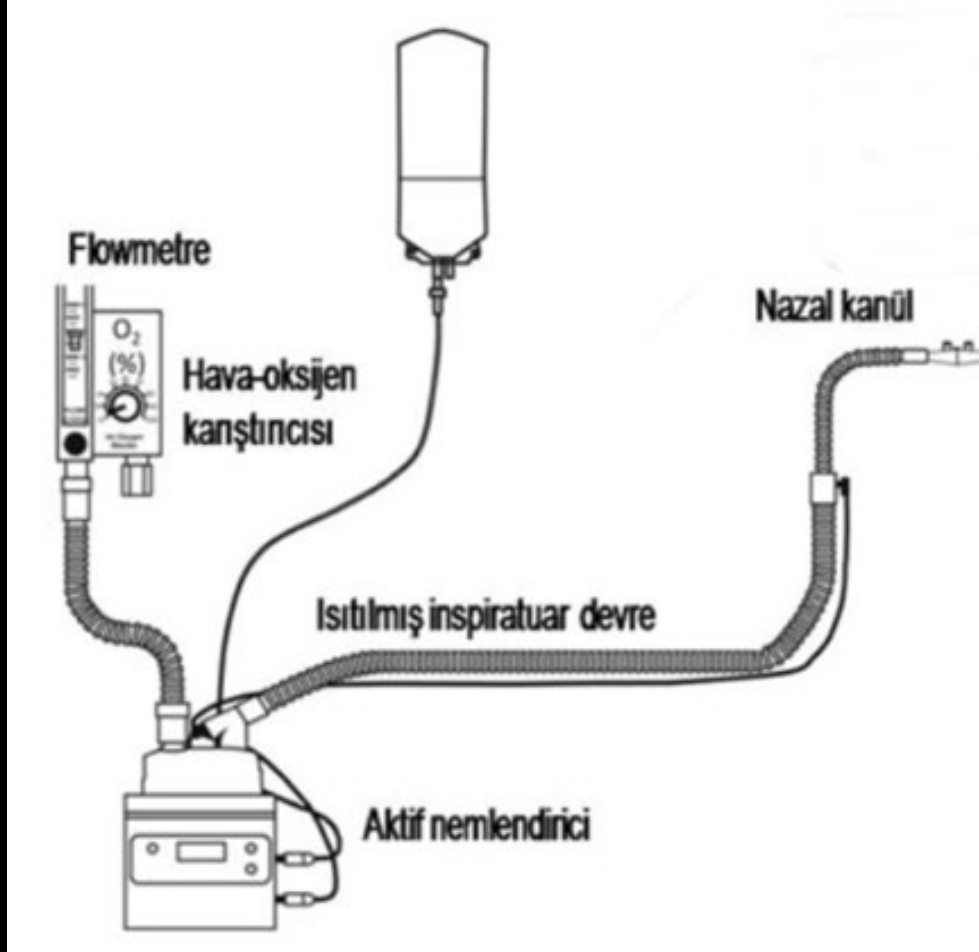


Güvenilir olmayan FiO₂
Nemlendirme eksik
P-SILI

Spontan Solunum: İki Ucu Keskin Bıçak



Yüksek Akışlı Nazal Oksijen



PEEP

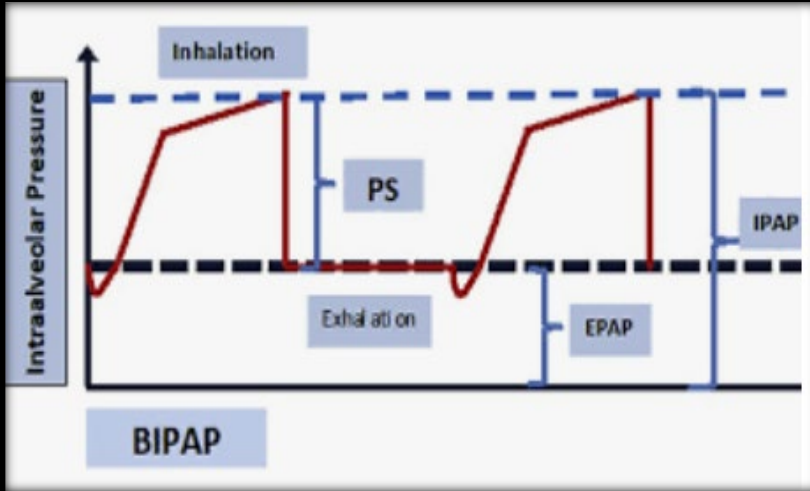
CO2 washout

Stabil FiO2

Nemlendirme

P-SILI azalması

NIV



CPAP ve BPAP

Maske / Helmet

Daha iyi PEEP

P-SILI riski

Positive Pressure

↑ ITP

↑ FRC

↓ Pre-load

↓ Venous return

↓ LV afterload

↓ PTM

↑ PaO₂

↓ WOB

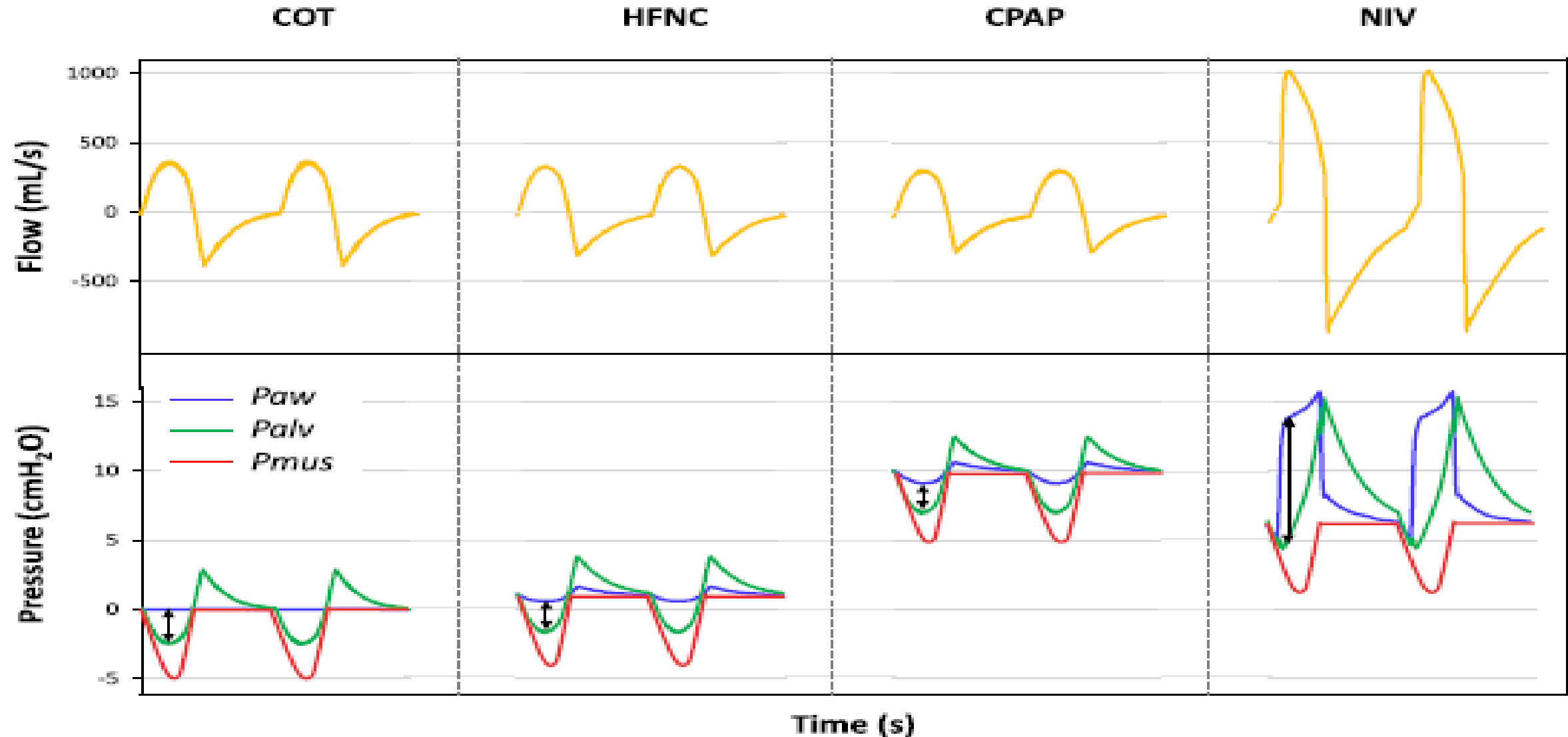
RESEARCH

Open Access

Influence of different noninvasive oxygenation support devices on tidal volume



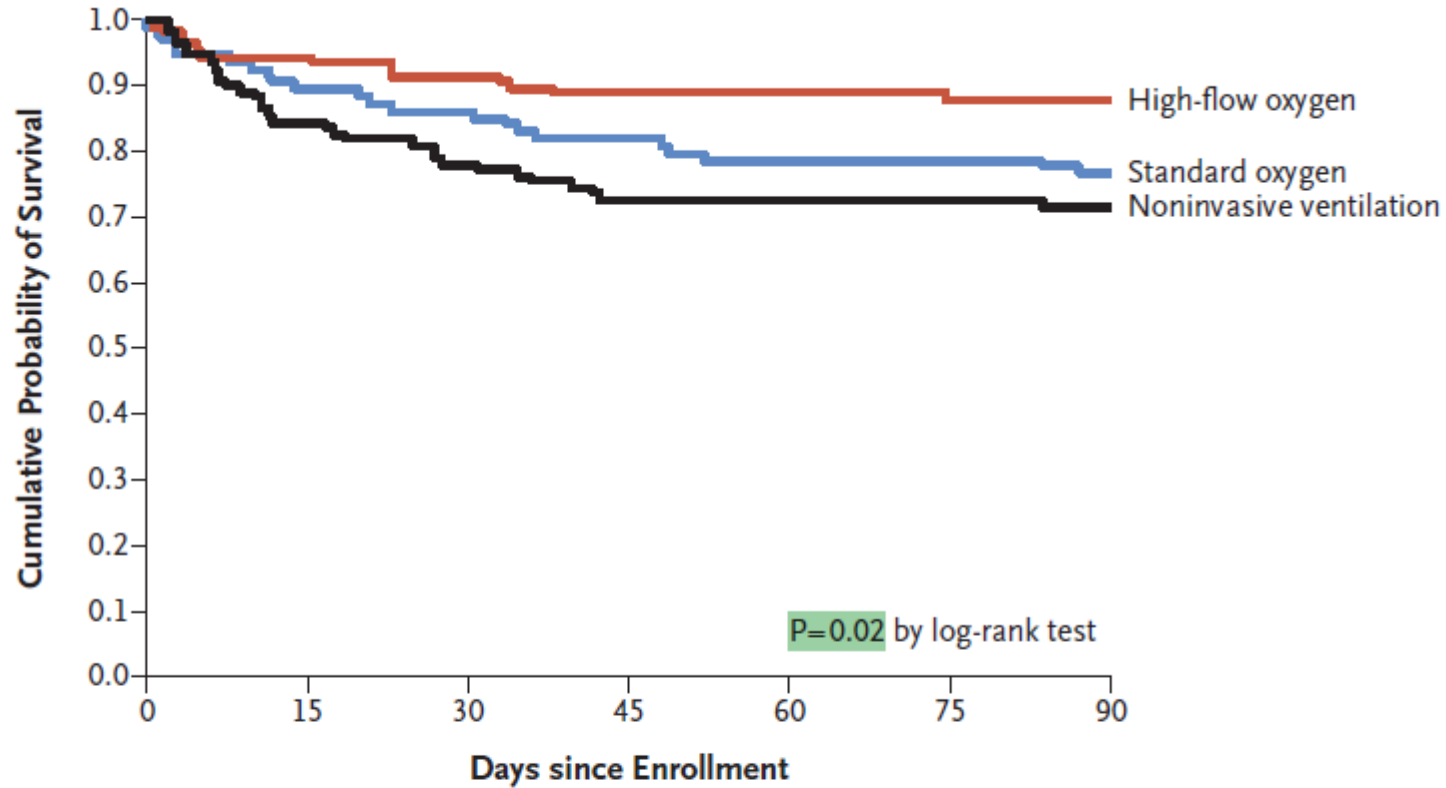
Inspiratuar basınç gradyanı üzerine etki:
en yüksek NIV



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

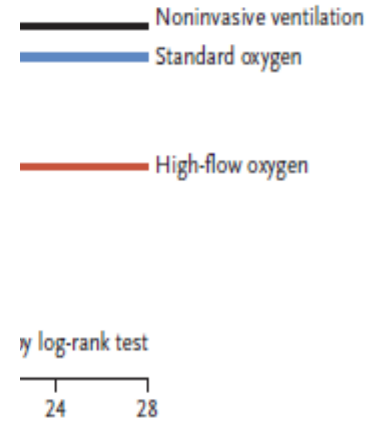
A Overall Population

Cumulative Incidence of Intubation



No. at Risk

No. at Risk	0	15	30	45	60	75	90
High-flow oxygen	106	100	97	94	94	93	93
Standard oxygen	94	84	81	77	74	73	72
Noninvasive ventilation	110	93	86	80	79	78	77



53	53
33	33
32	32

Figure 3. Kaplan–Meier Plot of the Probability of Survival from Randomization to Day 90.

HFNO / COT ?

SYSTEMATIC REVIEW

High flow nasal cannula compared with conventional oxygen therapy for acute hypoxemic respiratory failure: a systematic review and meta-analysis



- **9 RKCÇ, n= 2093**
- **Mortaliteyi azaltmaz**
- **Entübasyon ihtiyacını azaltır**

ventilation in either group) favouring HFNC-treated patients (RR 0.71, 95% CI 0.51–0.98), although certainty in both outcomes was low due to imprecision and issues related to risk of bias. HFNC had no effect on intensive care unit length of stay (mean difference [MD] 1.38 days more, 95% CI 0.90 days fewer to 3.66 days more, low certainty), hospital length of stay (MD 0.85 days fewer, 95% CI 2.07 days fewer to 0.37 days more, moderate certainty), patient reported comfort (SMD 0.12 lower, 95% CI 0.61 lower to 0.37 higher, very low certainty) or patient reported dyspnea (standardized mean difference [SMD] 0.16 lower, 95% CI 1.10 lower to 1.42 higher, low certainty). Complications of treatment were variably reported amongst included studies, but little harm was associated with HFNC use.

ERS clinical practice guidelines: high-flow nasal cannula in acute respiratory failure

Eur Respir J 2022; 59: 2101574

COT yerine HFNO tedavisi: koşullu öneri, orta düzeyde kanıt

Entübasyon riskini azaltabilir (0.89)

NIV'e eskalasyon gereksinimi HFNO ile biraz daha düşük

28 ve 90 günlük mortalitede fark yok

CONFERENCE REPORTS AND EXPERT PANEL

ESICM guidelines on acute respiratory distress syndrome: definition, phenotyping and respiratory support strategies



HFNO, COT'a göre entübasyon oranlarını düşürüyor (güçlü öneri)

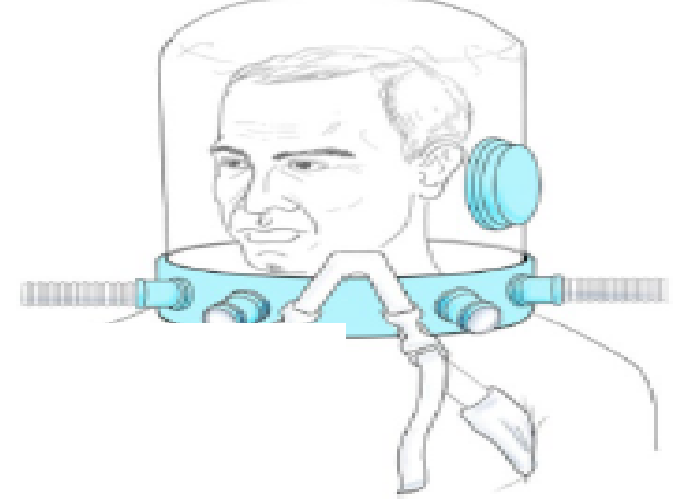
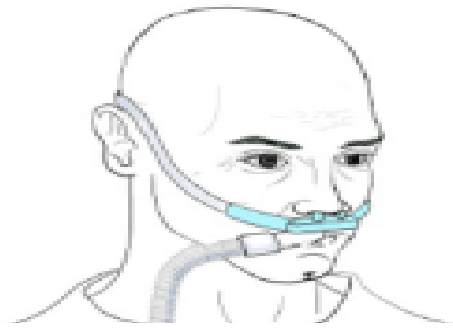
Mortalitede fark yok

Recommendation 3.1

We **recommend** that non-mechanically ventilated patients with AHRF not due to cardiogenic pulmonary edema or acute exacerbation of COPD receive HFNO as compared to conventional oxygen therapy to reduce the risk of intubation
Strong recommendation; moderate level of evidence in favor

We are **unable to make a recommendation** for or against the use of HFNO over conventional oxygen therapy to reduce mortality
No recommendation; high level of evidence of no effect

This recommendation applies also to AHRF from COVID-19
Strong recommendation; low level of evidence in favor for intubation and no recommendation; moderate level of evidence of no effect for mortality, for indirectness.



JAMA. 2021 Mar 25;325(17):1-13. doi: [10.1001/jama.2021.4682](https://doi.org/10.1001/jama.2021.4682)

Effect of Helmet Noninvasive Ventilation vs High-Flow Nasal Oxygen on Days Free of Respiratory Support in Patients With COVID-19 and Moderate to Severe Hypoxemic Respiratory Failure

The HENIVOT Randomized Clinical Trial

JAMA, 315(22), 2435-2441

Pitfalls

- Small amount of PEEP delivered

Daha düşük entübasyon oranı + ancak 28-gün ventilatörsüz gün sayısında fark yok

volume

- Poor tolerability: need for treatment interruptions

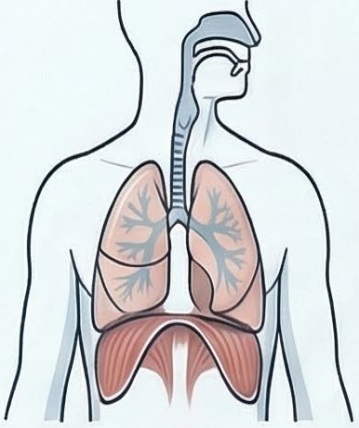
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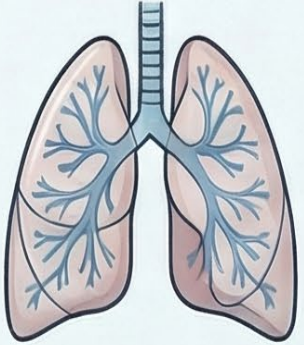
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Kask Arayüzünün Temel Avantajları



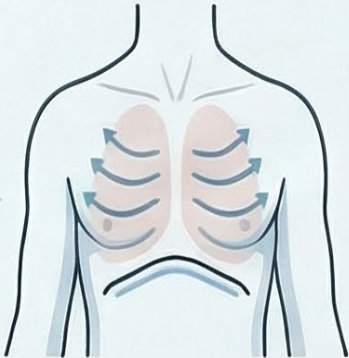
Kas İş Yükünün Azaltılması

Spesifik ayarlar sayesinde solunum kasları üzerindeki yükü belirgin şekilde hafifletir.



Homojen Ventilasyon ve Yüksek PEEP

Akciğerde eşit hava dağılımını sağlar ve 10-12+ cm H₂O PEEP seviyelerinde yüksek konfor sunar.



Spontan Eforun Korunması

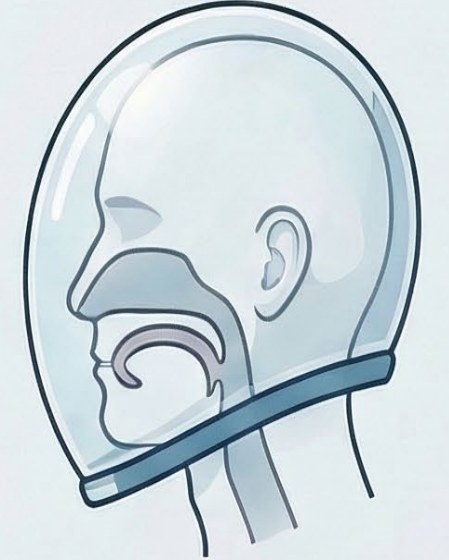
Yüksek PEEP uygulamasıyla hastanın spontan solunum çabasının akciğer üzerindeki yıkıcı etkilerini önler.



Performans Karşılaştırması: Kask vs. Yüz Maskesi

Basınç Desteği (PS) Etkinliği

Kask arayüzünde basınç dağılımı nedeniyle, aynı PS desteği maskeye göre daha düşük iş yükü azalması sağlar.



10 cm H₂O Basınç Desteği (PS) Altında Solunum İşindeki Azalma Oranları

Arayüz Tipi	Solunum İşindeki Azalma (%)	Temel Neden
Yüz Maskesi	50%	Doğrudan basınç iletimi
Kask (Helmet)	30%	Basınç dağılımı (dispersiyon)

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Helmet arayüzün yüz maskesine üstünlüğü konusunda öneri yapılamaz. (çok düşük düzey kanıt)

Recommendation 4.2

We are **unable to make a recommendation** for or against the use of helmet interface for CPAP/NIV as compared to face mask to prevent intubation or reduce mortality in patients with acute hypoxemic respiratory failure.
No recommendation; very low level of evidence in favor.

ERS clinical practice guidelines: high-flow nasal cannula in acute respiratory failure

Eur Respir J 2022; 59: 2101574

NIV yerine HFNO tedavisi: kořullu öneri, çok düşük kanıt

Kanıtlarda:



& aralıklı uygulama

Hospitalizasyonda fark yok

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HFNO ile CPAP/NIV arasında lehine ya da aleyhine öneri yapılamaz (mortalite: orta düzey kanıt, entübasyon: düşük düzey kanıt)

COVID-19'da entübasyonu azaltmak için CPAP/NIV düşünülebilir (Zayıf öneri; yüksek düzey kanıt)

Recommendation 3.2

We are **unable to make a recommendation** for or against the use of HFNO compared to continuous positive airway pressure (CPAP)/NIV to reduce intubation or mortality in the treatment of unselected patients with acute hypoxemic respiratory failure not due to cardiogenic pulmonary edema or acute exacerbation of COPD.

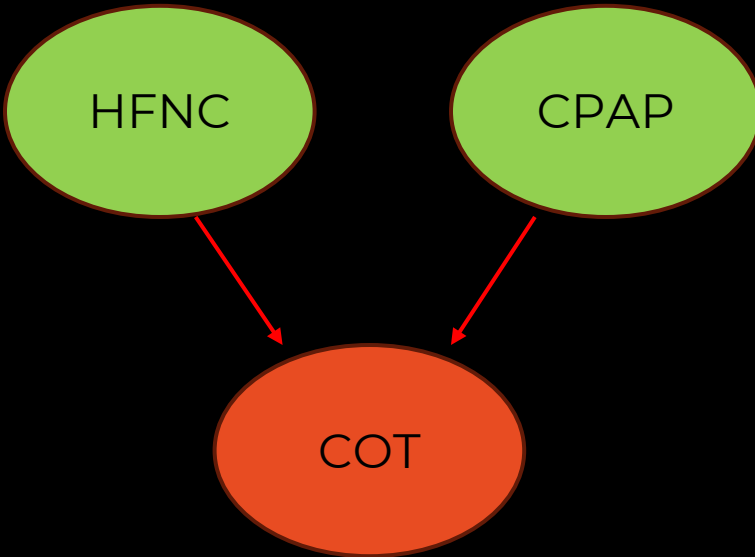
No recommendation; moderate level of evidence for mortality, low level of evidence for intubation, not in favor nor against.

We **suggest** that CPAP/NIV can be considered instead of HFNO for the treatment of AHRF due to COVID-19 to reduce the risk of intubation (*weak recommendation, high level of evidence*), but **no recommendation** can be made for whether CPAP/NIV can decrease mortality compared to HFNO in COVID-19.

No recommendation; high level of evidence of no effect.

Effect of Noninvasive Respiratory Strategies on Intubation or Mortality Among Patients With Acute Hypoxemic Respiratory Failure and COVID-19

The RECOVERY-RS Randomized Clinical Trial



n=1273

CPAP; entübasyon /ölümden oluşan kompozit sonlanımı yaklaşık %8 mutlak risk azalması ile azaltmıştır; bu etki esas olarak entübasyon oranındaki düşüştten kaynaklanmaktadır

HFNO'de fark yok

CONFERENCE REPORTS AND EXPERT PANEL

ESICM guidelines on acute respiratory distress syndrome: definition, phenotyping and respiratory support strategies



Recommendation 4.1

We are **unable to make a recommendation** for or against the use of CPAP/NIV compared to conventional oxygen therapy for the treatment of AHRF (not related to cardiogenic pulmonary edema or acute exacerbation of COPD) to reduce mortality or to prevent intubation.

No recommendation; high level of evidence for mortality, moderate level of evidence for intubation.

We **suggest** the use of CPAP over conventional oxygen therapy to reduce the risk of intubation in patients with acute hypoxemic respiratory failure due to COVID-19.

Weak recommendation; low level of evidence in favor.

In this population, we are **unable to make a recommendation** for or against the use of CPAP over conventional oxygen therapy to reduce mortality.

No recommendation; moderate level of evidence of no effect.

Mortaliteyi azaltmak veya entübasyonu önlemek için CPAP/NIV'nin COT'a üstünlüğü konusunda öneri yapılamaz.

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ESICM guidelines on acute respiratory distress syndrome: definition, phenotyping and respiratory support strategies



**AHRF tedavisinde NIV'nin CPAP'a üstün ya da daha kötü olduğuna dair öneri yapılamaz.
(kanıt yok)**

Recommendation 4.3

We are **unable to make a recommendation** for or against the use of NIV compared to CPAP for the treatment of AHRF.

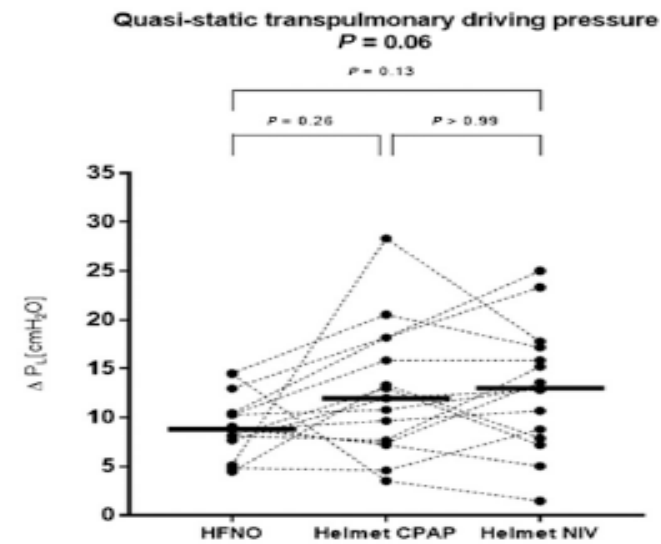
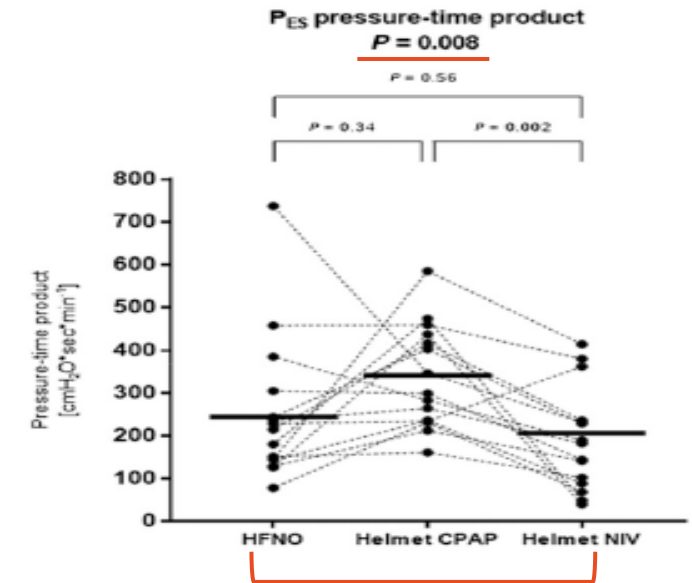
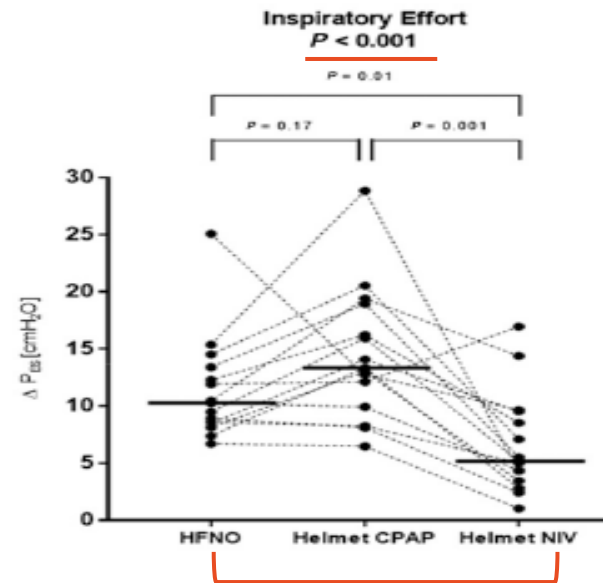
No recommendation; no evidence.

Respective Effects of Helmet Pressure Support, Continuous Positive Airway Pressure, and Nasal High-Flow in Hypoxemic Respiratory Failure

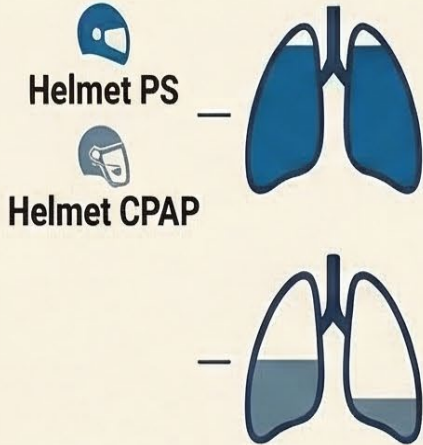
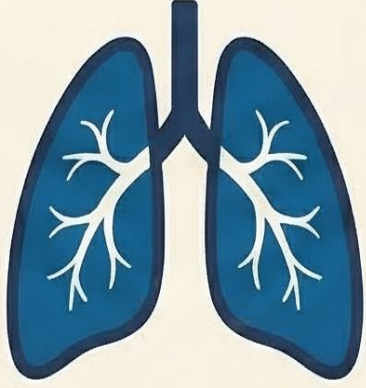
A Randomized Crossover Clinical Trial

HFNO- Helmet CPAP-
Helmet PS

N=15



Alveoler Rekrütmanda Helmet Üstünlüğü



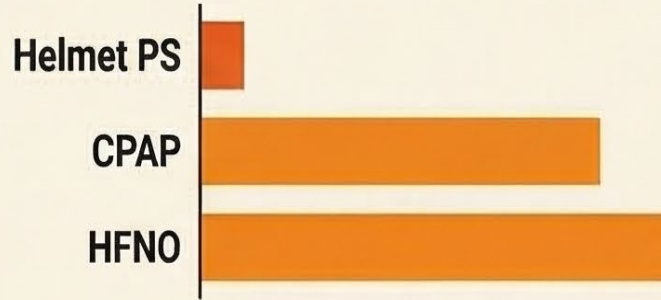
Hem Helmet CPAP hem de Helmet PS, HFNO'ya kıyasla EELV'de anlamlı artış sağlayarak daha etkin rekrütman gerçekleştirir.

En Düşük Solunum İş Yükü

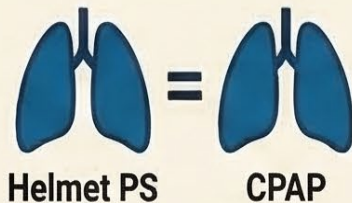


EN DÜŞÜK

Hastanın inspiratuar eforu ve toplam solunum iş yükü, Helmet PS modunda en düşük seviyeye iner.



PS vs. CPAP: Rekrütman Benzerliği



Akciğer rekrütmanı açısından Helmet PS ve CPAP arasında belirgin bir fark saptanmamıştır; fark efor yönetimindedir.

P-SILI Riskine Karşı En Koruyucu Strateji



«NIV herkeste kesin olarak daha güvenlidir» şeklinde düz bir sonuç çıkarılamaz. «hasta fenotipi»ne bağlı

Helmet PS, hem akciğer rekrütmanı sağlar hem de inspiratuar eforu en aza indirerek **P-SILI riskine karşı en güçlü korumayı sunar.**

High-Flow Nasal Oxygen vs Noninvasive Ventilation in Patients With Acute Respiratory Failure

JAMA

The RENOVATE Randomized Clinical Trial 2025 Mar 11;333(10):875-890

Noninferiority tasarımlı, RKÇ

Brezilya- 33 merkez ; n=1766

**Nonimmünsüprese
İmmünsüprese
Solunumsal asidozlu
KOAH Akut kardiyojenik
pulmoner ödem
COVID-19**

7 günde entübasyon / ölüm

High-Flow Nasal Oxygen vs Noninvasive Ventilation in Patients With Acute Respiratory Failure

JAMA

The RENOVATE Randomized Clinical Trial 2025 Mar 11;333(10):875-890

Nonimmünsüprese (n=485)
Solunumsal asidozlu KOAH alevlenmesi (n=77)
Akut kardiyojenik pulmoner ödem (n=272)
COVID-19 (n=882)

**Entübasyon / Ölümde
HFNO NIV'e non-inferior**

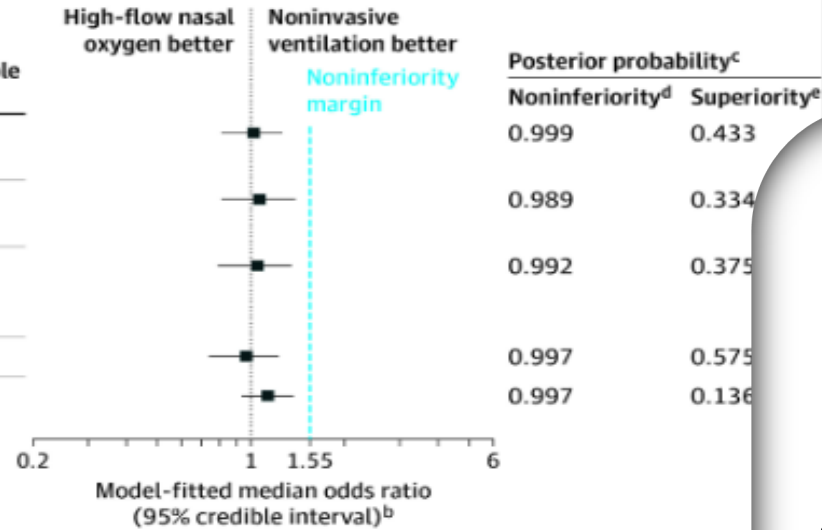
High-Flow Nasal Oxygen vs Noninvasive Ventilation in Patients With Acute Respiratory Failure

JAMA

The RENOVATE Randomized Clinical Trial 2025 Mar 11;333(10):875-890

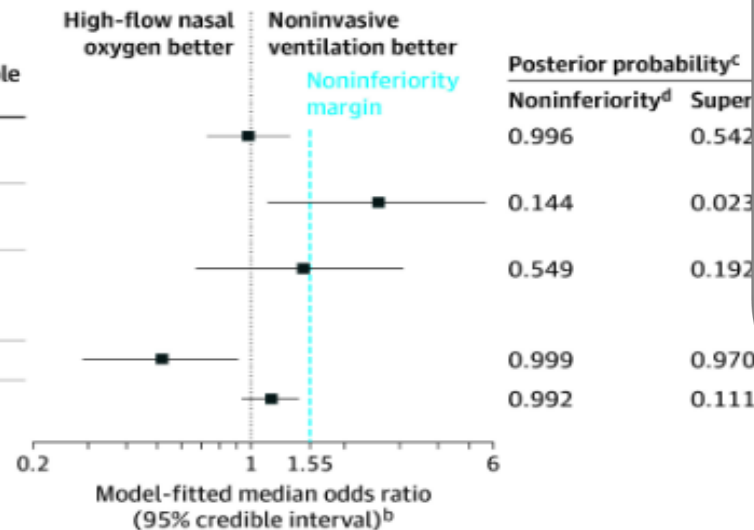
A Analysis of the primary outcome^a

Patients with acute respiratory failure	No./total (%)		Model-fitted median odds ratio (95% credible interval) ^b
	High-flow nasal oxygen	Noninvasive ventilation	
Nonimmunocompromised with hypoxemia	81/249 (32.5)	78/236 (33.1)	1.02 (0.81-1.26)
Immunocompromised with hypoxemia	16/28 (57.1)	8/22 (36.4)	1.07 (0.81-1.39)
Chronic obstructive pulmonary disease exacerbation with respiratory acidosis	10/35 (28.6)	11/42 (26.2)	1.05 (0.79-1.36)
Acute cardiogenic pulmonary edema	14/136 (10.3)	29/136 (21.3)	0.97 (0.73-1.23)
Hypoxemic COVID-19	223/435 (51.3)	210/447 (47.0)	1.13 (0.94-1.38)



B Post hoc analysis of the primary outcome^f

Patients with acute respiratory failure	No./total (%)		Model-fitted median odds ratio (95% credible interval) ^b
	High-flow nasal oxygen	Noninvasive ventilation	
Nonimmunocompromised with hypoxemia	81/249 (32.5)	78/236 (33.1)	0.98 (0.73-1.33)
Immunocompromised with hypoxemia	16/28 (57.1)	8/22 (36.4)	2.56 (1.14-5.68)
Chronic obstructive pulmonary disease exacerbation with respiratory acidosis	10/35 (28.6)	11/42 (26.2)	1.48 (0.67-3.09)
Acute cardiogenic pulmonary edema	14/136 (10.3)	29/136 (21.3)	0.52 (0.29-0.91)
Hypoxemic COVID-19	223/435 (51.3)	210/447 (47.0)	1.16 (0.94-1.43)



İmmüsuprese grup: Entübasyon /ölüm HFNO %57.1, NIV %36.4

Outcomes in immunocompromised patients with acute hypoxemic respiratory failure treated by high-flow nasal oxygen



RESPIR-OH, IVNICTUS, HIGH ve retrospektif kohort (EFRAIM) birleştirilmiştir.

n=986

28 günde IMV gereksinimi

28-gün mortalite ve ROX indeksinin öngörüsü

**HFNO sonrası IMV oranı %46, 28-gün mortalite %33
ROX için AUC \approx 0.67**

İmmünkompromize AHSY hastalar heterojen SOT'lı hastalar en iyi sonuçlara sahipken, hematolojik ve özellikle miyeloid malignitelerde HFNO başarısızlığı ve mortalite yüksektir.

Is there still a place for noninvasive ventilation in acute hypoxemic respiratory failure?

Table 1 Benefit–risk ratio assessment in favor or against NIV use in hypoxemic ARF patients

Indications for NIV use

- Acute exacerbation of COPD
- Acute cardiogenic pulmonary edema
- Hypoxemia post-abdominal surgery
- Chest trauma
- Preoxygenation before intubation

Against NIV use

- (Late or moderate–severe) ARDS
- High tidal volumes during the NIV session
- Leaks during the NIV session despite changes of interface
- Lack of patient adherence
- Dyspnea during NIV sessions
- Impossibility of close monitoring
- Absence of rapid clinical improvement (signs of respiratory distress including elevated respiratory rate) and gas exchange improvement after 1 h of NIV session

5 günden uzun NIV uygulaması sonrası entübasyon mortaliteyi %75'e yükseltmiştir.*

TV > 9–9.5 mL/kg PBW olması başarısızlık öngörücüsü

*Pneumologie. 2021 Jun;75(6):424–431. German. Epub 2021 May 11. PMID: 33975371.

TABLE 3

Outcomes

Primary outcome

Met the intubation criteria
(No. of events)

Secondary outcomes

Intubation with

Mortality within 30 days

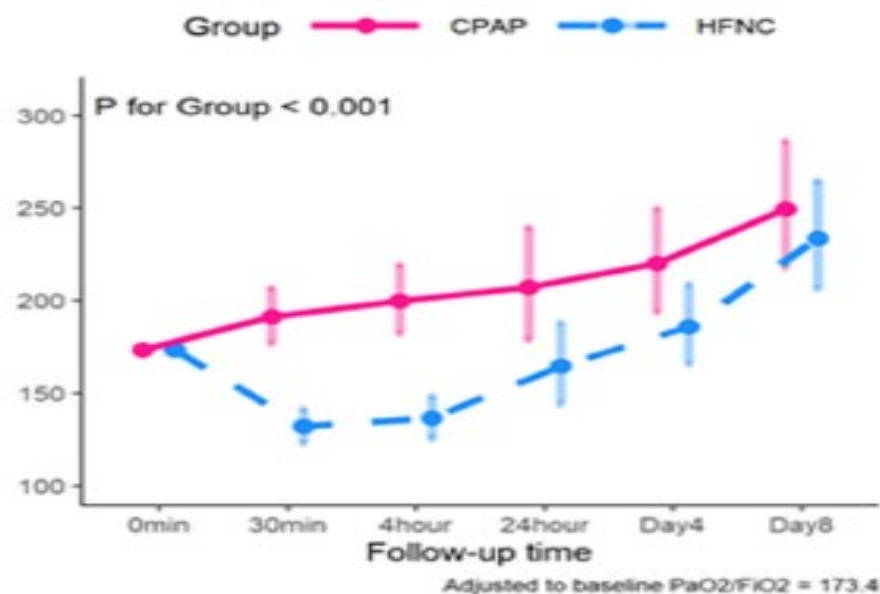
In-hospital mortality

Ventilator-free days

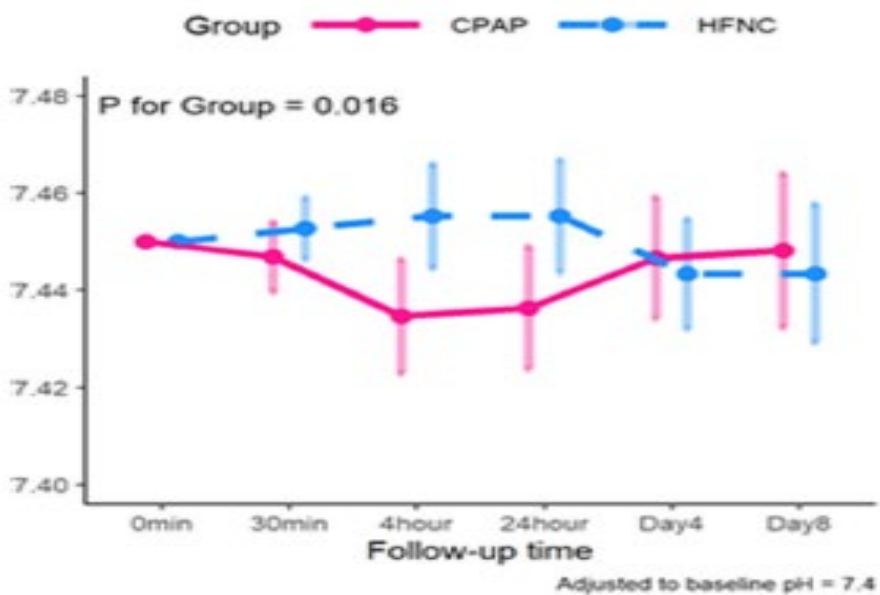
Duration of the first intubation

Duration of hospital stay

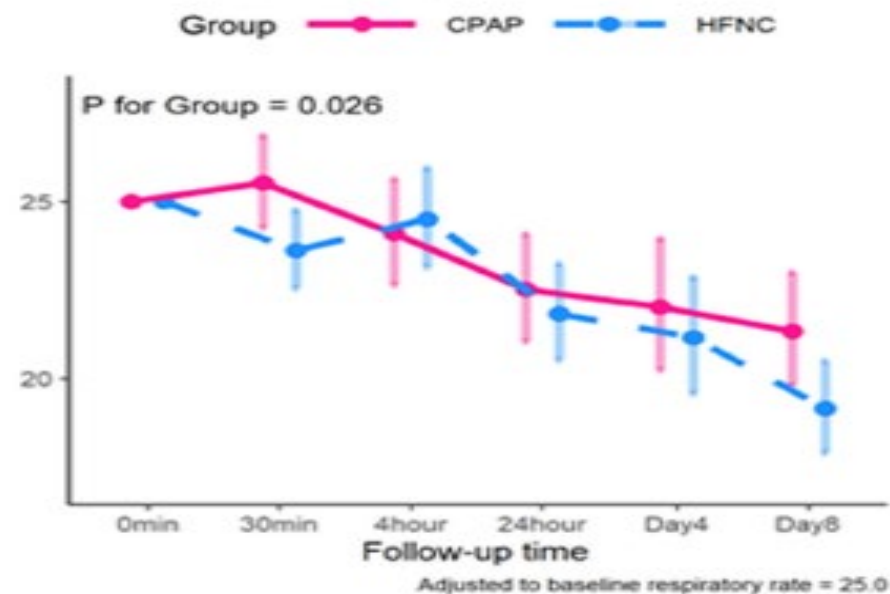
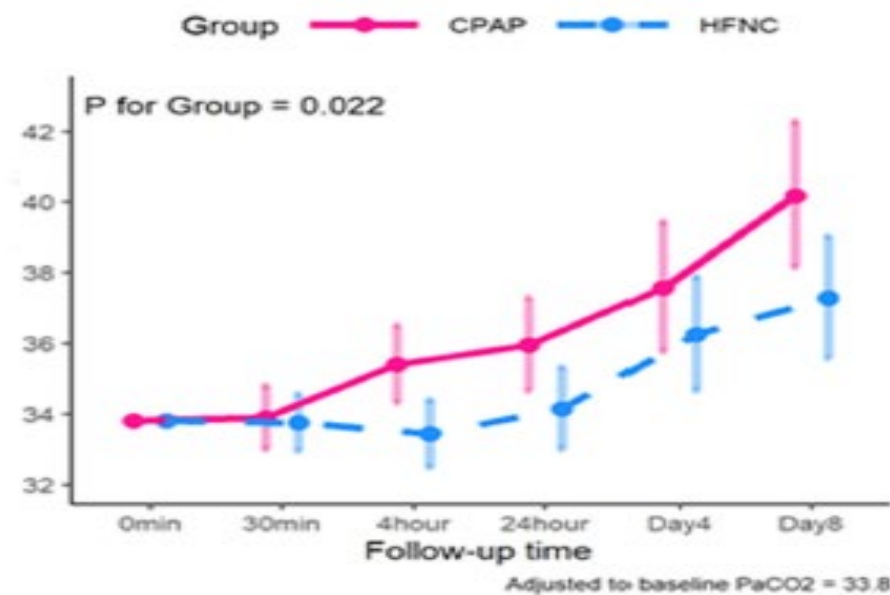
Need for continuous renal replacement therapy

(A) $\text{PaO}_2/\text{F}_1\text{O}_2$ ratio

(C) pH



(B) Respiratory rate

(D) PaCO_2 

P-value

0.006

0.385

0.363

0.378

0.744

0.405

0.741

Non-invasive ventilatory support and high-flow nasal oxygen as first-line treatment of acute hypoxemic respiratory failure and ARDS

Intensive Care Med (2021) 47:851–866

The role of non-invasive respiratory support (high-flow nasal oxygen and noninvasive ventilation) in the management of acute hypoxemic respiratory failure and acute respiratory distress syndrome is debated. The oxygenation improvement coupled with lung and diaphragm protection produced by non-invasive support may help to avoid endotracheal intubation, which prevents the complications of sedation and invasive mechanical ventilation. However, spontaneous breathing in patients with lung injury carries the risk that vigorous inspiratory effort, combined or not with mechanical increases in inspiratory airway pressure, produces high transpulmonary pressure swings and local lung overstretch. This ultimately results in additional lung damage (patient self-inflicted lung injury), so that patients intubated after a trial of noninvasive support are burdened by increased mortality. Reducing inspiratory effort by high-flow nasal oxygen or delivery of sustained positive end-expiratory pressure through the helmet interface may reduce these risks. In this physiology-to-bedside review, we provide an updated overview about the role of noninvasive respiratory support strategies as early treatment of hypoxemic respiratory failure in the intensive care unit. Noninvasive strategies appear safe and effective in mild-to-moderate hypoxemia ($\text{PaO}_2/\text{FiO}_2 > 150 \text{ mmHg}$), while they can yield delayed intubation with increased mortality in a significant proportion of moderate-to-severe ($\text{PaO}_2/\text{FiO}_2 \leq 150 \text{ mmHg}$) cases. High-flow nasal oxygen and helmet noninvasive ventilation represent the most promising

2026 SURVIVING SEPSIS CAMPAIGN GUIDELINES

Access the latest guidelines for sepsis treatment

COT yerine HFNO

NIV yerine HFNO

Dönüşümlü NIV+ HFNO yerine HFNO

Sedasyonsuz Uyanık Pron

Klasik Parametrelerin Sınırları

- Solunum sayısı, ROX indeksi ve HACOR skorları = inspiratuvar eforun **dolaylı** yansıması
 - Özefagus manometrisi (Δp_{pe}) **altın standart** , ama..

ROX İndeksi: Yüksek Akışlı Oksijen Tedavisinin Başarısını Öngörmek

ROX indeksi, Yüksek Akışlı Nazal Oksijen (HFNO) tedavisi gören hastalarda tedavinin başarısız olma olasılığını ve entübasyon ihtiyacını tahmin etmek için kullanılan basit, non-invaziv bir yatak başı aracıdır. Zaman içindeki değeri, hastanın klinik gidişatı hakkında önemli bilgiler sunar.



ROX İndeksi Nedir?

SpO₂/FiO₂'nin solunum sayısına bölünmesiyle elde edilen ve HFNO tedavisinin başarısını öngören bir araçtır.

Nasıl Hesaplanır?

$$\text{ROX İndeksi} = \frac{\text{SpO}_2}{\text{FiO}_2 \times \text{Solunum Sayısı (dakikada)}}$$



HFNO Başladıktan
2 Saat Sonra



ROX indeksi < 2.85 ise tedavi başarısızlığı ve entübasyon riski yüksektir.



HFNO Başladıktan
6 Saat Sonra



ROX indeksi < 3.47 ise tedavi başarısızlığı ve entübasyon riski yüksektir.



HFNO Başladıktan
12 Saat Sonra



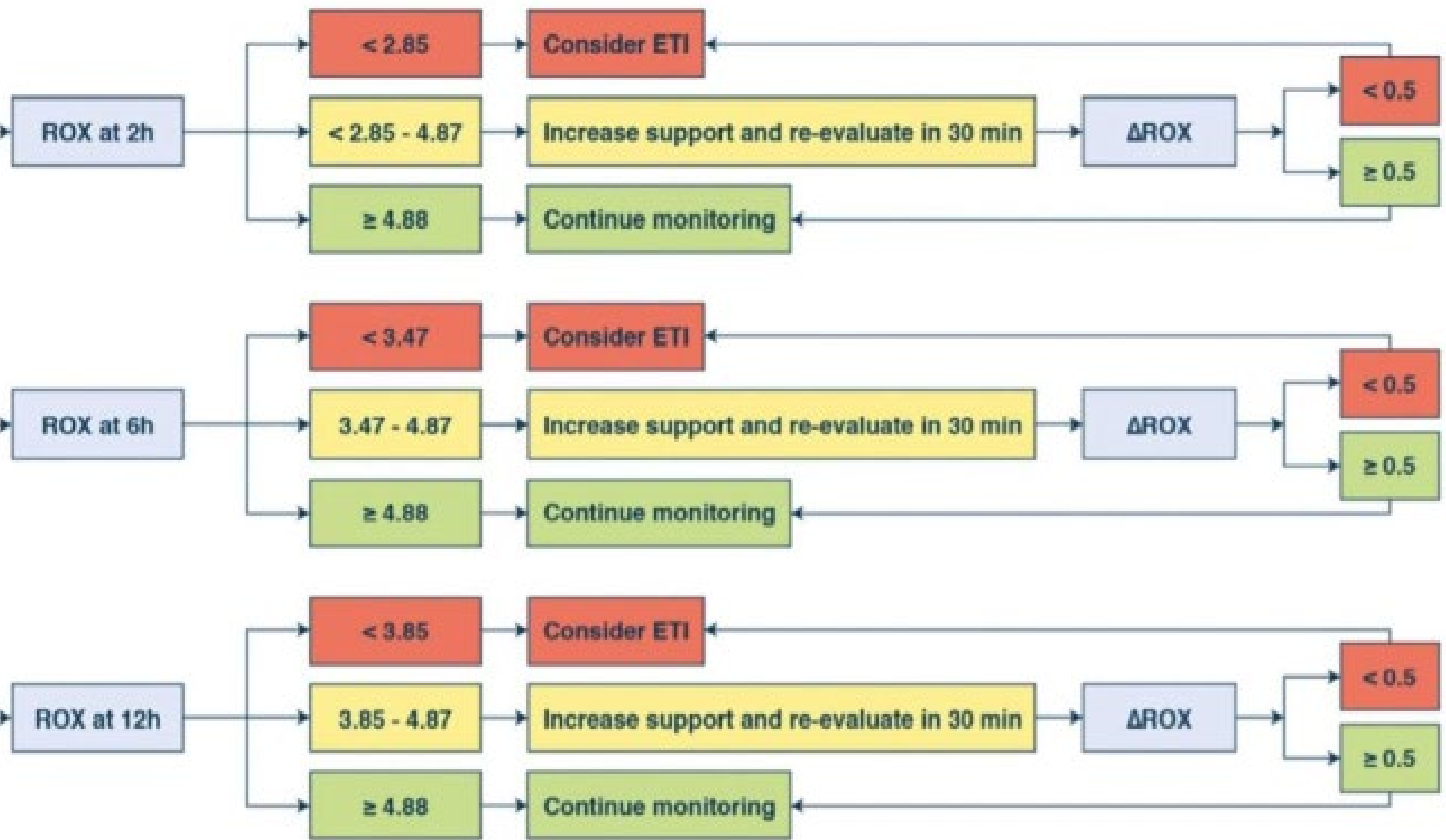
ROX indeksi < 3.85 ise tedavi başarısızlığı ve entübasyon riski yüksektir.

Önemli Not

Bu indeks yalnızca Yüksek Akışlı Nazal Oksijen (HFNO) tedavisi için doğrulanmıştır. Entübasyonu geciktirmemek için yakın takip zorunludur.



NHF onset



RESEARCH

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The ROX index as a predictor of high-flow nasal cannula outcome in pneumonia patients with acute hypoxemic respiratory failure: a systematic review and meta-analysis

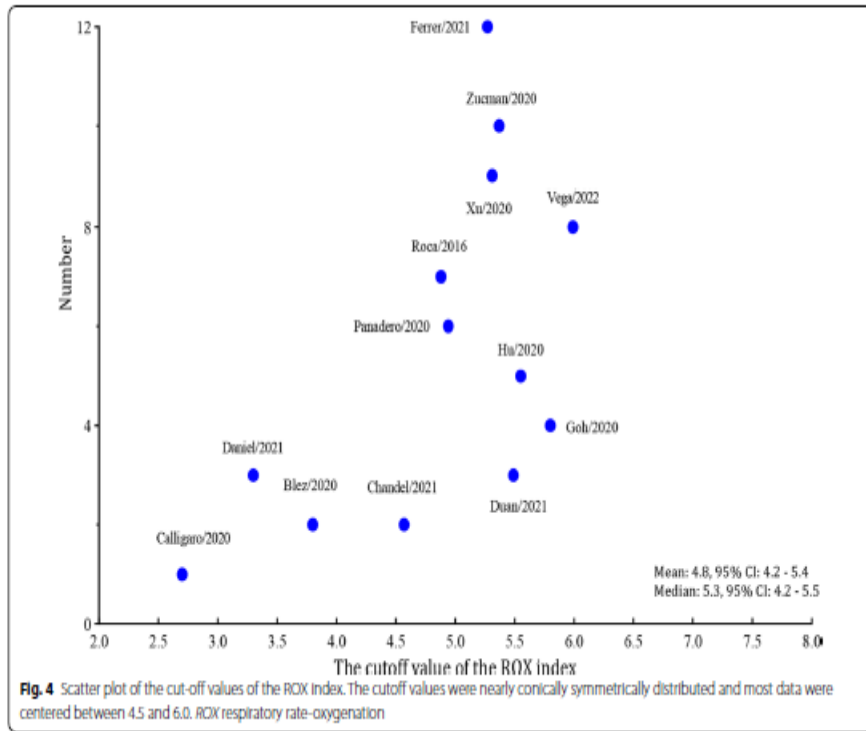


Fig. 4 Scatter plot of the cut-off values of the ROX index. The cutoff values were nearly conically symmetrically distributed and most data were centered between 4.5 and 6.0. ROX respiratory rate-oxygenation

6 prospektif, 7 retrospektif; n= 1751
10 çalışma COVID-19
ROX hem erken hem de 12 saate kadar benzer doğrulukta

ROX **<4.2** → entübasyon düşün

4.2–5.4 → **Dikkat**

• **>5.4** → iyi

AUHSROC = **0.81 (95% CI 0.77–0.84)**

• Pooled sensitivite = %71

• Pooled spesifisite = %78

• Tanısal odds ratio (DOR) = 8.3

RESEARCH

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Association between the ROX index and mortality in patients with acute hypoxemic respiratory failure: a retrospective cohort study

**ROX: ilk 24 saatin ortalaması alınmış-
Fizyolojik?**

n= 813

28 günlük mortalite

3-6-12 ay mortalite, ICU ve hastane mortalitesi

**Sonuç: nonlinear- güçlü ilişki
*ROX>8.28 ilişki kayboluyor.**

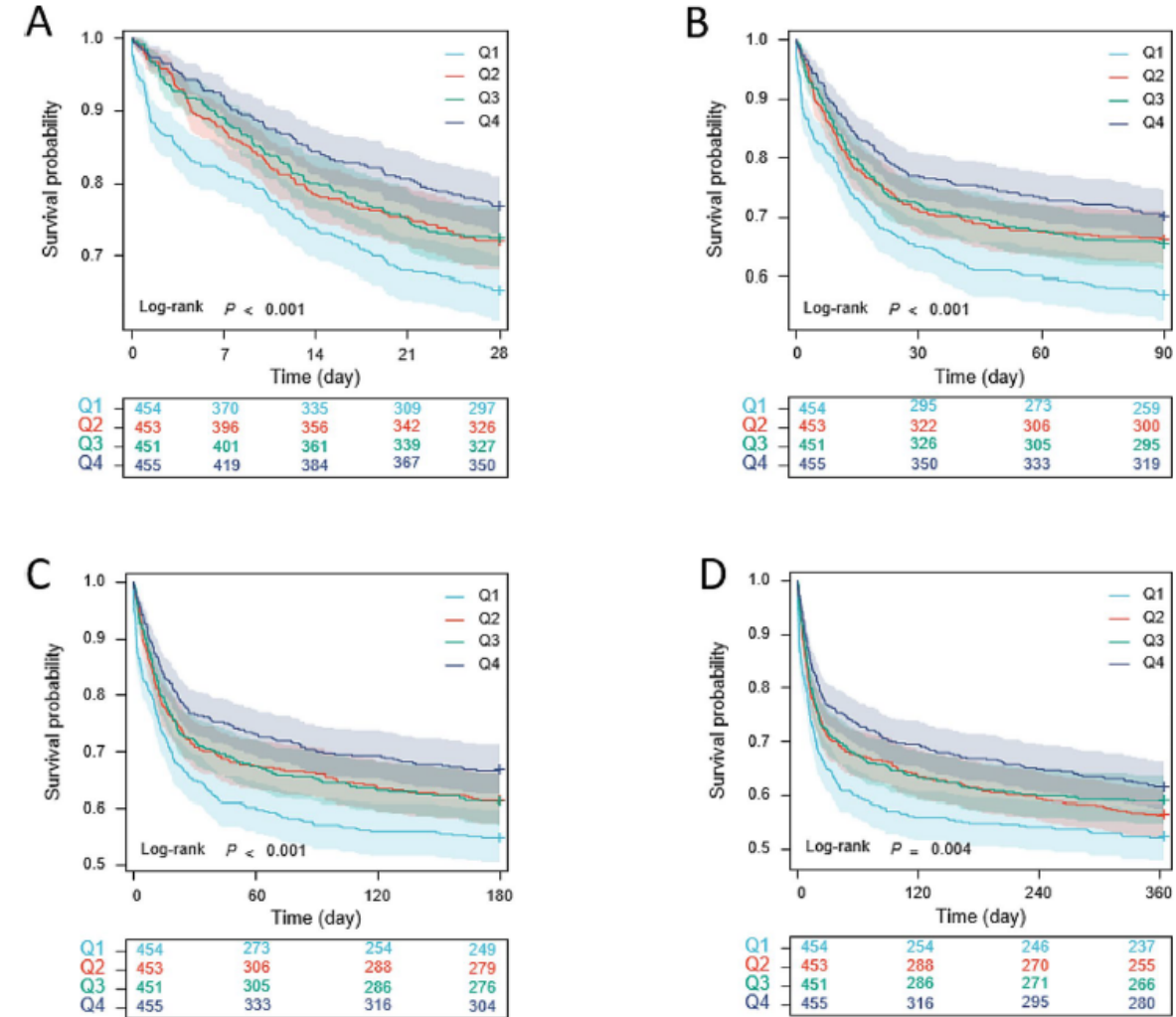


Fig. 4 Kaplan–Meier survival analysis curves for all-cause mortality. ROX index: Q1 (ROX ≤ 5.89), Q2 (5.89 < ROX ≤ 8.28), Q3 (8.28 < ROX ≤ 11.24), Q4 (ROX > 11.24). Kaplan–Meier curves showing cumulative probability of all-cause mortality according to groups at 28 days (A), 3 months (B), 6 months (C) and 1 year (D)

$$VOX = \frac{SpO_2/FiO_2}{VT}$$

Tek merkez, n=62

ROX ilk saatlerde yetersiz..

TV, HFNO başarısız olan hastalarda başlangıçtan itibaren belirgin yüksek. Özellikle >10 ml/kg PBW!

Table 1. Diagnostic Accuracy of Different Respiratory Variables at Different Time Points of Need for IMV in Patients Treated With HFNC

	Time (h)	AUROC	95% CI	P Value
SpO ₂ /FiO ₂	0	0.76	0.64–0.89	<0.001
	2	0.82	0.71–0.93	<0.001
	6	0.84	0.73–0.95	<0.001
Respiratory rate	0	0.50**	0.35–0.64	0.944
	2	0.42**	0.28–0.56	0.271
	6	0.51**	0.36–0.66	0.893
ROX index	0	0.66**	0.52–0.80	0.034
	2	0.70*	0.56–0.85	0.006
	6	0.79**	0.68–0.91	0.001
VT (ml/kg of PBW)	0	0.83	0.71–0.94	<0.001
	2	0.87	0.77–0.97	<0.001
	6	0.87	0.76–0.98	<0.001
VOX index	0	0.84	0.75–0.94	<0.001
	2	0.88	0.79–0.97	<0.001
	6	0.93	0.86–0.99	<0.001

HACOR skoru

Variable	Value	Score
Heart rate (HR)	≤ 120	0
	> 120	1
Acidosis (pH)	≥ 7.35	0
	7.30 - 7.34	2
	7.25 - 7.29	3
	< 7.25	4
Consciousness (GCS)	15	0
	13 - 14	2
	11 - 12	5
	≤ 10	10
Oxygenation (PaO ₂ /FiO ₂)	> 200	0
	176 - 200	2
	151 - 175	3
	126 - 150	4
	101 - 125	5
	≤ 100	6
Respiratory rate (RR)	≤ 30	0
	31 - 35	1
	36 - 40	2
	41 - 45	3
	> 45	4

>5

1 Saatte >5 Puan Yüksek Başarısızlık Riski Anlamına Gelir.

Bu eşik değer, farklı hasta gruplarında NIV başarısızlığını öngörmeye etkilidir.

Zaman İçinde Tahmin Gücü



RESEARCH

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An updated HACOR score for predicting the failure of noninvasive ventilation: a multicenter prospective observational study



Variable	Value	Score
Heart rate (HR)	≤ 120	0
	> 120	1
Acidosis (pH)	≥ 7.35	0
	7.30 - 7.34	2
	7.25 - 7.29	3
	< 7.25	4
Consciousness (GCS)	15	0
	13 - 14	2
	11 - 12	5
	≤ 10	10
Oxygenation (PaO ₂ /FiO ₂)	> 200	0
	176 - 200	2
	151 - 175	3
	126 - 150	4
	101 - 125	5
	≤ 100	6
Respiratory rate (RR)	≤ 30	0
	31 - 35	1
	36 - 40	2
	41 - 45	3
	> 45	4

18 merkez, prospektif çalışma (n=2179)

6 NIV öncesi değişken eklenmiştir:

SOFA skoru

Pnömoni varlığı

Kardiyojenik pulmoner ödem

Pulmoner ARDS

İmmünsüpresyon

Septik şok

RESEARCH

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An updated HACOR score for predicting the failure of noninvasive ventilation: a multicenter prospective observational study



Güncellenmiş HACOR skorunun **NIV başladıktan 1–2 saat sonra** hesaplanması, en güçlü sonuçları vermiştir.

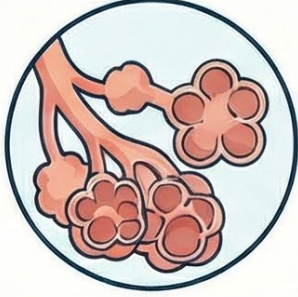
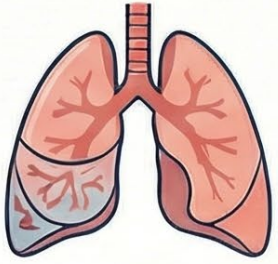
AUC \approx 0.85

- ≤ 7 puan \rightarrow Düşük risk (NIV başarısızlığı %12)
- 7.5–10.5 \rightarrow Orta risk (%38)
- 11–14 \rightarrow Yüksek risk (%67)
- 14 \rightarrow Çok yüksek risk (%84)

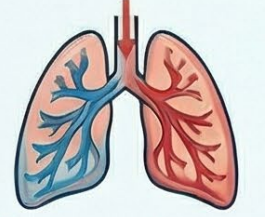
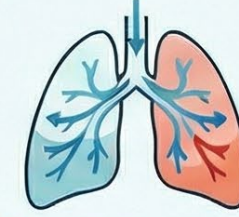
Pron Pozisyonunun Fizyolojik Gerekçeleri

Hastalarda yüzüstü pozisyon uygulamasının akciğer fonksiyonları üzerindeki temel fizyolojik faydaları ve etki mekanizmaları.

Mekanik ve Yapısal İyileşme



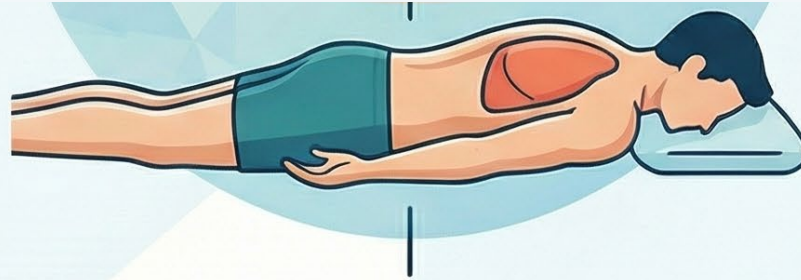
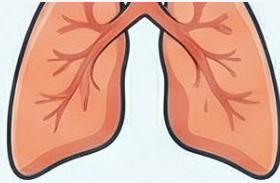
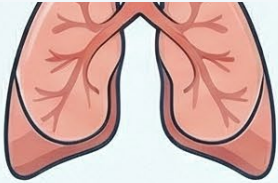
Fonksiyonel Faydalar ve Klinik Yanıt



V/Q Uyumunun İyileşmesi

Perfüzyonun havalandırılan bölgelere yönelmesiyle ventilasyon-perfüzyon eşleşmesi belirli şekilde düzelir.

Kılavuz Önerisi: ATS/ESICM/SCCM Klinik Uygulama Kılavuzu (2017) — Orta-ağır ARDS'de ($PaO_2/FiO_2 < 150$ mmHg) günde ≥ 12 saat pron pozisyon için güçlü öneri.



Yer çekimi desteğiyle drenaj kolaylaşır, küçük hava yolları açılır ve enfeksiyon riski azalır.



Homojen Stres Dağılımı

Transpulmoner basınç eşitlenerek ventilatör kaynaklı akciğer hasarı (VILI) riski düşürülür.



1-4 Saat İçinde Klinik Yanıt

Fizyolojik yanıt veren hastalarda oksijenasyon iyileşmesi genellikle bu sürede gözlemlenir.

- «HFNO mı, NIV mi?» tercihte belirsizlikler devam etmekte.
- **De novo hipoksemik solunum yetmezliğinde, sepsise bağlı AHSY’de**
HFNO hala ilk öneri, konforlu
- **Tedavi başarısı sadece oksijenasyonla değil, inspiratuar efor, WOB ve**
pulmoner heterojeniteye göre incelenmelidir.
- **Yüksek inspiratuar efor → Helmet + Pressure Support koruyucu**
olabilir;yine de çalışmalara ihtiyaç var
 - **İmmünsuprese grupta belirsizlikler devam ediyor**
- **Oksijenasyon zayıfsa → Helmet NIV, HFNO’den daha etkili olabilir; hastaya göre**
karar vermek gerekir.
 - **Başarısızlığı ERKEN tanınmalı.**

TEŐEKKÜRLER