

# 10 věcí, kterých bych se měl u pacienta v těžké sepsi vyvarovat



Beneš Jan

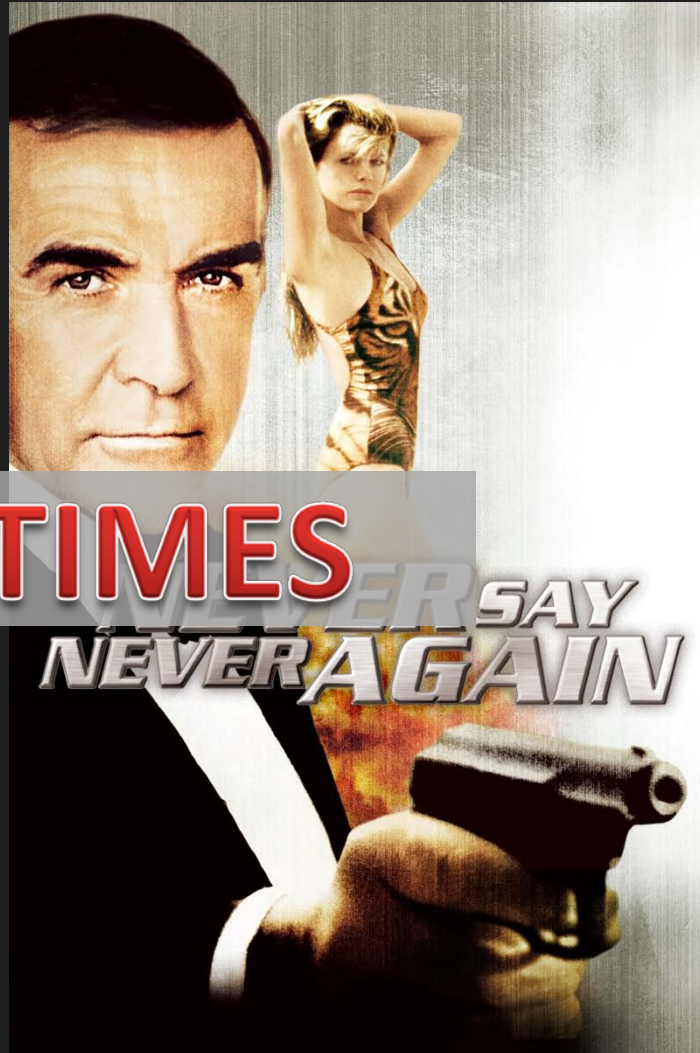


Anesteziologicko-resuscitační klinika, Fakultní nemocnice a Lékařská Fakulta v Plzni  
Univerzity Karlovy v Praze



## Konflikt zájmu:

- **Práce na dlouhodobém výzkumu méně invazivních metod monitorace krevního oběhu na ARK byly podpořeny výzkumným záměrem MSM 0021620819 a Projektem 36 (PRVOUK)**
- **Monitorovací prostředky pro výzkum byly zapůjčeny fy. CNSystems (Graz, Rakousko) a Edwards Lifescience Inc (Irvine, USA)**
- **Člen** Advisory board fy Edwards Lifesciences Inc.



**10 TIMES**  
**NEVER SAY**  
**NEVER AGAIN**

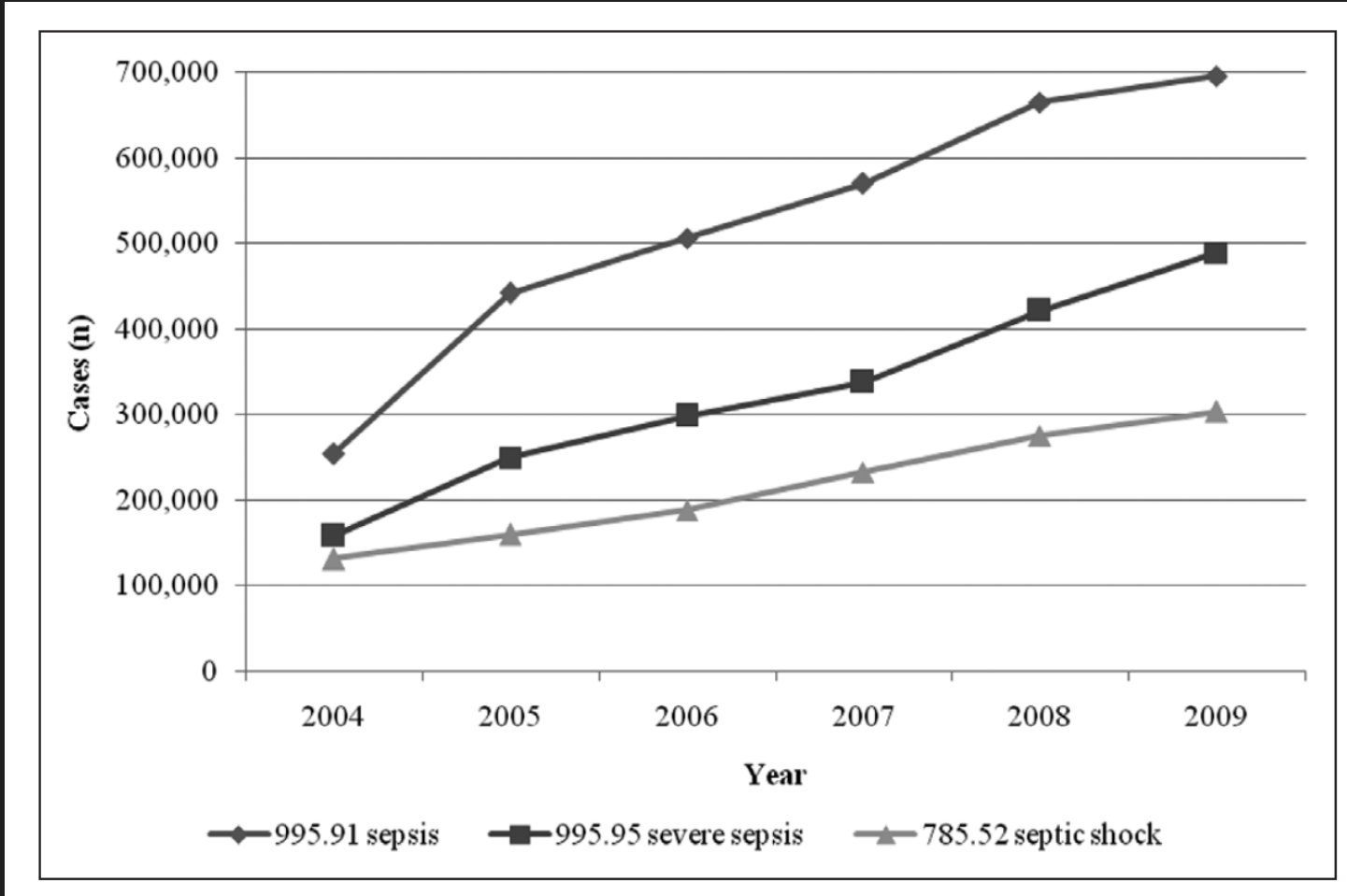
# 1. Nevzdávat to



# Benchmarking the Incidence and Mortality of Severe Sepsis in the United States\*

(*Crit Care Med* 2013; 41:1167-1174)

David F. Gaieski MD<sup>1</sup>; J. Matthew Edwards, MD<sup>1</sup>; Michael J. Kallan, MS<sup>2</sup>; Brendan G. Carr, MD, MA, MS<sup>1-3</sup>



# Mortality Related to Severe Sepsis and Septic Shock Among Critically Ill Patients in Australia and New Zealand, 2000-2012

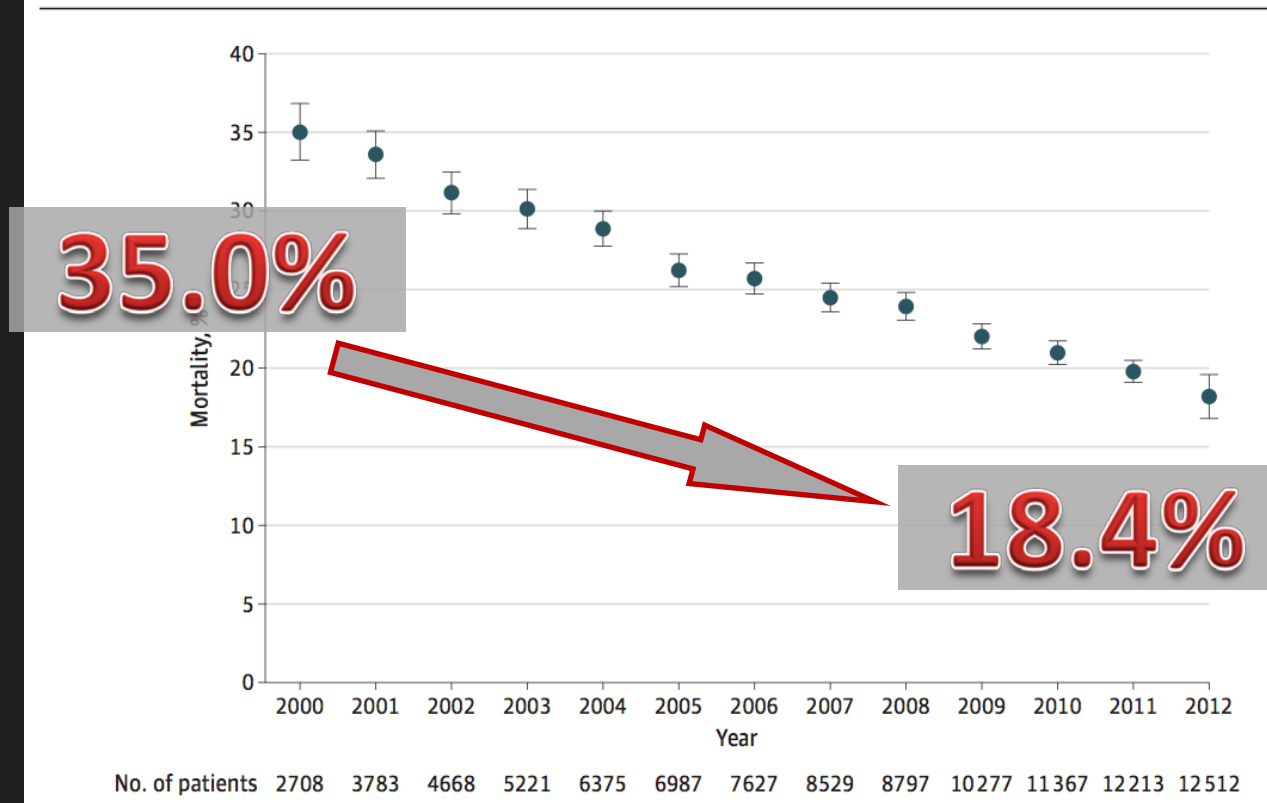
Kirsi-Majja Kaukonen, MD, PhD, EDIC<sup>1,2</sup>; Michael Bailey, PhD<sup>1</sup>; Satoshi Suzuki, MD<sup>3</sup>; David Pilcher, FCICM<sup>1,4,5</sup>; Rinaldo Bellomo, MD, PhD<sup>1,3</sup>

[+] Author Affiliations

JAMA. 2014;311(13):1308-1316. doi:10.1001/jama.2014.2637.

Text Size: A A A

Figure 1. Mean Annual Mortality in Patients With Severe Sepsis



## 2. Neztráčet čas



GOAL-DIRECTED

EARLY GOAL-DIRECTED THERAPY IN THE TREATMENT OF SEVERE SEPSIS AND SEPTIC SHOCK

EMANUEL RIVERS, M.D., MICHAEL M. COLE, M.D., SUZANNE HAVSTAD, M.A., JULIE RESSLER, B.S., ALEXANDRIA MUZZIN, B.S., BERNHARD KNOBLOCH, M.D., JEFFREY S. STERNSON, PH.D., AND MICHAEL TOMLANOVICH, M.D., FOR THE EARLY GOAL-DIRECTED THERAPY GROUP\*

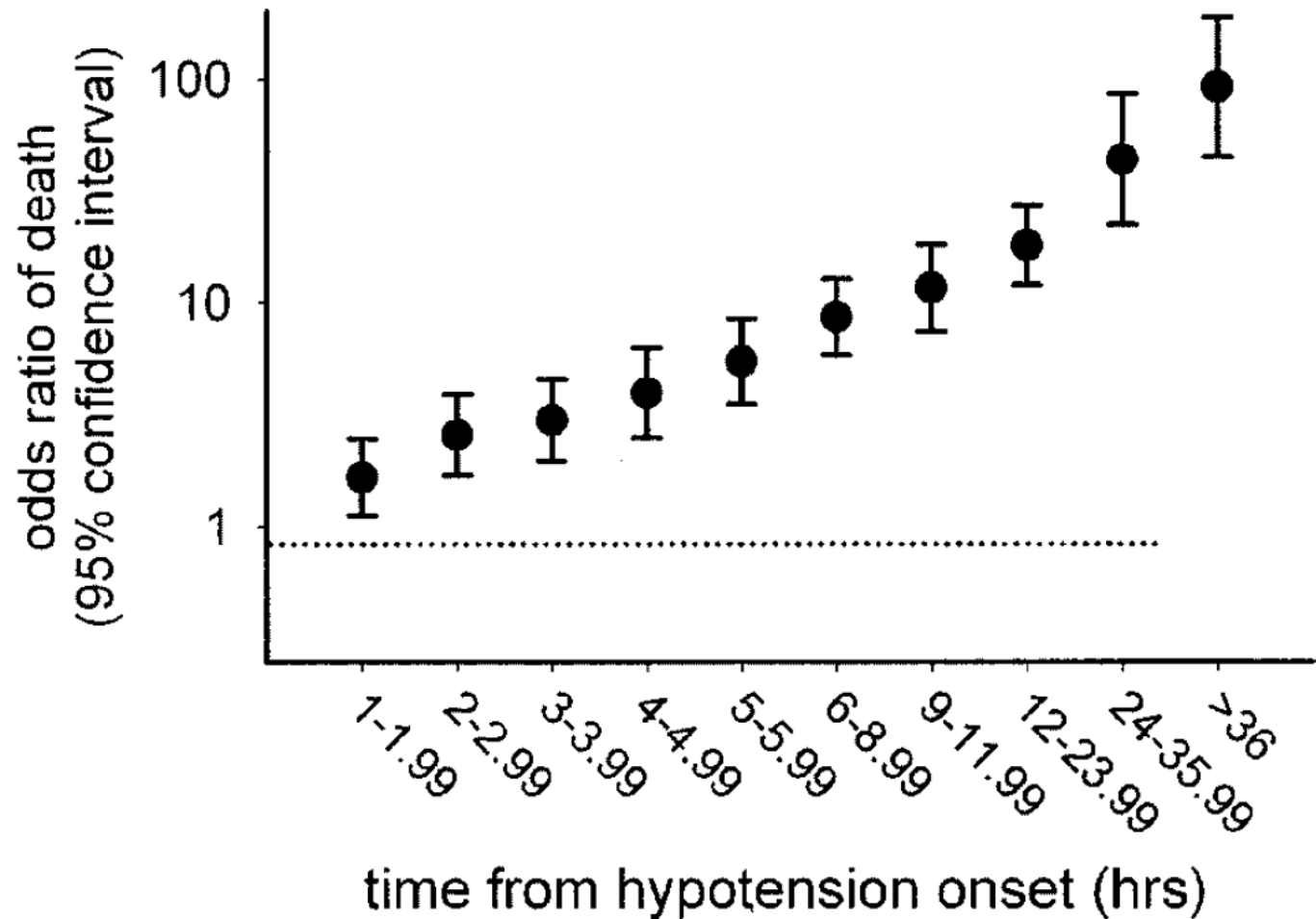
263 pts

Mortality 46% vs 30%

EARLY GO

Duration of hypotension before initiation of effective antimicrobial therapy is the critical determinant of survival in human septic shock\*  
(Crit Care Med 2006; 34:1589-1596)

Anand Kumar, MD; Daniel Roberts, MD; Kenneth E. Wood, DO; Bruce Light, MD; Joseph E. Parrillo, MD; Satendra Sharma, MD; Robert Suppes, BSc; Daniel Feinstein, MD; Sergio Zanotti, MD; Leo Taiberg, MD; David Gurka, MD; Aseem Kumar, PhD; Mary Cheang, MSc



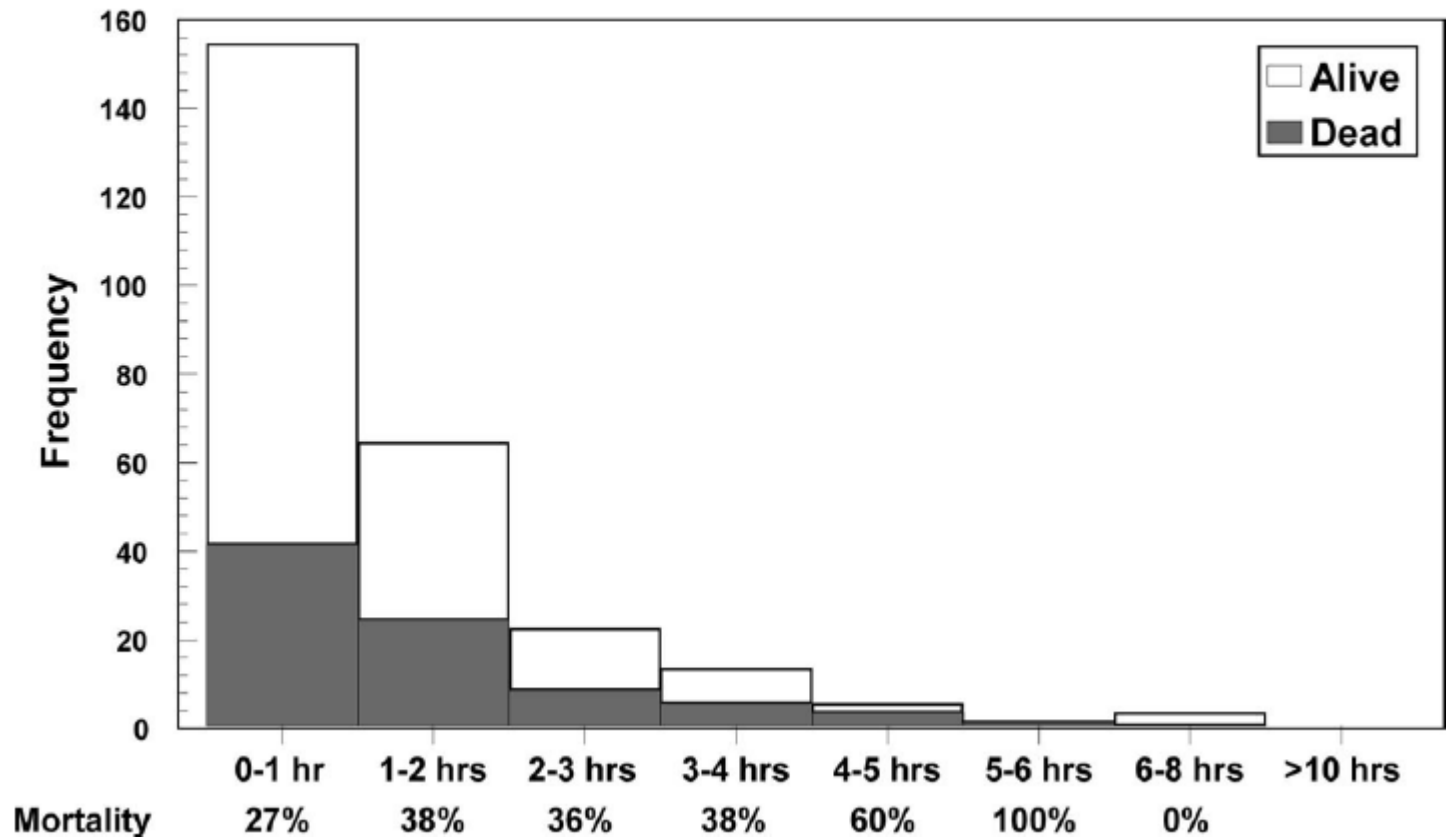


Impact of time to antibiotics on survival in patients with severe sepsis or septic shock in whom early goal-directed therapy was initiated in the emergency department

(*Crit Care Med* 2013; 41:1167-1174)

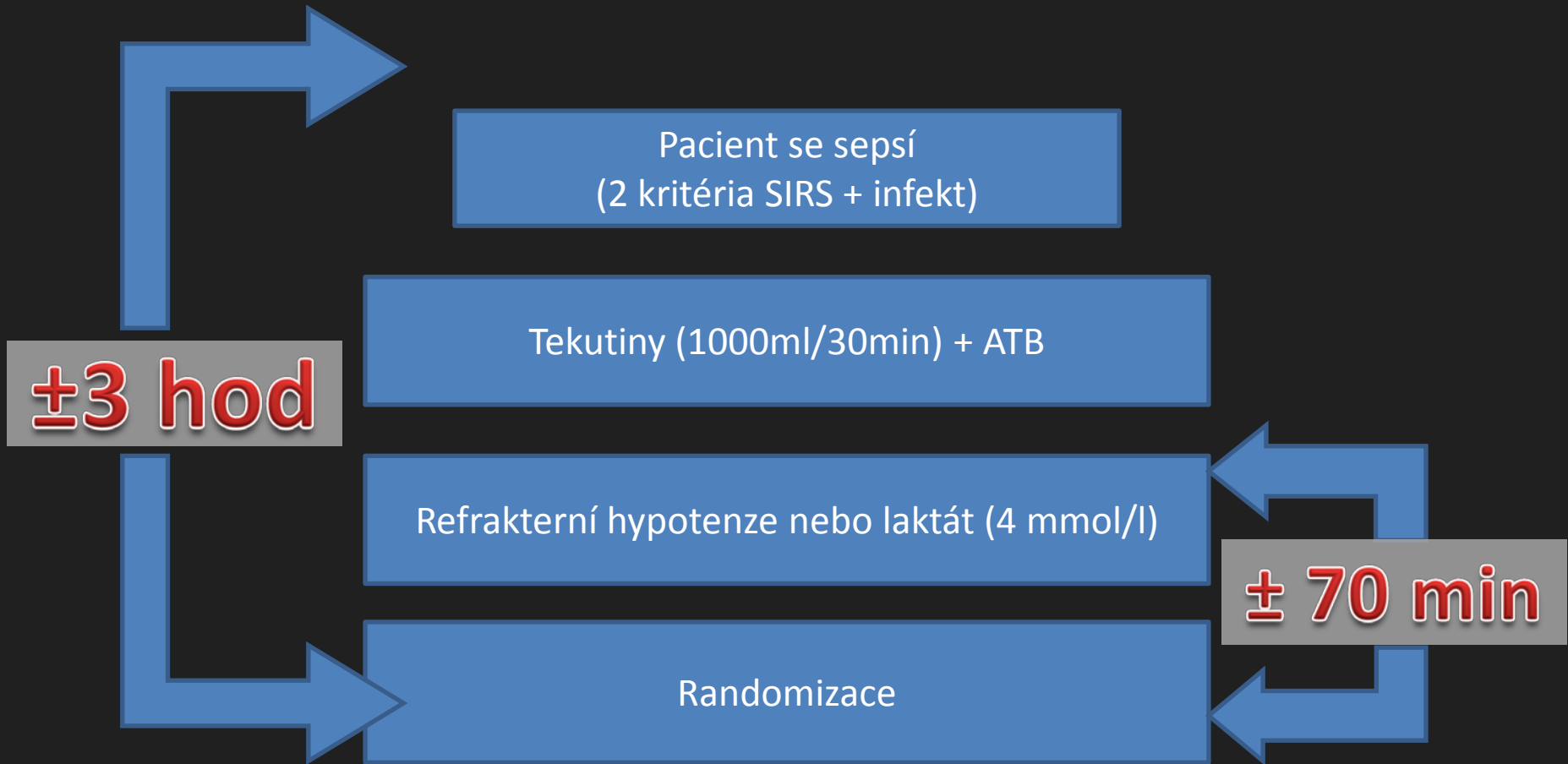
David F. Gaieski, MD; Jesse M. Pines, MD, MBA, MSCE; Roger A. Band, MD;  
Mark E. Mikkelsen, MD, MSCE; Richard Massone, MD; Frances F. Furia, MD; Frances S. Shofer, PhD;  
Munish Goyal, MD

## Time from Qualification for EGDT to Antibiotics



# A Randomized Trial of Protocol-Based Care for Early Septic Shock

The ProCESS Investigators\*



## A Randomized Trial of Protocol-Based Care for Early Septic Shock

The ProCESS Investigators\*

In summary, in our multicenter, randomized trial, **U PACIENTŮ, KTEŘÍ BYLI VČAS IDENTIFIKOVÁNÍ NA URGENTNÍM PŘÍJMU A KTEŘÍ DOSTALI RYCHLE ANTIMIKROBIÁLNÍ TERAPII A ZÁKLADNÍ OBJEMOVOU LÉČBU NEBYL PROKÁZÁN VÝZNAMNÝ EFEKT REUSCITAČNÍCH PROTOKOLŮ**

use of central venous catheterization and central hemodynamic monitoring in all patients.

### 3. Nebát se počáteční racionální terapeutické agresivity

A Randomized Trial of Protocol-Based Care for Early Septic Shock

The ProCESS Investigators\*

In summary, in our multicenter, randomized trial, in which patients were identified early in the emergency department as having septic shock and received antibiotics and other nonresuscitation aspects of care promptly, we found no significant advantage, with respect to mortality or morbidity, of protocol-based resuscitation over bedside care that was provided accord-

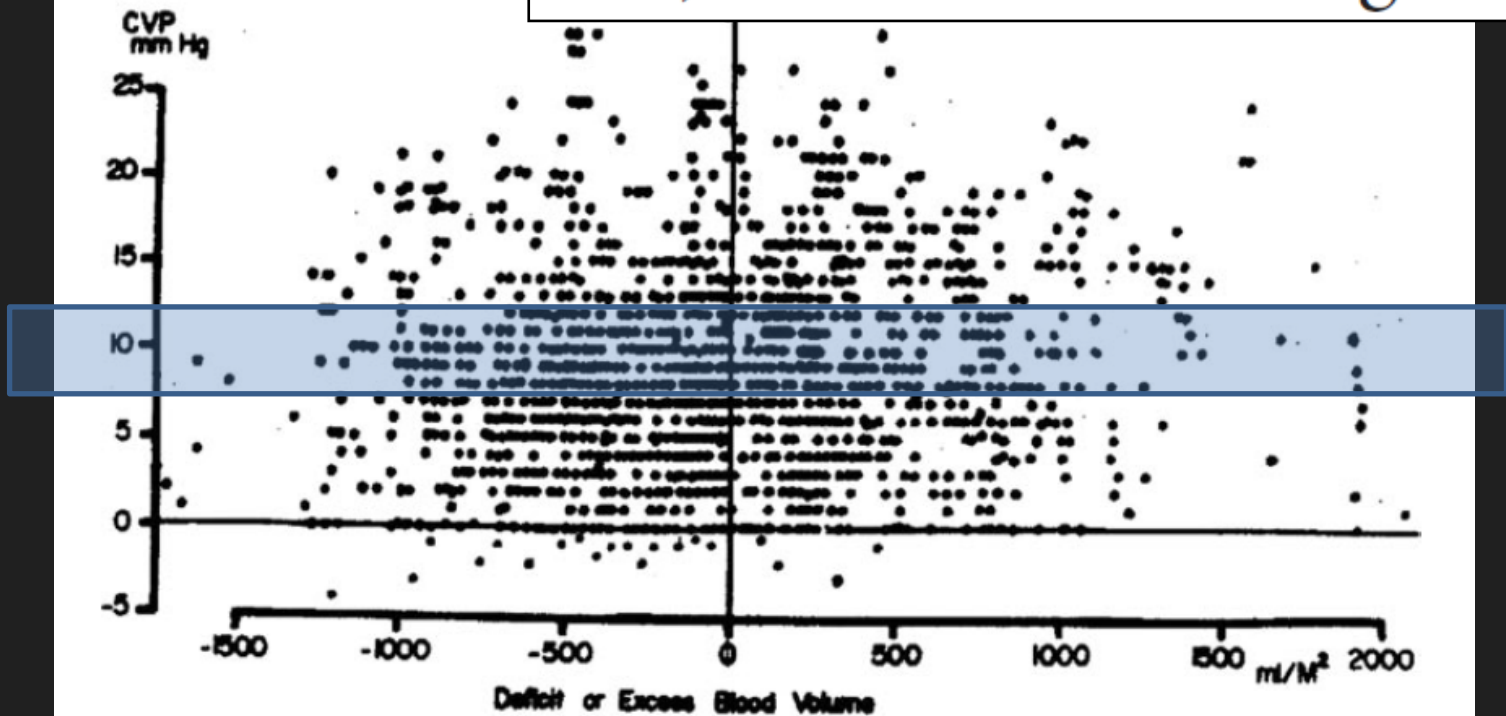
**CENTRÁLNÍ ŽILNÍ KATETRIZACE SE  
NEUKÁZALA VÝHODNÁ U VŠECH  
NEMOCNÝCH**

# Does Central Venous Pressure Predict Fluid Responsiveness?\*

A Systematic Review of the Literature and the Tale of Seven Mares  
(*CHEST* 2008; 134:172–178)

Paul E. Marik, MD, FCCP; Michael Baram, MD, FCCP; and Bobbak Vahid, MD

a) CVP 8–12 mm Hg



# ŠOK

PÉČE O ADEKVÁTNÍ DODÁVKU  
KYSLÍKU DO TKÁNÍ

ODSTRANĚNÍ PŘÍČINY ŠOKU

ODSTRANĚNÍ **PŘÍČINY** ŠOKU

**ATB**

**FOKUS**

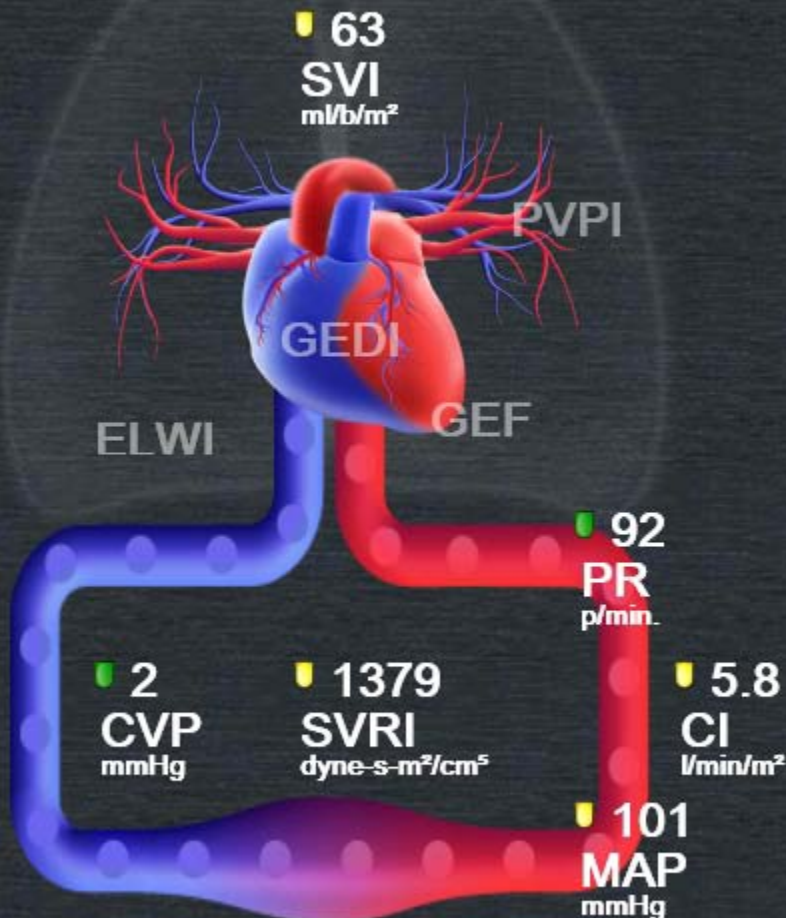
PÉČE O ADEKVÁTNÍ DODÁVKU  
KYSLÍKU DO TKÁNÍ

**PERFUZE**

**OXYGENACE**



ScvO<sub>2</sub>  
%



07:08:28

25.05.2012

20

BT: 37.4°C

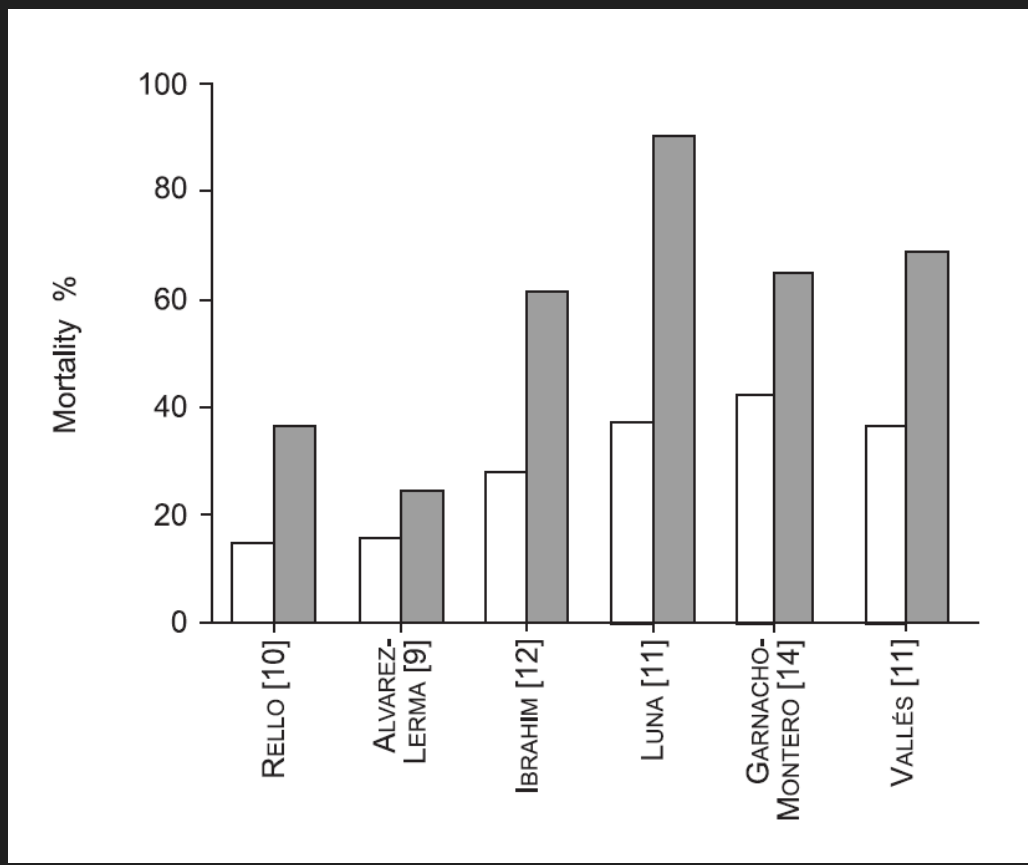


Chyba oxymetrie: Odpojení OM



# Importance of appropriate initial antibiotic therapy and de-escalation in the treatment of nosocomial pneumonia

J. Rello



# Benefit of Appropriate Empirical Antibiotic Treatment: Thirty-day Mortality and Duration of Hospital Stay

Abigail Fraser, MPH,<sup>a</sup> Mical Paul, MD,<sup>a,b</sup> Nadja Almanasreh, MD,<sup>c</sup> Evelina Tacconelli, MD,<sup>d</sup> Uwe Frank, MD,<sup>c</sup> Roberto Cauda, MD,<sup>d</sup> Sara Borok, MD,<sup>a</sup> Michal Cohen, MD,<sup>e</sup> Steen Andreassen, PhD,<sup>f</sup> Anders D. Nielsen, MSc,<sup>f</sup> Leonard Leibovici, MD,<sup>a,b</sup> on behalf of the TREAT Study Group

**Table 3** Duration of Hospitalization by Medical Center and Appropriateness of Antibiotic Treatment

	Appropriate Antibiotic Treatment	Inappropriate Antibiotic Treatment
Rabin Medical Center, Israel		
No. of patients	282	151
Mean (SD)	7.97 (8.58)	10.77 (13.90)
Median (range)	5 (0-62)	7 (1-125)
Freiburg University Hospital, Germany		
No. of patients	90	31
Mean $\pm$ SD	15.63 (8.83)	18.42 (13.83)
Median (range)	14 (1-36)	14 (3-65)
A. Gemelli University Hospital, Italy		
No. of patients	84	39
Mean $\pm$ SD	14.25 (11.63)	16.62 (12.20)
Median (range)	10.5 (1-62)	12 (2-48)
Total		
No. of patients	456	221
Mean $\pm$ SD	10.64 (13.90)	12.87 (9.86)
Median (range)*	7 (0-62)	8 (1-125)

SD = standard deviation.

\*Mann Whitney test,  $P = .048$ .

## 4. ... a její ekonomické / personální náročnosti

### SEPSE

Augmentin  
500,-/den  
Meronem  
6 000,-/den

HD monitorace  
5-10 tis

### AIM

ReoPro  
20 000,-  
Simdax  
30 000,-

TCA + stent  
cca 100 – 150 tis

### CMP

Actilyse  
20 000,-

Mechanická  
trombolýza  
, extrakce  
100 – 130 tis

# 5. Nedělat léčebnou kosmetiku



110/60  
(76)

99%

95/min

85 ml





<b>Astrup</b>		
B'-pH	7,22	
B'-pO2	7,6	
B'-pCO2	5,9	
B-BE	<b>-8,6</b>	
B-HCO3 aktualni	17,8	
B-Oxyhemoglobin	0,83	
B-Saturace Hb kys	0,85	
B-Hemoglobin	<b>106</b>	
B'-Teplota aktualní	36,5	
<b>Spec. bioch. vyšetře</b>		
P-Laktat		<b>7,59</b>

# Re-thinking resuscitation: leaving blood pressure cosmetics behind and moving forward to permissive hypotension and a tissue perfusion-based approach

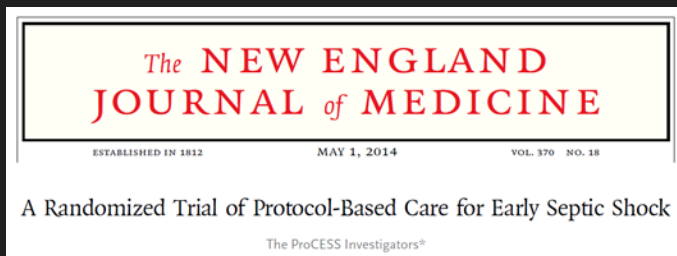
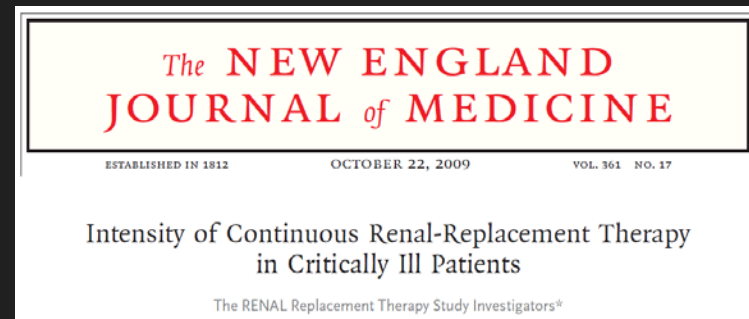
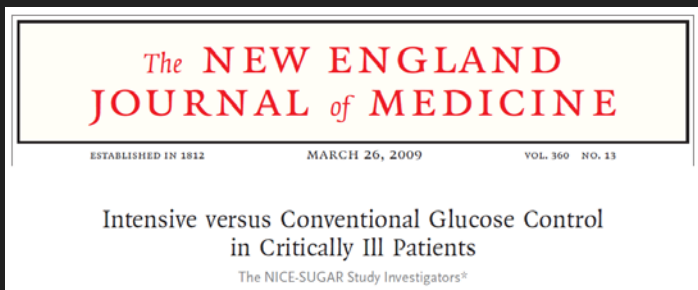
Martin W Dünser<sup>\*1</sup>, Jukka Takala<sup>2</sup>, Andreas Brunauer<sup>1</sup> and Jan Bakker<sup>3</sup>

Dünser *et al. Critical Care* 2013, **17**:326

**Table 1. Tissue perfusion-based resuscitation endpoints**

Category	Parameter	Endpoint	Suggested measurement interval	Pathophysiologic background	Therapeutic implications
Peripheral perfusion	Capillary refill time	<4.5 seconds	15-60 minutes <sup>a</sup>	Inadequate systemic blood flow	Fluids, red blood cells, inotropes, vasodilator.  Vasopressors only to ensure minimum mean arterial blood pressure for coronary and cerebral perfusion!
	Skin mottling	Absent			
	Peripheral temperature	Warm			
	Peripheral perfusion index	≥1.4	Continuous		
	Tissue oxygen saturation	≥70%			
Venous oxygen saturation	Central	≥65%-70%	Continuous		
	Mixed	≥60%-65%			
Arterial lactate	Absolute value	<2 mmol/L	2 hours	Inadequate systemic blood flow or excessive vasodilation	Fluids, blood, inotropes, vasodilators, and/or vasopressors
	Clearance	>20%/2 hours			
Urine output		≥0.5 mL/kg per hour	15-60 minutes <sup>a</sup>		

# 6. Nepodléhat módním trendům



The New England  
Journal of Medicine

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VOLUME 345      NOVEMBER 8, 2001      NUMBER 19

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INTENSIVE INSULIN THERAPY IN CRITICALLY ILL PATIENTS

GREET VAN DEN BERGHE, M.D., Ph.D., PETER WOUTERS, M.Sc., FRANK WEEKERS, M.D., CHARLES VERWAEST, M.D.,  
FRANS BRUYNINCKX, M.D., MET SCHEZ, M.D., Ph.D., DIRK VLASSELAERS, M.D., PATRICK FERDINANDE, M.D., Ph.D.,  
PETER LAUWERS, M.D., AND ROGER BOULLON, M.D., Ph.D.



*The* **NEW ENGLAND**  
**JOURNAL** *of* **MEDICINE**

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ESTABLISHED IN 1812      MARCH 26, 2009      VOL. 360 NO. 13

Intensive versus Conventional Glucose Control  
in Critically Ill Patients

The NICE-SUGAR Study Investigators\*



*The* **NEW ENGLAND JOURNAL of MEDICINE**

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**ORIGINAL ARTICLE**

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Hypoglycemia and Risk of Death  
in Critically Ill Patients

The NICE-SUGAR Study Investigators\*

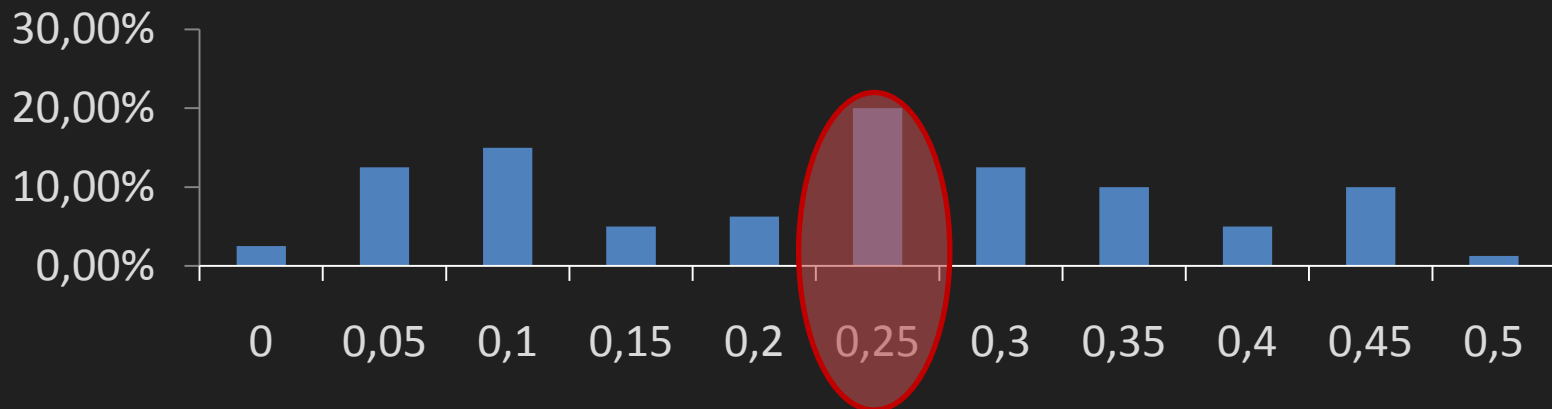


# 7. Opomíjet limitace klinických studií

Představte si phase 3 studii  
naradrenalinu ...

... pacient se sepsí indukovanou  
refrakterní hypotenzí (STK pod 90mmHg  
po podání 1000ml) dostane  
0,25ug/kg/min testované látky...

Bude tohle fungovat ?



# The NEW ENGLAND JOURNAL of MEDICINE

ESTABLISHED IN 1812

APRIL 24, 2014

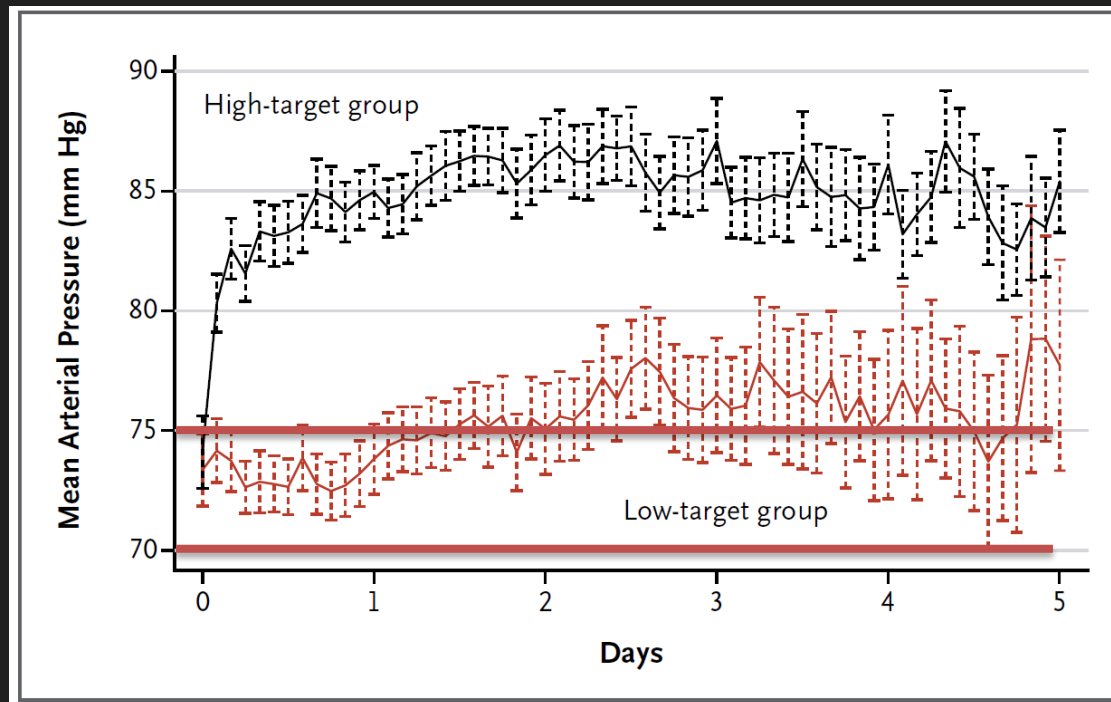
VOL. 370 NO. 17

## High versus Low Blood-Pressure Target in Patients with Septic Shock

Pierre Asfar, M.D., Ph.D., Ferhat Meziani, M.D., Ph.D., Jean-François Hamel, M.D., Fabien Grelon, M.D.,

### CONCLUSIONS

Targeting a mean arterial pressure of 80 to 85 mm Hg, as compared with 65 to 70 mm Hg, in patients with septic shock undergoing resuscitation did not result in significant differences in mortality at either 28 or 90 days. (Funded by the French Ministry of Health; SEPSISPAM ClinicalTrials.gov number, NCT01149278.)



# The NEW ENGLAND JOURNAL of MEDICINE

ESTABLISHED IN 1812

APRIL 24, 2014

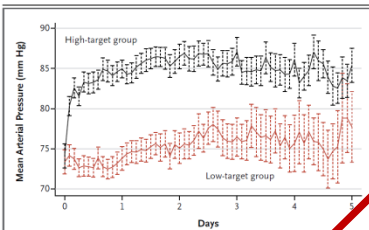
VOL. 370 NO. 17

## High versus Low Blood-Pressure Target in Patients with Septic Shock

Pierre Asfar, M.D., Ph.D., Ferhat Meziani, M.D., Ph.D., Jean-François Hamel, M.D., Fabien Grelon, M.D.,

### Secondary outcomes — no./total no. (%)

Death at day 90†	164 (42.3)	170 (43.8)	0.74
Survival at day 28 without organ support‡	241 (62.1)	235 (60.6)	0.66
Doubling of plasma creatinine	161 (41.5)	150 (38.7)	0.42
No chronic hypertension	71/215 (33.0)	85/221 (38.5)	0.32
Chronic hypertension	90/173 (52.0)	65/167 (38.9)	0.02
Renal-replacement therapy from day 1 to day 7	139 (35.8)	130 (33.5)	0.50
No chronic hypertension	66/215 (30.7)	77/221 (34.8)	0.36
Chronic hypertension	73/173 (42.2)	53/167 (31.7)	0.046



**Figure 2. Mean Arterial Pressure during the 5-Day Study Period.**

Mean arterial pressures were significantly lower in the low-target group than in the high-target group during the 5 protocol-specified days (P=0.02 by repeated-measures regression analysis), although the values exceeded the target values of 80 to 85 mm Hg in the high-target group and 65 to 70 mm Hg in the low-target group. The I bars represent 95% confidence intervals.

fluid balance, and the fluid balance was lower than those reported previously,<sup>7,8</sup> possibly because our population of patients differed from those in previous studies or because of more restrictive protocols for fluid administration in our study. In addition, there were no significant between-group differences in the overall rates of organ dysfunction or death at 90 days. However, in patients with a history of chronic arterial hypertension, targeting a mean arterial pressure of 80 to 85 mm Hg reduced both the incidence of a doubling of the blood creatinine level and the rate of renal-replacement therapy. There was no significant between-group difference in the overall rate of serious adverse events, but patients in the high-target group had significantly more episodes of atrial fibrillation.

No differences in the primary and secondary outcomes were observed between the two groups. Our study was prospectively powered to detect an absolute difference of 10 percentage points in the rate of death on the basis of an expected rate of 45% in the low-target group, at an alpha level of 0.05 and a beta level of 0.20, with the use of a two-tailed test. The expected overall death rate in our study was consistent with the rates among patients with septic shock that were reported in previous multicenter trials (37%,<sup>5</sup> 39%,<sup>8</sup> 47%,<sup>4</sup> and 49%<sup>6</sup>) at the time the

trial was designed. The absolute reduction of 10 percentage points in mortality was not reported in the literature when the patients in our study because the patients in the literature when the patients in 2008 had tested the high-target group. The high-target group had reductions of 7 percentage points, and 10 percent of death, two other trials after we started recruiting patients. The hypothesis of an absolute difference of 10 percentage points<sup>5,15</sup> and 10 percentage points. Hence, the anticipated risk reduction in our study was close to the risk reduction in previous studies. However, the rate of death at 28 days was lower than in other studies, although it was similar to the rate in more recent trials, ranging from 25 to 57%.<sup>16</sup> Nevertheless, the lower-target group had a lower rate of death led to an underpowered study. We may not have detected a difference in the incidence of some adverse events such as myocardial infarction.

Septic shock is a major cause of death in our study. Septic shock was significantly more common in the high-target group than in the low-target group. An adverse effect might be that the high-target group had significantly higher doses of catecholamine and the longer duration of catecholamine infusions in the high-target group. However, given the small number of episodes of atrial fibrillation, other confounding factors cannot be ruled out. The association between atrial fibrillation and septic shock should be considered only as a hypothesis-generating concept for future trials.

At randomization, according to the protocol, the high-target group had significantly more episodes of atrial fibrillation, which was reported in previous studies.<sup>18</sup> Atrial fibrillation, a common complication in patients with septic shock, means that the patient could have a lower rate of perfusion. The overall rates of atrial fibrillation were similar between the two groups. The rates of atrial fibrillation were similar between those with septic shock and those with other causes of shock.

organ dysfunction or death at 90 days. However, in patients with a history of chronic arterial hypertension, targeting a mean arterial pressure of 80 to 85 mm Hg reduced both the incidence of a doubling of the blood creatinine level and the rate of renal-replacement therapy. There was no

## Studie

## Relativní riziko

CHEST

1,21 (1,00-1,45)

6 S

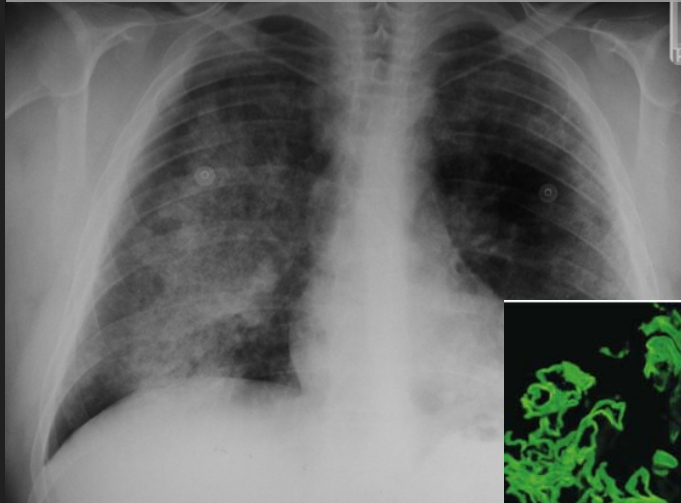
1,28 (0,96-1,72)

SEPSISPAM

1,23 (0,91-1,67)

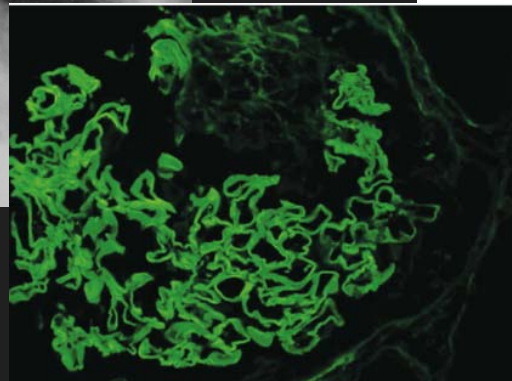
# 8. Nemyslet si, že už vím o pacientově stavu všechno

**PNEUMONIE**



**S-AKI**

I've let you down.

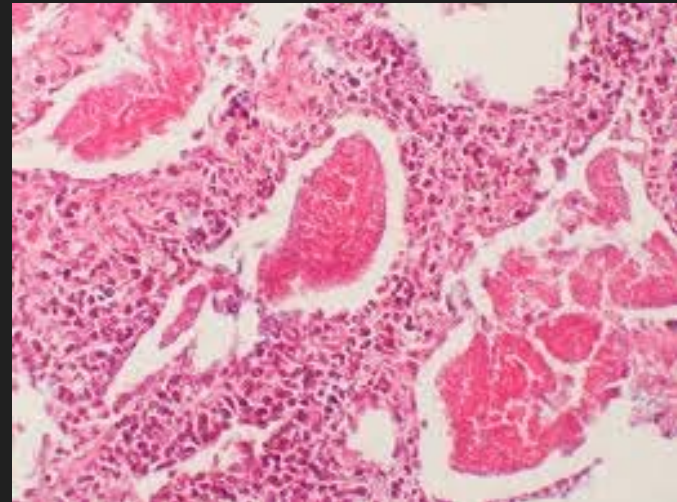


**GOODPASTURE SY.**

# REZISTENTNÍ TBC



**HIV/AIDS**



**PNEUMOCYSTIS**

# „EXOTICKÁ“ PNEUMONIE

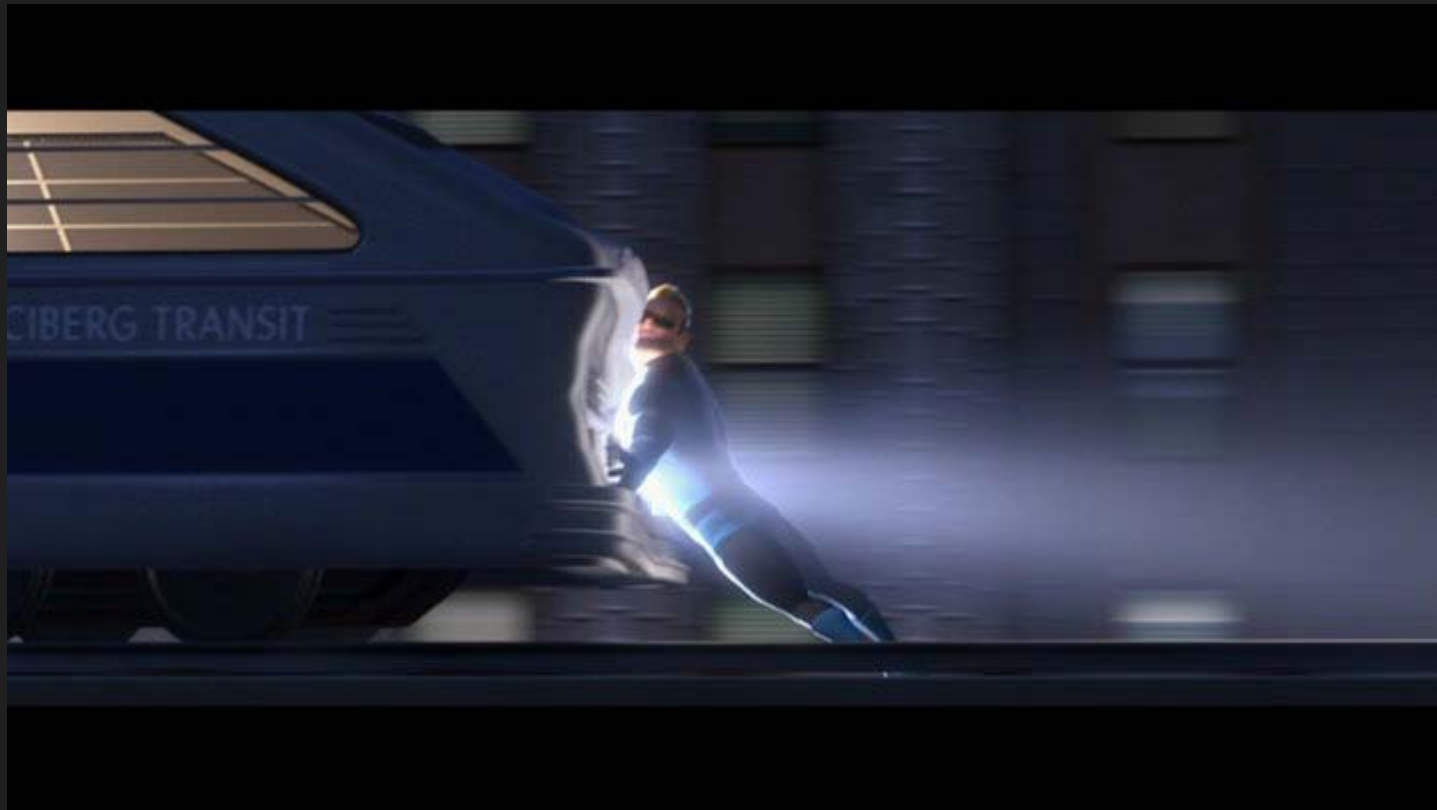


**MYXEDEM**



**LEGIONELLA**

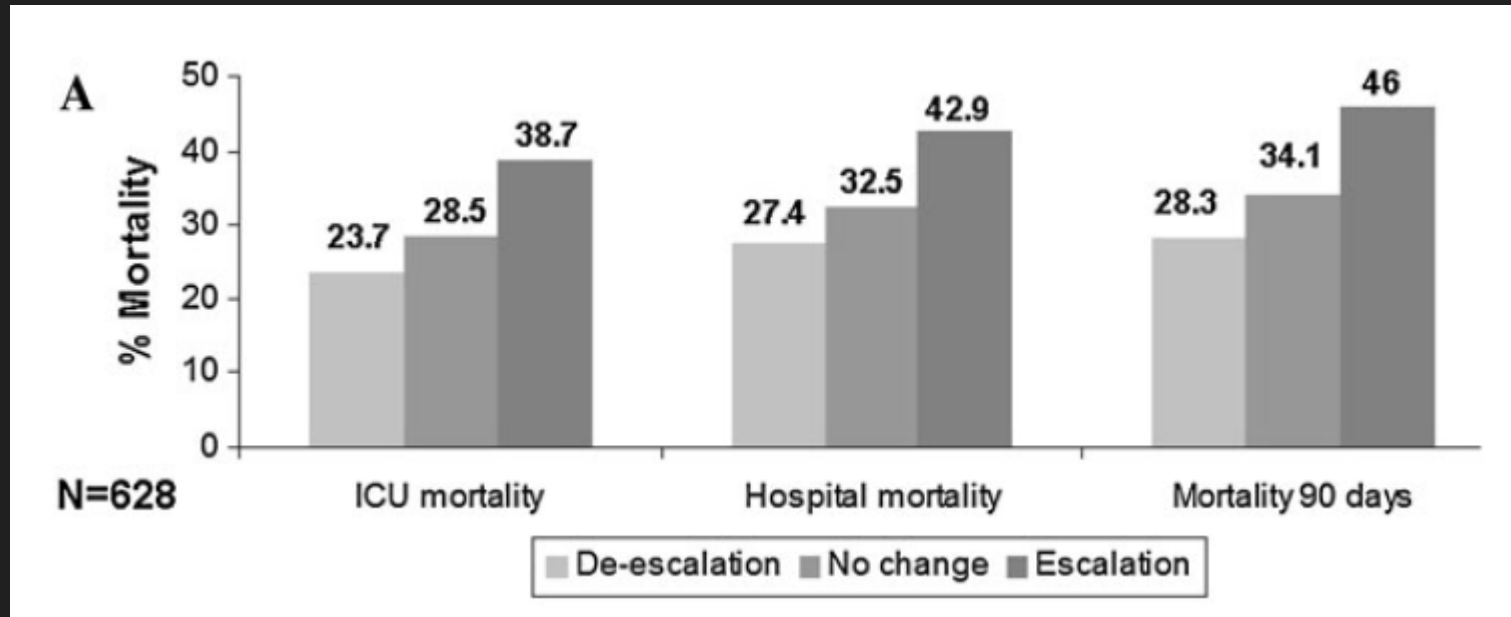
# 9. Neopomenout de-eskalovat





J. Garnacho-Montero  
A. Gutiérrez-Pizarra  
A. Escoreca-Ortega  
Y. Corcia-Palomo  
Esperanza Fernández-Delgado  
I. Herrera-Melero  
C. Ortiz-Leyba  
J. A. Márquez-Vácaro

## De-escalation of empirical therapy is associated with lower mortality in patients with severe sepsis and septic shock



# The Importance of Fluid Management in Acute Lung Injury Secondary to Septic Shock

CHEST / 136 / 1 / JULY, 2009

Claire V. Murphy, PharmD; Garrett E. Schramm, PharmD;  
Joshua A. Doherty, BS; Richard M. Reichley, RPh; Ognjen Gajic, MD, FCCP;  
Bekele Afessa, MD, FCCP; Scott T. Micek, PharmD; and  
Marin H. Kollef, MD, FCCP

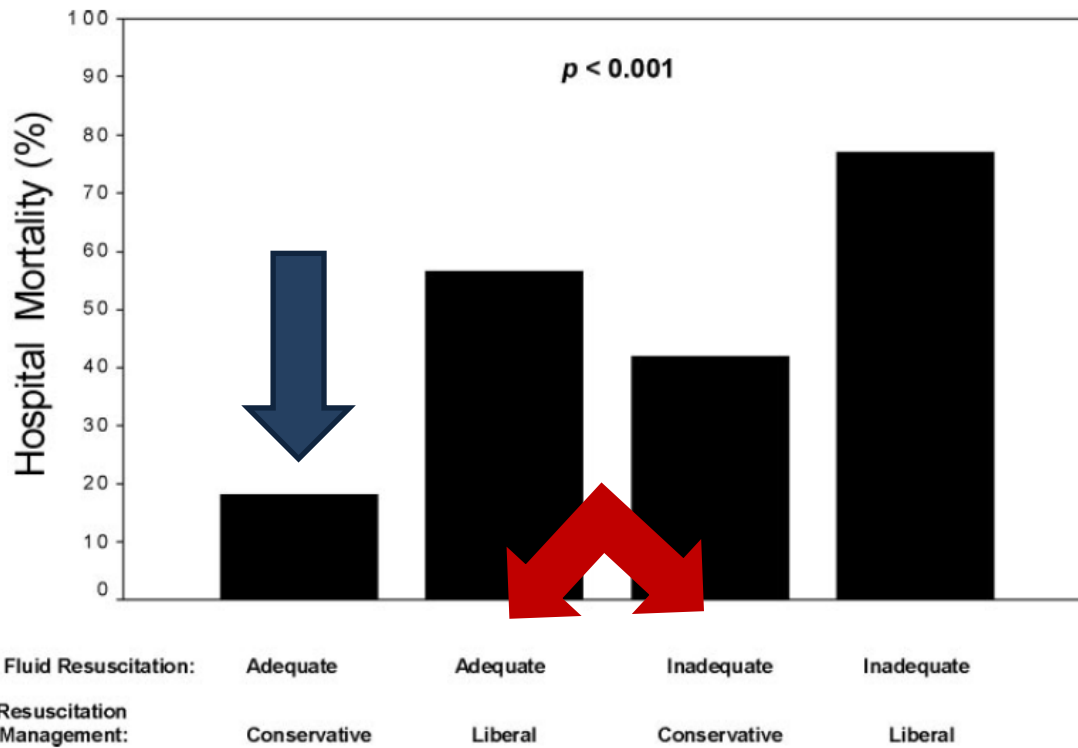


FIGURE 3. Hospital mortality according to whether or not patients achieved AIFR, CLFM, both, or neither.

# 10. „... někdy už to vzdát ...“

## **DOPORUČENÍ PŘEDSTAVENSTVA ČLK č. 1/2010**

**k postupu při rozhodování o změně léčby intenzivní na léčbu paliativní u pacientů v terminálním stavu, kteří nejsou schopni vyjádřit svou vůli**

- c) Zahájení nebo pokračování jakéhokoliv léčebného postupu, který není odborně odůvodněný, kde neexistuje racionální předpoklad jeho příznivého účinku na celkový průběh onemocnění a kde rizika komplikací, strádání, útrap a bolesti převažují nad reálným klinickým přínosem zvoleného postupu, je v rozporu s etickými principy medicíny a Chartou práv umírajících. Neexistuje povinnost zahajovat marnou a neúčelnou léčbu nebo v ní pokračovat, pokud je probíhající léčba odůvodněně za takovou označena.
- d) Zabezpečení fyzických, psychických, sociálních a duchovních potřeb pacienta, odstranění pocitů bolesti, strádání a utrpení s respektováním lidské důstojnosti jsou základní priority paliativní péče.



UDA  
Ultra Digital Animation

# 10 times NEVER AGAIN

1. Nevzdávat to
2. Neztrácet čas
3. Nebát se počáteční racionální terapeutické agresivity
4. ... a její ekonomické / personální náročnosti
5. Nedělat léčebnou kosmetiku
6. Nepodléhat módním trendům
7. Opomíjet limitace klinických studií
8. Nemyslet si, že už vím o pacientově stavu všechno
9. Neopomenout de-eskalovat
10. „... někdy už to vzdát ...“

Robert

**Fulghum**

VŠECHNO, CO

OPRAVDU

POTŘEBUJU

ZNÁT.

JSEM SE

NAUČIL

V MATEŘSKÉ

ŠKOLCE



NEOPOMENOUT  
SI NA CESTU  
PŘIBALIT  
ZDRAVÝ SELSKÝ  
ROZUM A  
TROCHU  
ZNALOSTÍ

**DĚKUJI  
ZA  
POZORNOST**

**Práce byla podpořena  
VZ MSM0021620819 a projektem P36 „PRVOUK“**

**ATB**

**OBJEM**

**ECHO**

**NA**







ORIGINAL ARTICLE

# High-Frequency Oscillation in Early Acute Respiratory Distress Syndrome

Niall D. Ferguson, M.D., Deborah J. Cook, M.D., Gordon H. Guyatt, M.D.,

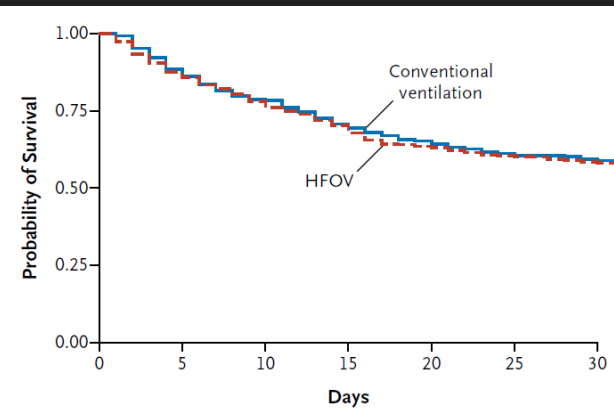
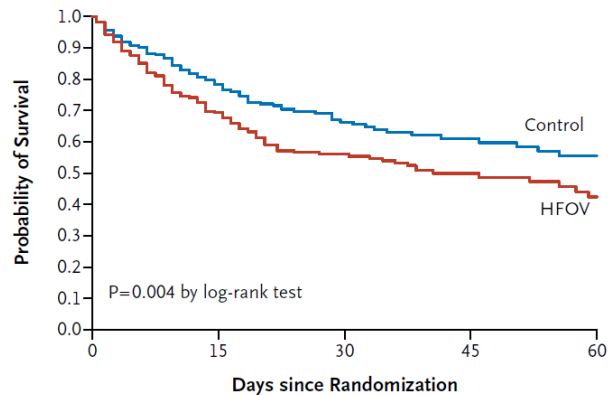
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# High-Frequency Oscillation for Acute Respiratory Distress Syndrome

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## High-Frequency Oscillation in Early Acute Respiratory Distress Syndrome

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In conclusion, in a large effectiveness study, we were unable to find any benefit or harm from the use of HFOV in adult patients with ARDS. We recommend that this mode of ventilation not be used for routine care.