



MODALITÉS DE L'ÉPURATION EXTRARÉNALE DANS LE CHOC SEPTIQUE

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Service de Réanimation Médicale

CHU Ibn El Jazzar Kairouan

ATR Novembre 2024

PHYSIOPATHOLOGIE

Pathogène

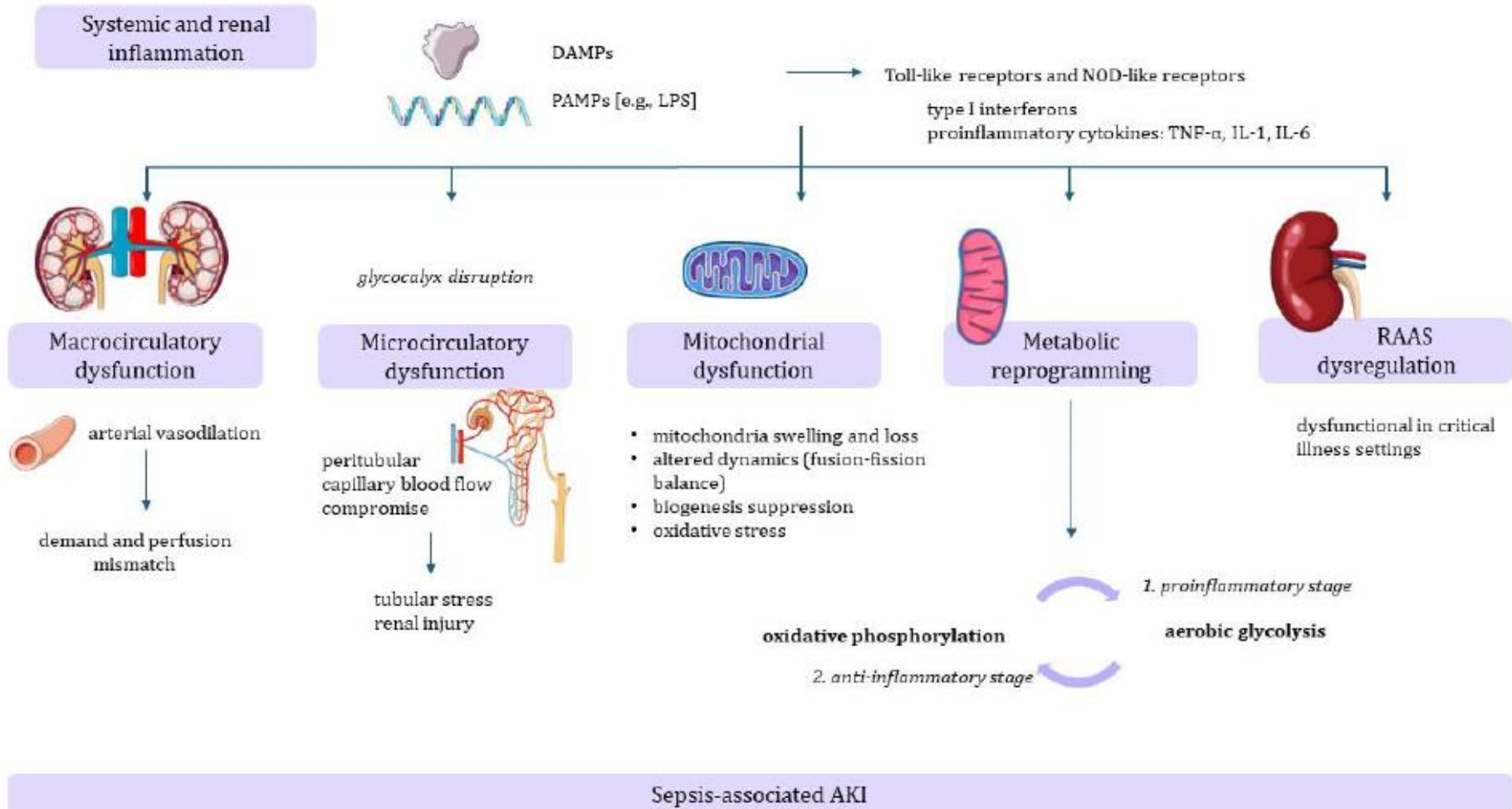
Infection

**Orage
cytokinique**

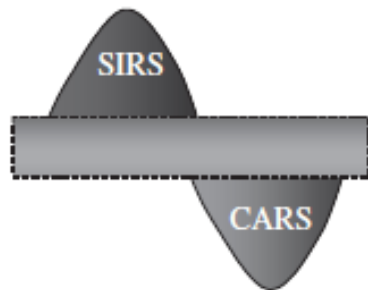
**Défaillance
Multiviscérale**

Décès

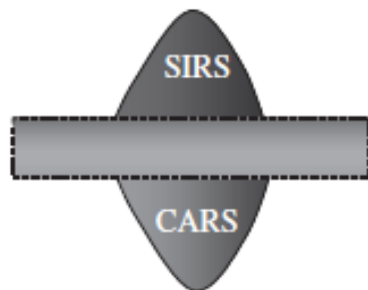
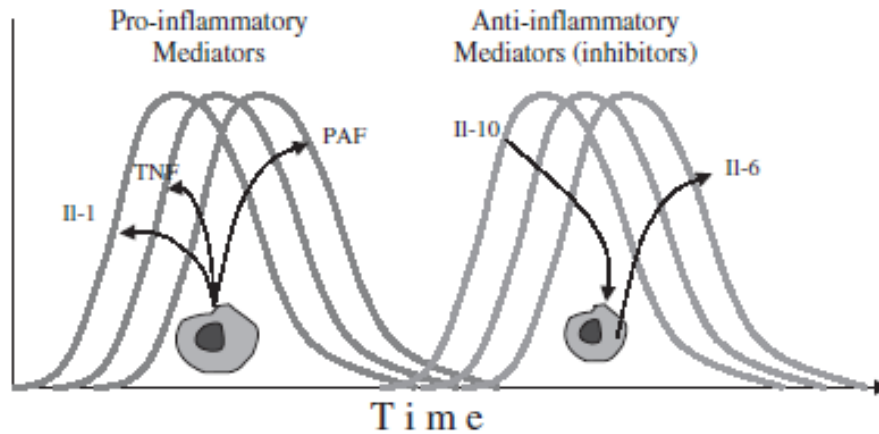
PHYSIOPATHOLOGIE SA-AKI



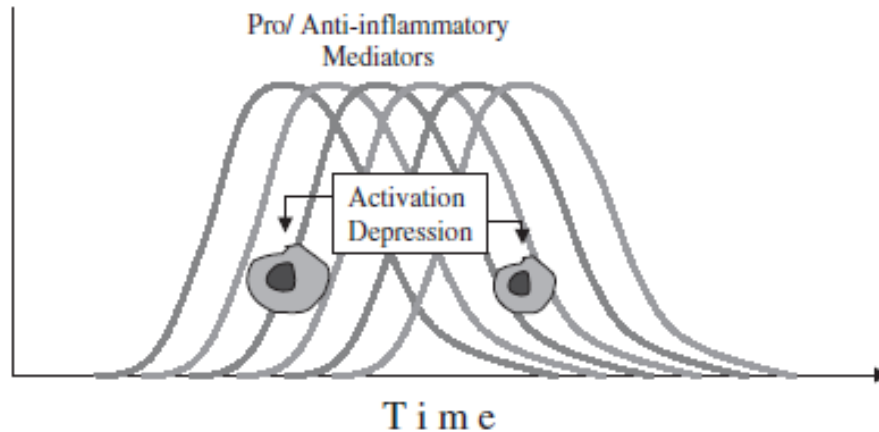
PHYSIOPATHOLOGIE CYTOKINE STORM



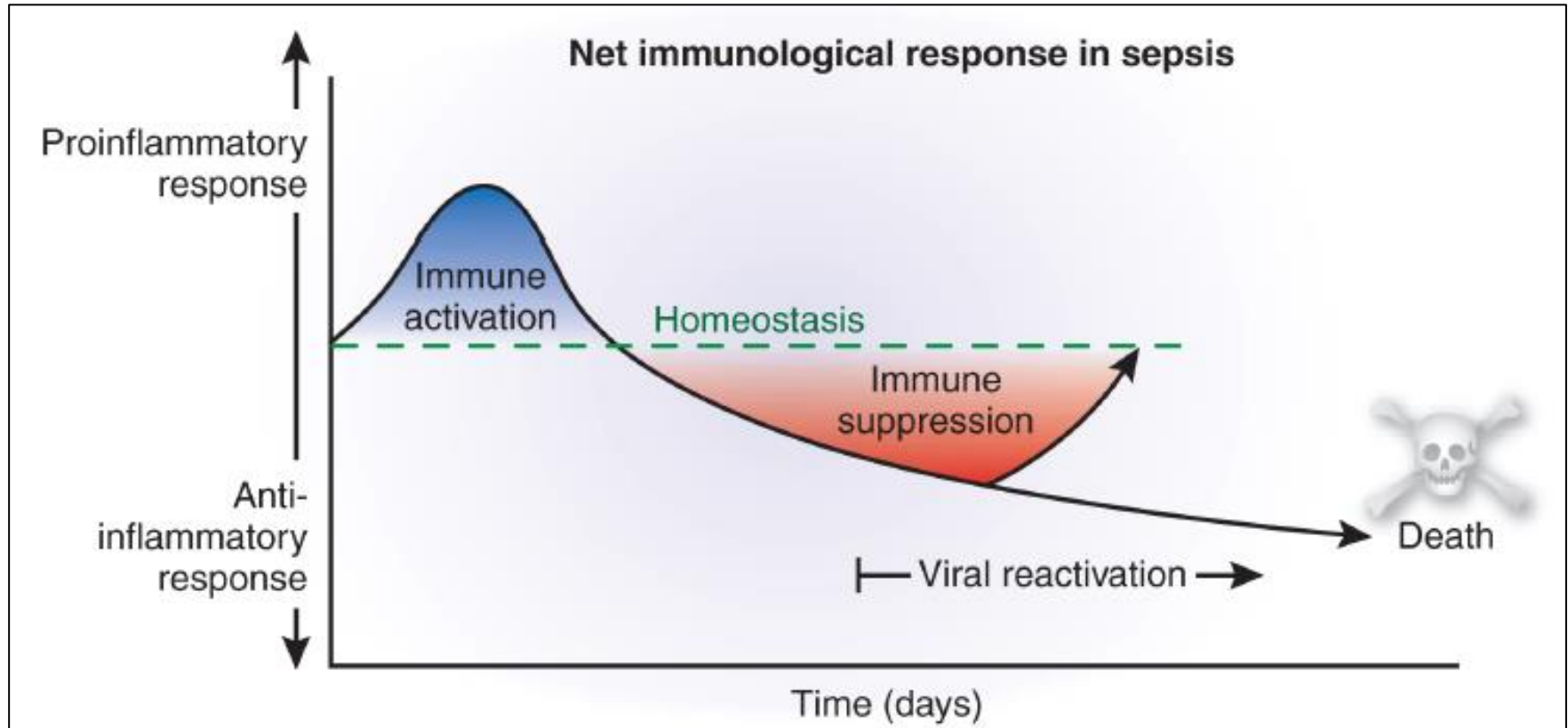
Mediators' levels (Arbitr. Units)



Mediators' levels (Arbitr. Units)

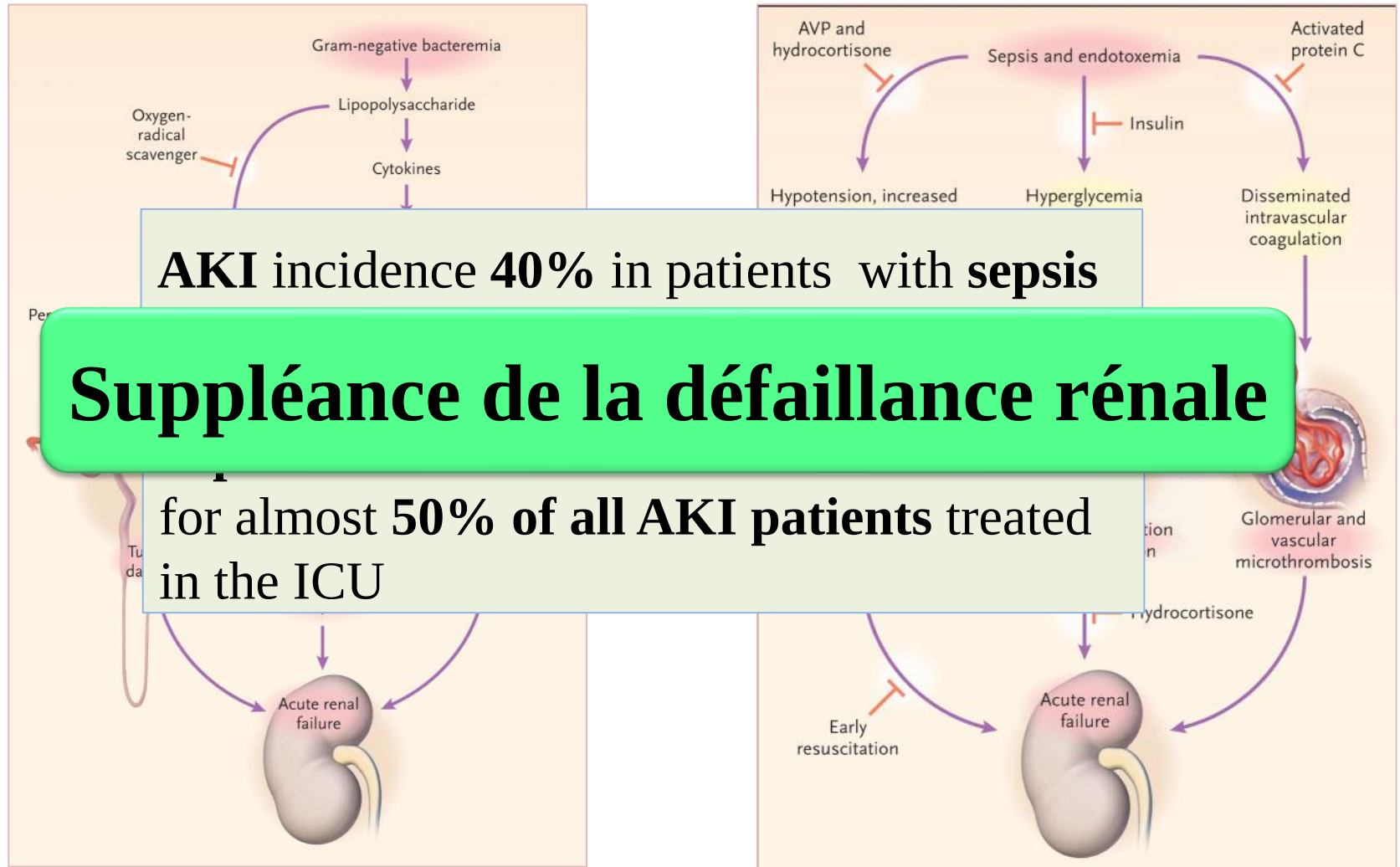


PHYSIOPATHOLOGIE CYTOKINE STORM

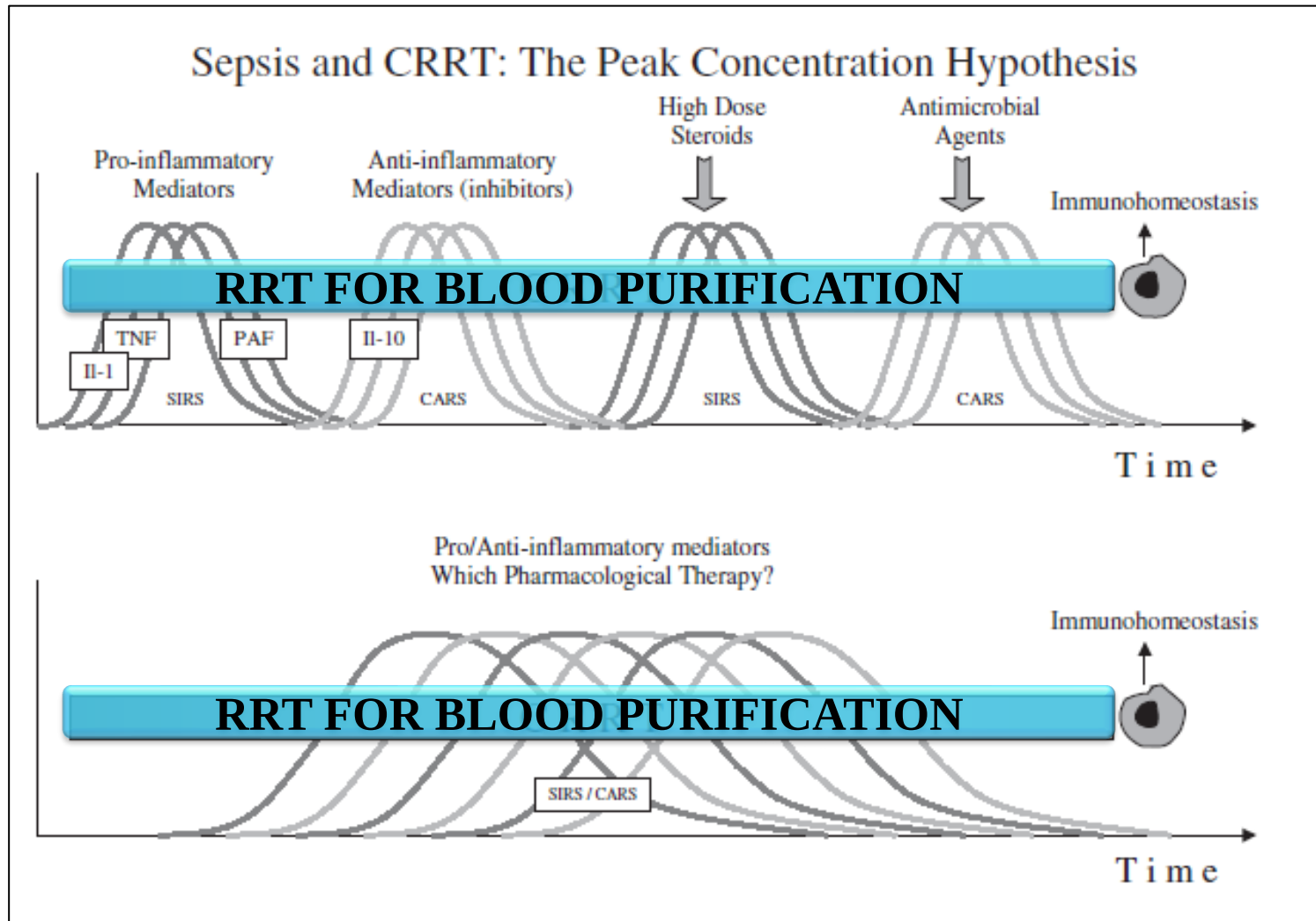


INTÉRÊT DE L'ÉPURATION EXTRA-RÉNALE AU COURS DU CHOC SEPTIQUE ?

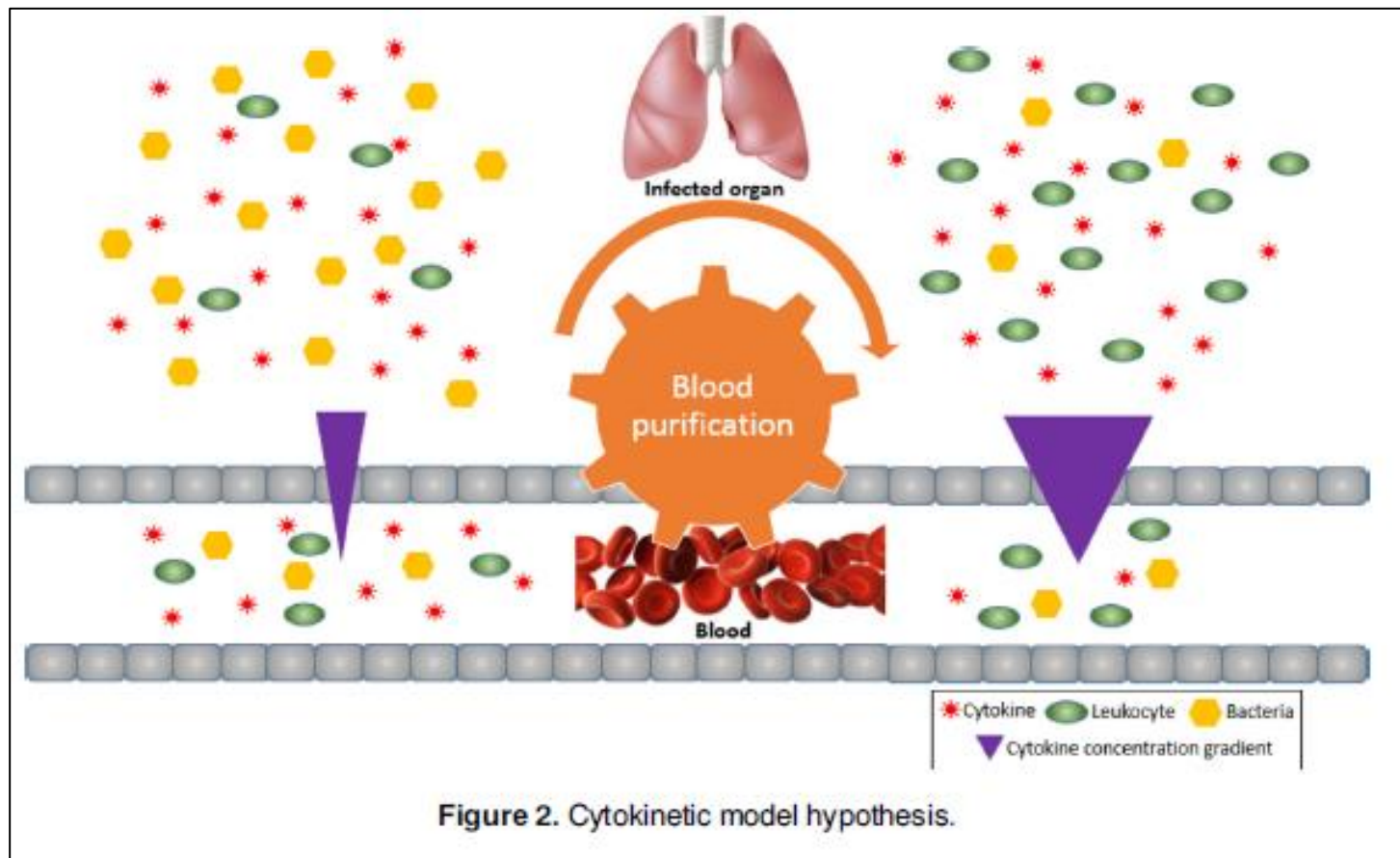
RRT FOR SA-AKI



RRT FOR BLOOD PURIFICATION



RRT FOR BLOOD PURIFICATION



**EER POUR SUPPLÉANCE DE LA
DÉFAILLANCE RENALE ASSOCIÉE
AU SEPSIS**

SA-AKI

RRT FOR SA-AKI / QUELLE MODALITÉ ??

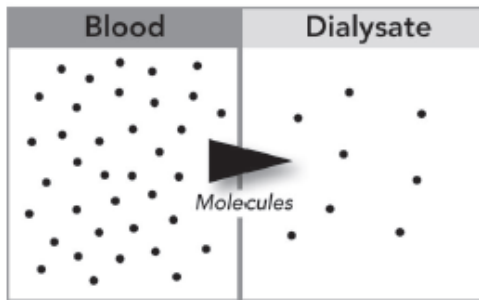
**Modalité : Continue vs intermittente?
Convection vs Diffusion ?**

Timing précoce ou tardive ?

Dose de dialyse?

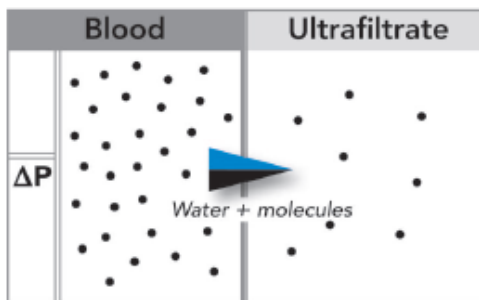
EER / RAPPEL THEORIQUE

A Diffusion



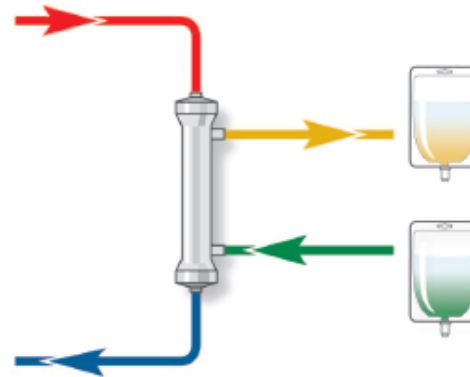
Concentration gradient

B Convection

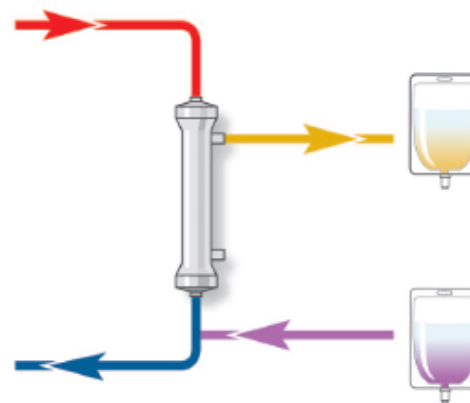


Hydrostatic pressure gradient

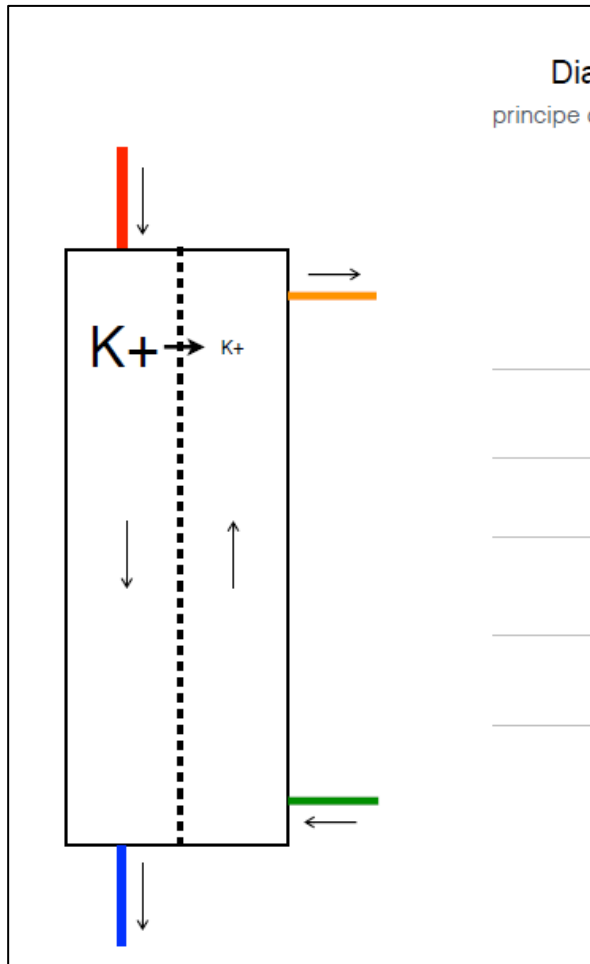
C Hemodialysis circuit



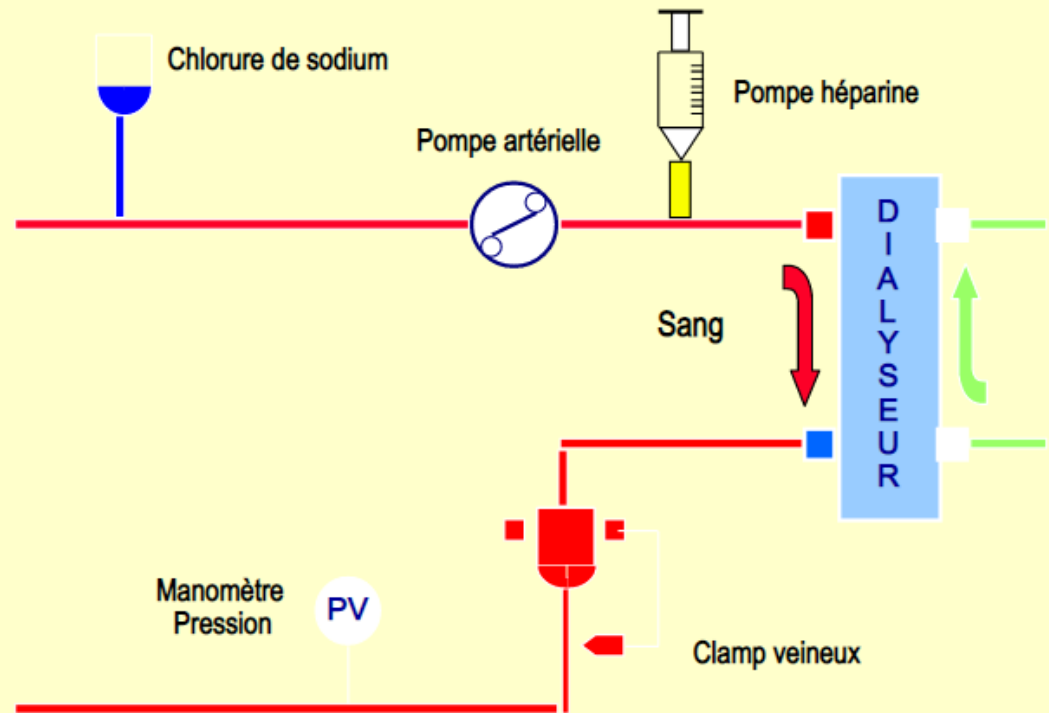
D Hemofiltration circuit



EER / RAPPEL THEORIQUE

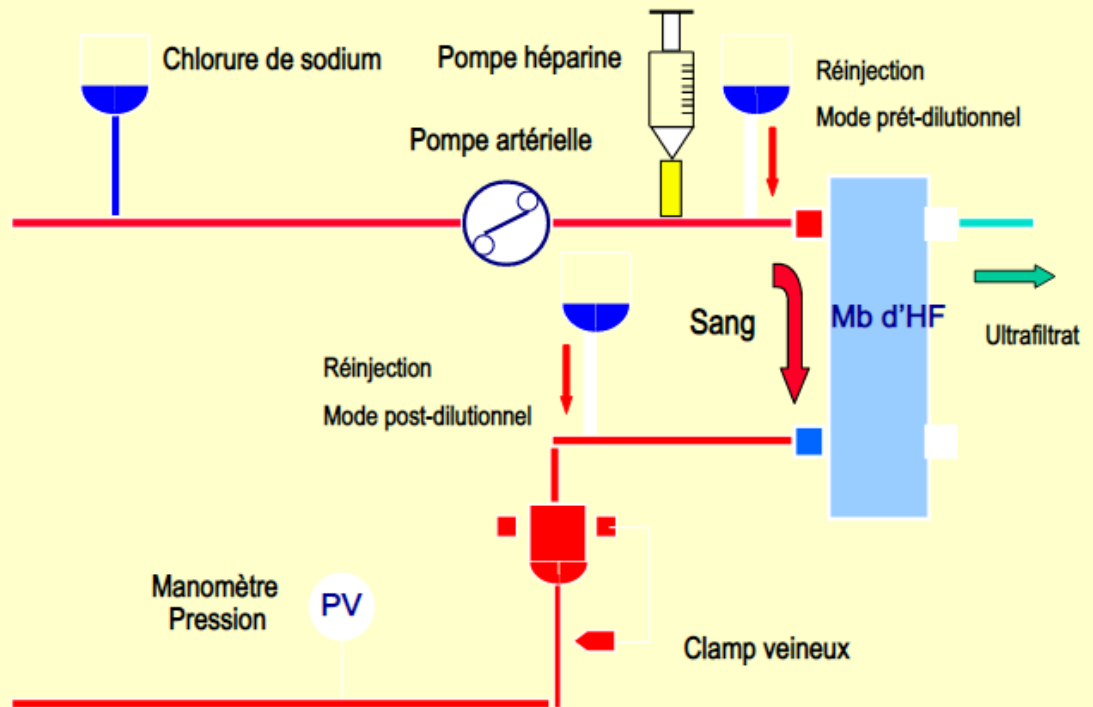
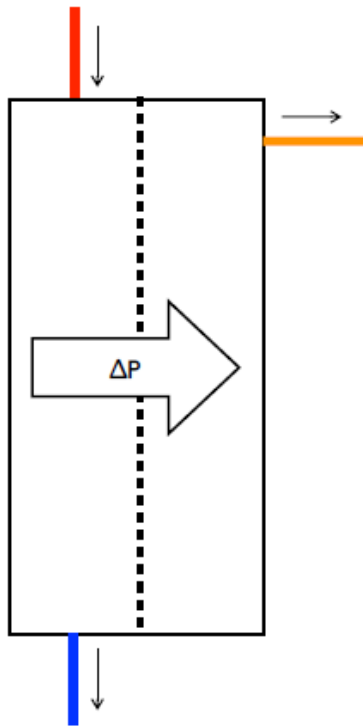


TECHNIQUE HEMODIALYSE



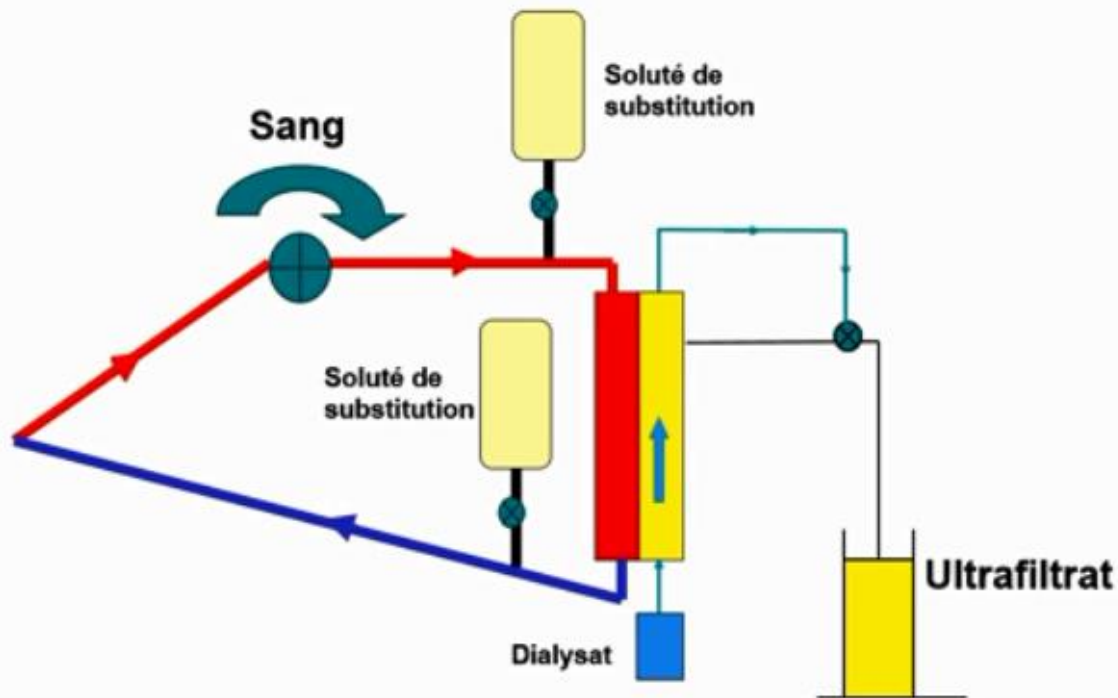
EER / RAPPEL THEORIQUE

TECHNIQUE HEMOFILTRATION



EER / RAPPEL THEORIQUE

TECHNIQUE HEMODIAFILTRATION



Continuous venovenous haemodiafiltration versus intermittent haemodialysis for acute renal failure in patients with multiple-organ dysfunction syndrome: a multicentre randomised trial

Lancet 2006; 368: 379–85

Jean-Louis Pallot,

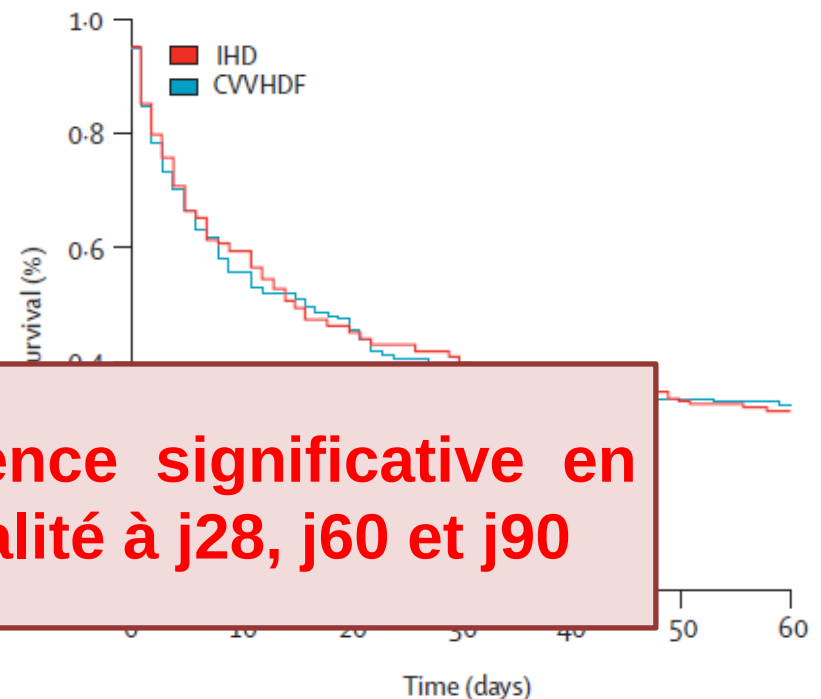
	Intermittent haemodialysis (n=184)	Continuous venovenous haemodiafiltration (n=175)
Age (years)	65 (63–67)	65 (63–67)
Weight (kg)	79 (76–82)	79 (76–82)
Sex		
Male	132 (72%)	132 (75%)
Female	52 (28%)	43 (25%)
Reason for admission		
Medical	134 (73%)	134 (76%)
Surgical	50 (27%)	41 (24%)
Previous health status		
No or moderate limitation	106 (58%)	106 (60%)
Serious limitation or bedridden	78 (42%)	69 (40%)
SAPS II score	64 (62–66)	64 (62–66)
LOD score	10 (9–11)	10 (9–11)
Catecholamine	158 (86%)	155 (89%)
Mechanical ventilation	174 (95%)	171 (98%)
Delayed ARF	123 (67%)	123 (70%)
Median time from admission to inclusion (days)	2	3
Oliguria	101 (55%)	107 (61%)
Presence of sepsis	127 (69%)	98 (56%)
Urea (mmol/L)	31 (29–33)	29 (26–31)
Serum creatinine (μmol/L)	432 (407–457)	422 (381–464)

	Intermittent haemodialysis	Continuous venovenous haemodiafiltration
Duration of sessions (h)	5.2 (5.1–5.3)	continuous
Blood flow (mL per min)	278 (275–281)	146 (145–147)
Dialysate flow*	500	1099 (1068–1128)
Ultrafiltration flow (mL per h)		1278 (1255–1301)
Net ultrafiltration† (mL per day)	2213 (2141–2285)	2107 (2011–2203)
Mean urea (mmol/L)	15.7 (7.5)	14.8 (9.1)

Continuous venovenous haemodiafiltration versus intermittent haemodialysis for acute renal failure in patients with multiple-organ dysfunction syndrome: a multicentre randomised trial

Lancet 2006; 368: 379–85

Christophe Vinsonneau, Christophe Camus, Alain Combes, Marie Alyette Costa de Beauregard, Kada Klouche, Thierry Boulain, Jean-Louis Pallot, Jean-Daniel Chiche, Pierre Taupin, Paul Landais, Jean-François Dhainaut, for the H



Intermittent haemodialysis

Survival

Day 28

Day 60 (primary endpoint)

Day 90

Renal support duration (days)

Length of ICU stay (days)

Length of hospital stay (days)

Values are mean (95% CI). ICU=intensive-care unit.

Table 3: Outcomes according to treatment group

Numbers at risk

IHD	184	85	68	58
CVVHDF	175	83	62	57

Continuous venovenous haemodiafiltration versus intermittent haemodialysis for acute renal failure in patients with multiple-organ dysfunction syndrome: a multicentre randomised trial

Lancet 2006; 368: 379–85

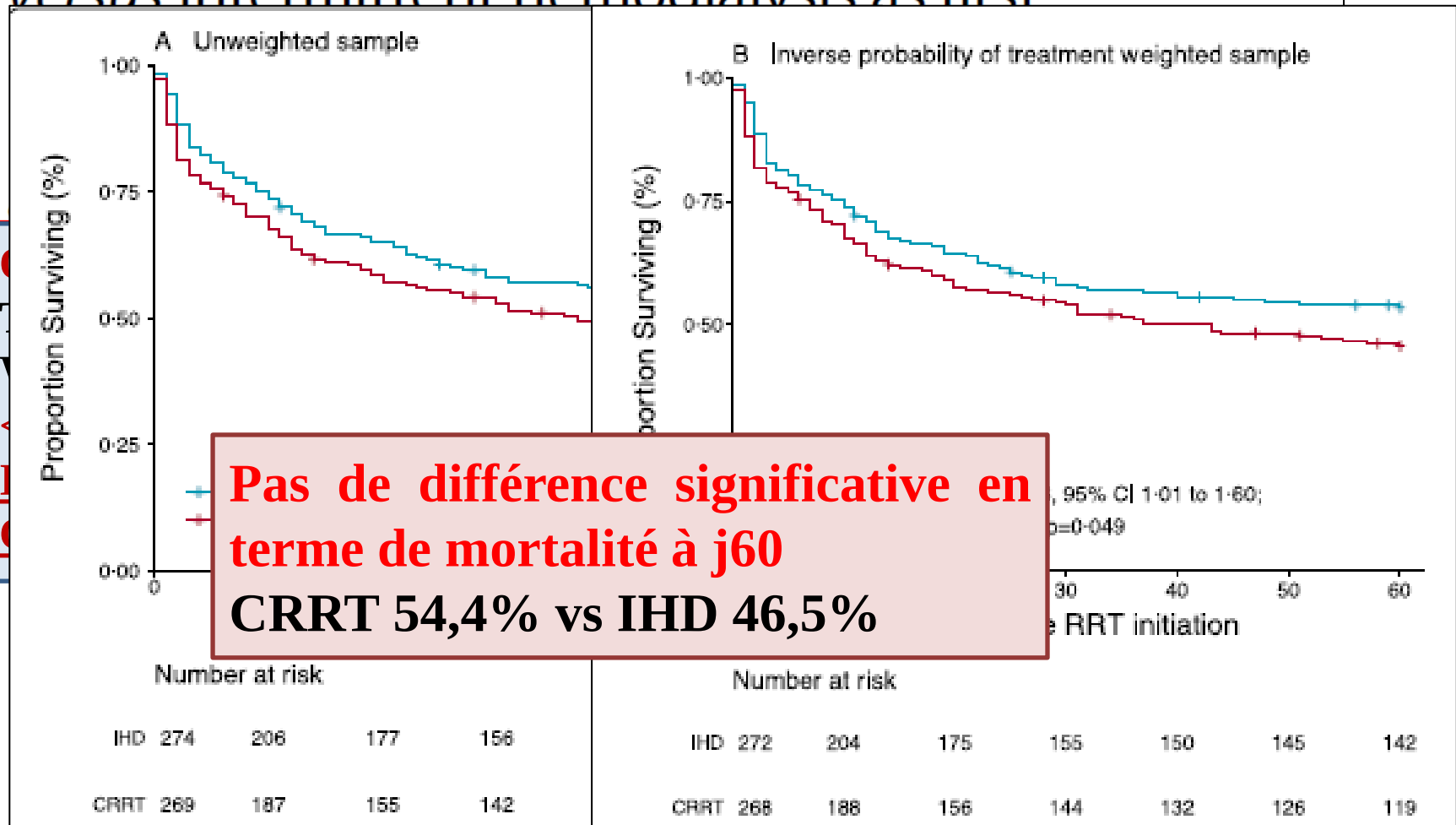
*Christophe Vinsonneau, Christophe Camus, Alain Combes, Marie Alyette Costa de Beauregard, Kada Klouche, Thierry Boulain, Jean-Louis Pallot, Jean-Daniel Chiche, Pierre Taupin, Paul Landais, Jean-François Dhainaut, for the Hemodiafe Study Group**

	Intermittent haemodialysis (n=184)	Continuous venovenous haemofiltration (n=175)	p value
Hypotension*	72 (39%)	61 (35%)	0.47
Bleeding event†	13 (7%)	12 (7%)	0.89
Thrombocytopenia	22 (12%)	12 (7%)	
Hypoglycaemia	12 (7%)	12 (7%)	
Hypophosphataemia	13 (7%)	12 (7%)	
Hypothermia	10 (5%)	12 (7%)	
Arrhythmia	18 (10%)	9 (5%)	0.15
Catheter infection	2 (1%)	3 (2%)	0.95

Pas de différence significative en terme de tolérance hémodynamique



Continuous renal replacement therapy *versus* intermittent hemodialysis as first



Initiation of continuous renal replacement therapy versus intermittent hemodialysis in critically ill patients with severe acute kidney injury: a secondary analysis of STARRT-AKI trial

Results: We identified 1590 trial participants who initially received CRRT and 606 who initially received IHD. The composite outcome of death or RRT dependence at 90-days occurred in 823 (51.8%) patients who commenced CRRT and 329 (54.1%) who commenced IHD. In multivariable adjusted analyses, initiation of CRRT, as compared to IHD, was associated with a significant reduction in the composite outcome of death or RRT dependence at 90-days (OR 0.61; 95% CI 0.39-0.94). In sensitivity analyses, initial receipt of CRRT was associated with a lower risk of the composite outcome at 90-days compared with initial receipt of IHD (OR 0.61; 95% CI 0.39-0.94), predominantly driven by a lower risk of RRT dependence at 90-days (OR 0.61; 95% CI 0.39-0.94).

Conclusions: In critically ill patients with severe AKI, initiation of CRRT, as compared to IHD, was associated with a significant reduction in the composite outcome of death or RRT dependence at 90-days.

LES RECOMMANDATIONS / TECHNIQUE

Réanimation
DOI 10.1007/s13546-014-0917-6

RÉFÉRENTIEL / *GUIDELINES*

2014

Les techniques d'EER, **continues et intermittentes**, peuvent être **utilisées indifféremment**, mais en tenant compte de **la disponibilité de la technique** et de l'expérience de l'équipe.

Accord fort

et de la Société francophone de dialyse (SFD)

Les techniques d'EER **diffusives ou convectives** peuvent être **utilisées indifféremment**, mais en tenant compte de **la disponibilité de la technique** et de l'expérience de l'équipe.

Accord fort

Venry, David Osman, Ly Van Vong

LES RECOMMANDATIONS

Intensive Care Med (2021) 47:1181–1247
<https://doi.org/10.1007/s00134-021-06506-y>

Intensive Care Med 2021

GUIDELINES

Surviving sepsis campaign: international guidelines for management of sepsis and septic shock 2021



In adults with sepsis or septic shock and AKI who require renal replacement therapy, we suggest using either continuous or intermittent renal replacement therapy
Weak recommendation, low quality of evidence

Theodore Iwashyna³³, Shevin Jacob³⁴, Ruth Kleinpell³⁵, Michael Klompas^{36,37}, Younsuck Koh³⁸, Anand Kumar³⁹, Arthur Kwizera⁴⁰, Suzana Lobo⁴¹, Henry Masur⁴², Steven McGloughlin⁴³, Sangeeta Mehta⁴⁴, Yatin Mehta⁴⁵,

Initiation Strategies for Renal-Replacement Therapy in the Intensive Care Unit

AKIKI NEJM 2016

Stéphane Gaudry, M.D., David Hajage, M.D., Frédérique

Table 1. Characteristics of the Patients at Baseline.*

Characteristic	Early Strategy (N = 311)	Delayed Strategy (N = 308)
Age — yr	64.8±14.2	67.4±13.4
Physiological support		
Invasive mechanical ventilation	266 (86)	267 (87)
Vasopressor support	265 (85)	263 (85)
Sepsis status — no. (%)		
Sepsis	25 (8)	21 (7)
Severe sepsis	16 (5)	19 (6)
Septic shock	209 (67)	204 (66)
Patients with oliguria or anuria — no. (%)	202 (65)	191 (62)
Serum creatinine — mg/dl	3.25±1.40	3.20±1.32
Blood urea nitrogen — mg/dl	53±24	54±24
Serum potassium — mmol/liter	4.4±0.7	4.4±0.7
Serum bicarbonate — mmol/liter	18.7±5.1	18.8±5.5

EARLY : KDIGO 3 , 4 h après
DELAYED : KDIGO 3 , 57 h
après
Techniques : HDI ou CCRT

Initiation Strategies for Renal-Replacement Therapy in the Intensive Care Unit

AKIKI NEJM 2016

Stéphane Gaudry, M.D., David Hajage, M.D., Frédérique

Laurent Martin-Lefevre, M.D., Bertrand Pons, M.D., Eric Boulet, M.D.

Alexandre Boyer, M.D., G

Dorothee Carpi

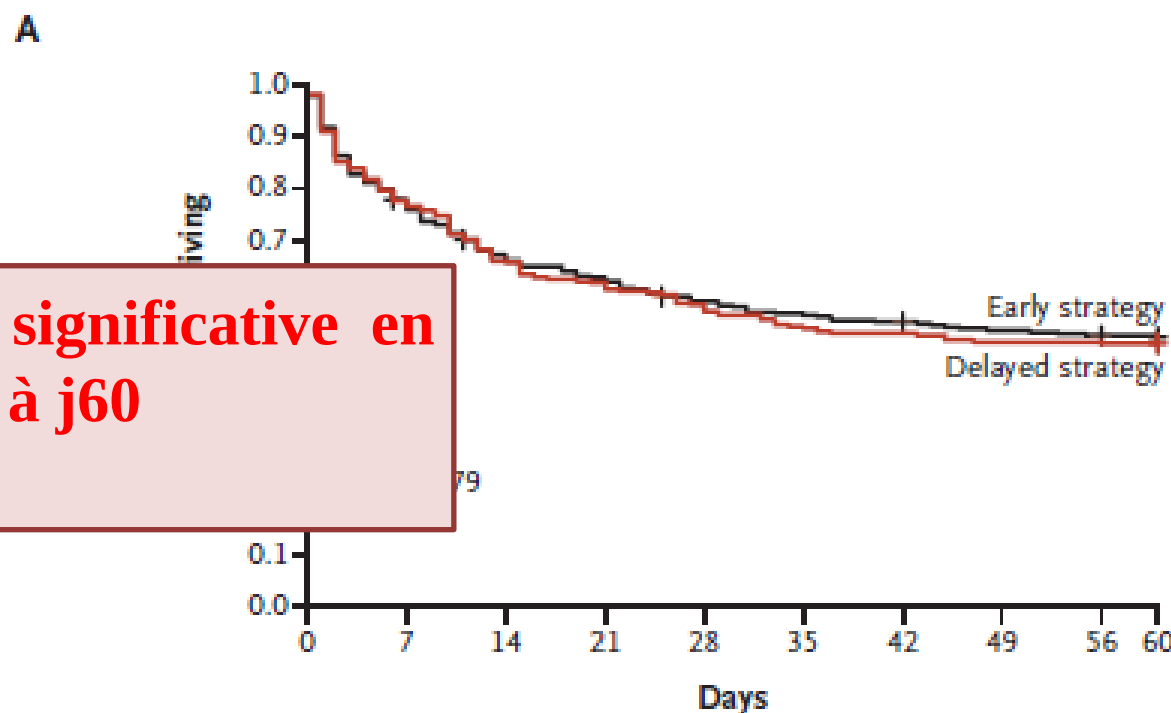
Alexandre Lautrette, I

Saad Nseir, M.D., Ph.D.,

Jean-Marie Forel, M.D.

**Pas de différence significative en
terme de mortalité à j60**

48,5% vs 49,7%



No. at Risk

Early strategy	311	241	207	194	179	172	167	161	158	157
Delayed strategy	308	239	204	191	178	165	161	156	156	155

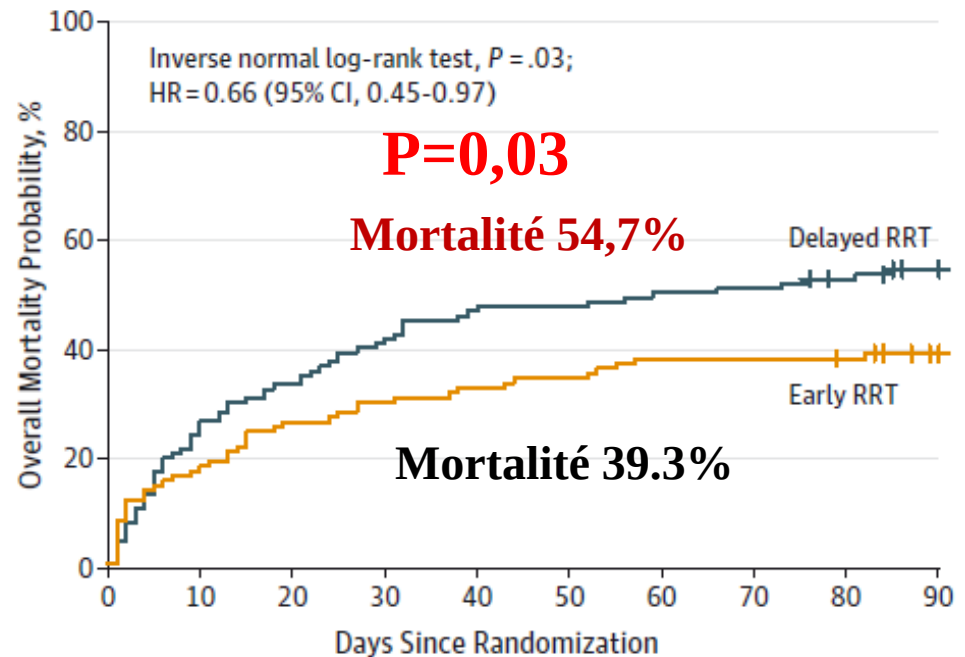
Effect of Early vs Delayed Initiation of Renal Replacement Therapy on Mortality in Critically Ill Patients With Acute Kidney Injury

The ELAIN Randomized Clinical Trial

ELAIN JAMA 2016

Alexander Zarbock, MD; John A. Kellum, MD; Christiaan A. Pinsky, MD; John D. Tobias, MD; Hermann Pavenstädt, MD; Andreea Boanta, MD; Joerg J. W. Uhl, MD

EARLY: 112 patients
KDIGO 2, 6 h après
DELAYED: 119 patients
KDIGO 3, 25 h après
TECHNIQUE: CVVHDF



No. at risk										
Early RRT	112	92	82	78	75	73	69	69	66	55
Delayed RRT	119	90	79	70	63	62	59	58	54	48

Effect of Early vs Delayed Initiation of Renal Replacement Therapy on Mortality in Critically Ill Patients With Acute Kidney Injury

The ELAIN Randomized Clinical Trial

ELAIN JAMA 2016

Alexander Zarbock, MD; John A. Kellum, MD; Christoph Schmidt, MD; Hugo Van Aken, MD; Carola Wempe, PhD; Hermann Pavenstädt, MD; Andreea Boanta, MD; Joachim Gerß, PhD; Melanie Meersch, MD

Table 3. Clinical Outcomes for Early vs Delayed Renal Replacement Therapy (RRT) Among Critically Ill Patients

	Absolute			Relative, %	OR or HR (95% CI)
Primary Outcome, No. (%)					
90-d All-cause mortality				-2.6)	HR: 0.66 (0.45 to 0.97)
Secondary Outcomes, No. (%)					
Recovery of renal function at day 90 ^e					
Yes	60 (53.6)	46 (38.7)	.02	14.9 (2.2 to 27.6)	OR: 0.55 (0.32 to 0.93) ^f
No ^g	52 (46.4)	73 (61.3)			

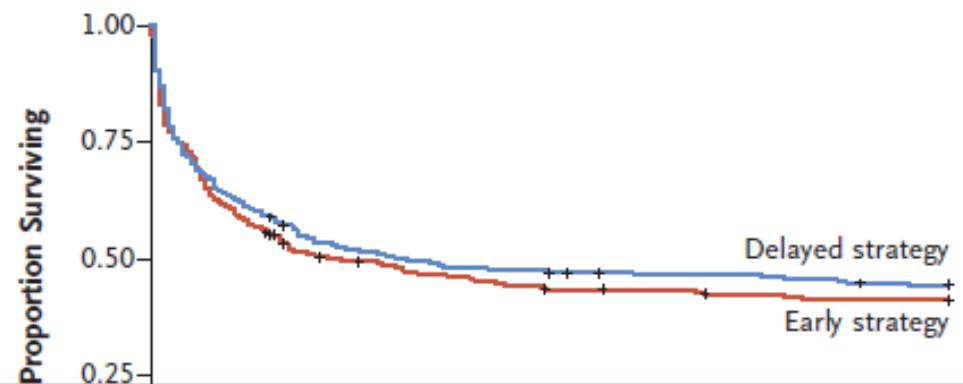
Meilleure récupération de la fonction rénale à j90 dans le groupe Précoce

Timing of Renal-Replacement Therapy in Patients with Acute Kidney Injury and Sepsis

IDEAL ICU NEJM 2018

S.D. Barbar, R. Clere-Jehl, A. Bourredjem, R. He
C. Lebert, J. Bohé, J. Badie, J.-P. Eraldi, J.-P. Rigaud, B. Levy, S. Siami,
G. Louis, L. Bouadma, J.-M. Constantin, E. Mercier, K. Klouche, D. du Cheyron,
G. Piton, D. Annane, S. Leber, T. van der Linden, C. Blesse, J. P. Mira,
C. Schwebel, L. Chimot, P. B
B. Louart, R. Trusson
for the IDEAL-ICU Trial Inve

EARLY: 246 patients
KDIGO 3, 12 h après
DELAYED: 242 patients
indication clinique ou
après 48 h si non
résolution de l'IRA



Pas de différence significative en terme de mortalité à j90

58% vs 54%

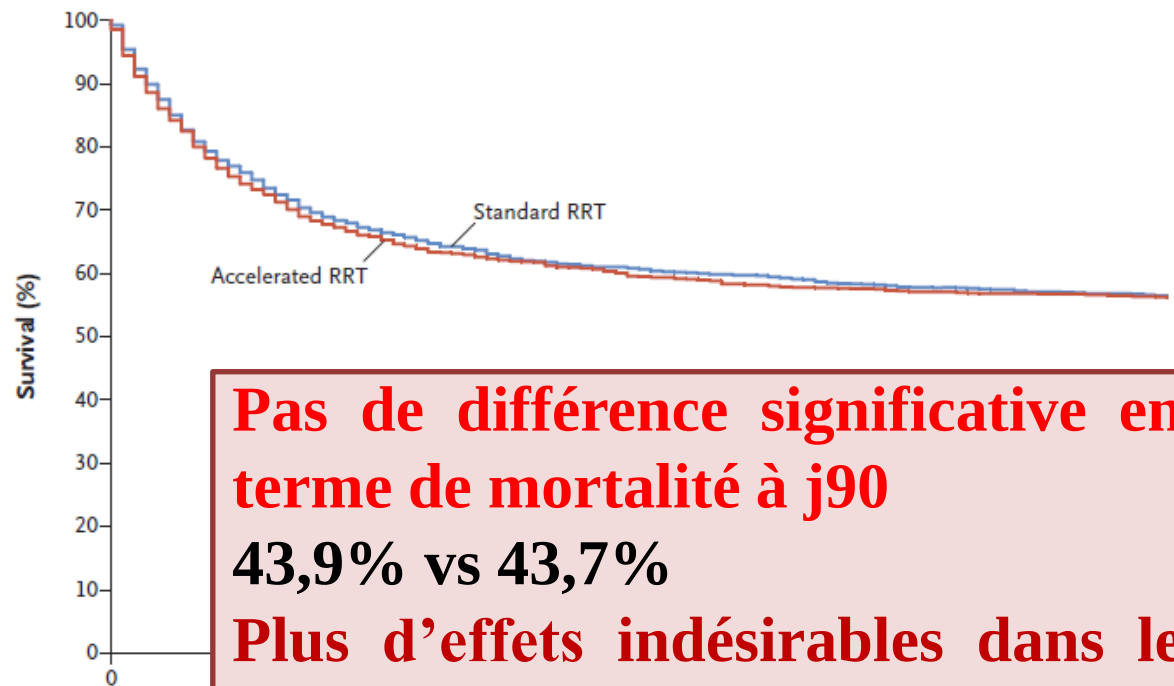
No. at R							
Delayed strategy	242	137	117	112	107	105	101
Early strategy	246	127	109	99	98	92	92

Timing of Initiation of Renal-Replacement Therapy in Acute Kidney Injury

STARRT AKI NEJM 2020

The STARRT-AKI Investigators, for the Canadian Society of Intensive Care, the Australian and New Zealand Intensive Care Society Clinical Trials Group, the United Kingdom Intensive Care Society, and the French Society of Intensive Care Medicine

EARLY: KDIGO
dans les 12 h
DELAYED: atte
jusqu'à $k > 6$, pH
surcharge hydrique
AKI > 72 h l'IRA
44% des patients
choc septique



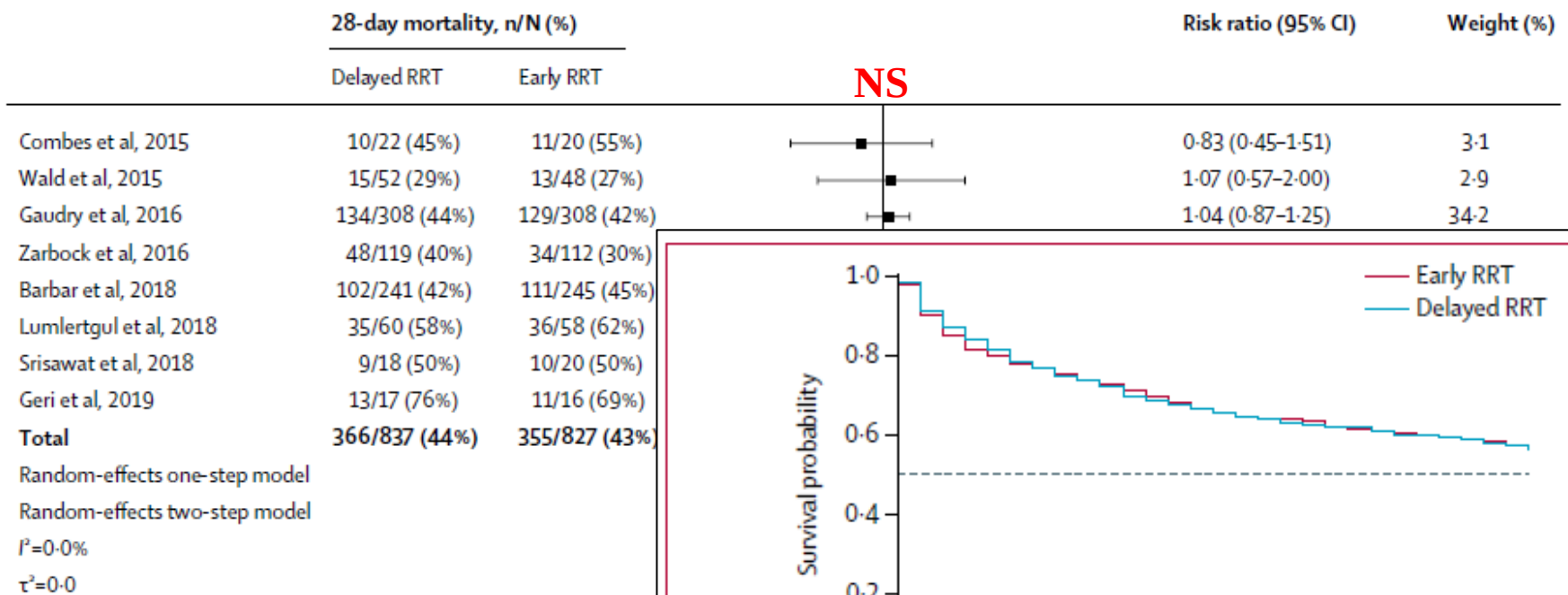
Pas de différence significative en terme de mortalité à j90
43,9% vs 43,7%
Plus d'effets indésirables dans le groupe précoce 23% vs 16,5%

No. at Risk										
Standard RRT	1462	1138	999	939	897	878	862	844	833	823
Accelerated RRT	1465	1122	985	925	892	865	846	835	830	823

Delayed versus early initiation of renal replacement therapy for severe acute kidney injury: a systematic review and individual patient data meta-analysis of randomised clinical

LANCET 2020

A



B

	28-day mortality, n/N (%)						
	Delayed RRT	Early RRT					
Number at risk							
Early RRT	831	636	552	509	474		
Delayed RRT	840	645	557	517	478		

Comparison of two delayed strategies for renal replacement therapy initiation for severe acute kidney injury (AKIKI 2): a multicentre, open-label, randomised, controlled trial

Stéphane Gaudry, David Hajage, Laurent Martin-L, Béatrice La Combe, Bertrand Pons, Nicolas de Prost, Guillaume Chevrel, Julien Bohé, Elisabeth Coupez, M, Eric Boulet, Karim Lakhal, Nadia Aissaoui, Steven C, Karim Asehnoune, Guillaume Geri, Kada Klouche, C, Jean-Damien Ricard*, Jean-Pierre Quenot†, Didier L

DELAYED: KDIGO 3
urée<40, oligurie>72 h
MORE DELAYED:
attendre jusqu'à
hyperK, urée>50,
acidose métab,
surcharge hydrique

	Delayed RRT strategy group (n=137)	More-delayed RRT strategy group (n=141)	p value
RRT-free days			
All patients	12 (0-25)	10 (0-24)	0.93
Survivors	24 (15-27)	23 (14-28)	0.54
Number of patients who actually received RRT	134 (98%)	111 (79%)	<0.0001
Time from randomisation to RRT, h	3 (2-5)	33 (24-60)	<0.0001
Number of RRT sessions*	5 (2-10)	5 (2-10)	0.75
Duration of RRT days*	5 (2-10)	5 (2-10)	0.75
Modality, first day*			
Intermittent RRT	81 (60%)	64 (58%)	0.53
Continuous RRT	52 (39%)	44 (40%)	..
Both modalities	1 (1%)	3 (3%)	..
Mortality			
At day 28	52 (38%)	63 (45%)	0.26
At day 60	60 (44%)	77 (55%)	0.071
At ICU discharge	55 (40%)	66 (47%)	0.26
At hospital discharge	61 (45%)	75 (53%)	0.15
RRT dependence‡			
At day 28	13 (16)	7 (11)	0.33
At day 60	3 (4)	1 (2)	0.62

LES RECOMMANDATIONS / TIMING

Réanimation

DOI 10.1007/s13546-014-0917-6

RÉFÉRENTIEL / *GUIDELINES*

2014

Il faut considérer « **précoce** » l'initiation d'une EER, au stade **KDIGO 2** ou dans les 24 heures suivant l'apparition d'une insuffisance rénale aiguë dont la réversibilité semble peu probable.

(Avis d'experts) Accord faible

et de la Société francophone de dialyse (SFD)

Il faut considérer « **tardive** » l'initiation de l'EER à **plus de 48 heures** de la survenue d'une **insuffisance rénale aiguë KDIGO 3** ou lors de l'apparition d'une situation mettant en jeu le pronostic vital et en rapport avec l'insuffisance rénale aiguë.

(Avis d'experts) Accord faible

LES RECOMMANDATIONS / TIMING

Réanimation

DOI 10.1007/s13546-014-0917-6

RÉFÉRENTIEL / *GUIDELINES*

2014

Il faut initier sans délai l'EER dans les situations mettant en jeu le pronostic vital (hyperkaliémie, acidose métabolique, syndrome de lyse, oedème pulmonaire réfractaire au traitement médical).

(Avis d'experts) Accord fort

et de la Société francophone de dialyse (SFD)

Les données disponibles sont insuffisantes pour définir le délai optimal avant instauration de l'EER en dehors des situations mettant en jeu le pronostic vital.

(Avis d'experts) Accord fort

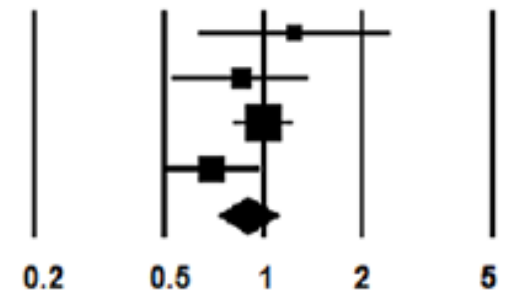
BMJ Open High-dose versus low-dose haemofiltration for the treatment of critically ill patients with acute kidney injury: an updated systematic review and meta-analysis

2017

(A) 90-day mortality

Pas de difference
mortalité j90

Study name	Statistics for each study				
	Odds ratio	Lower limit	Upper limit	Z-Value	p-Value
Joannes-Boyau (2013)	1.24	0.63	2.43	0.63	0.53
Zhang (2012)	0.85	0.53	1.38	-0.64	0.52
Bellomo (2009)	1.00	0.81	1.22	-0.03	0.97
Ronco (2000)	0.70	0.50	0.97	-2.12	0.03
Pooled	0.90	0.73	1.11	-1.00	0.32



Heterogeneity test: $Q = 4.10$, $P = 0.25$, $I^2 = 26.7\%$

Favors high-dose

Favors low-dose

8 st
HV
SV
Ma
Per
pat

LES RECOMMANDATIONS / DOSE DE DIALYSE

Réanimation

DOI 10.1007/s13546-014-0917-6

RÉFÉRENTIEL / *GUIDELINES*

2014

En épuration extrarénale intermittente il faut probablement que la dose de dialyse minimale délivrée soit de

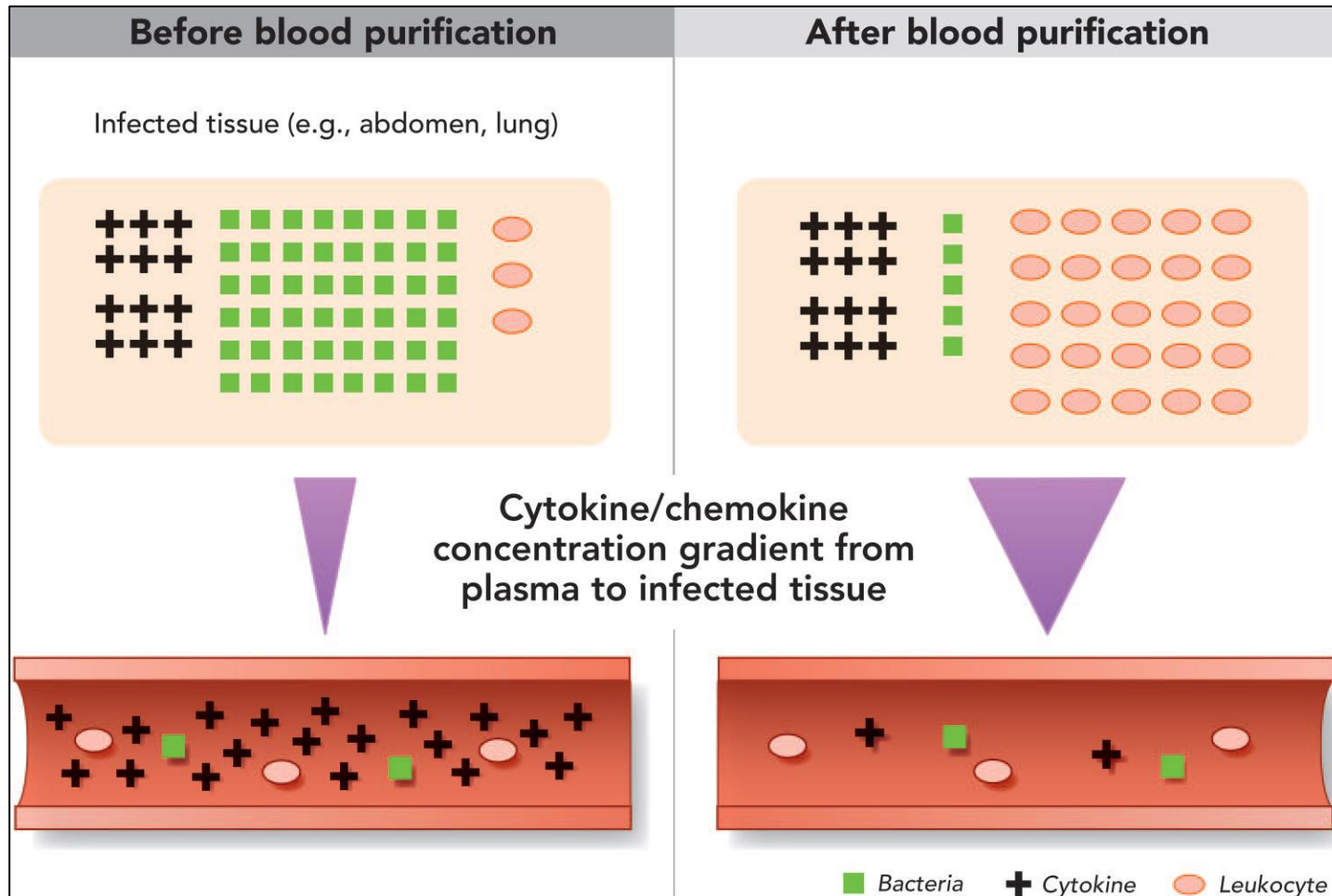
- 1) **trois séances par semaine de quatre heures au moins** avec un **débit sang > 200 ml/min** et un **débit dialysat > 500 ml/min**,
 - 2) l'obtention d'un **$Kt/V > 3,9$** par semaine,
 - 3) le maintien d'une concentration en **urée pré-dialytique de 20–25 mmol/l**.
- Accord fort**

En épuration extrarénale continue, il faut probablement que la dose de dialyse minimale délivrée soit de **20–25 ml/kg par heure** d'effluent, obtenus par filtration et/ou diffusion.

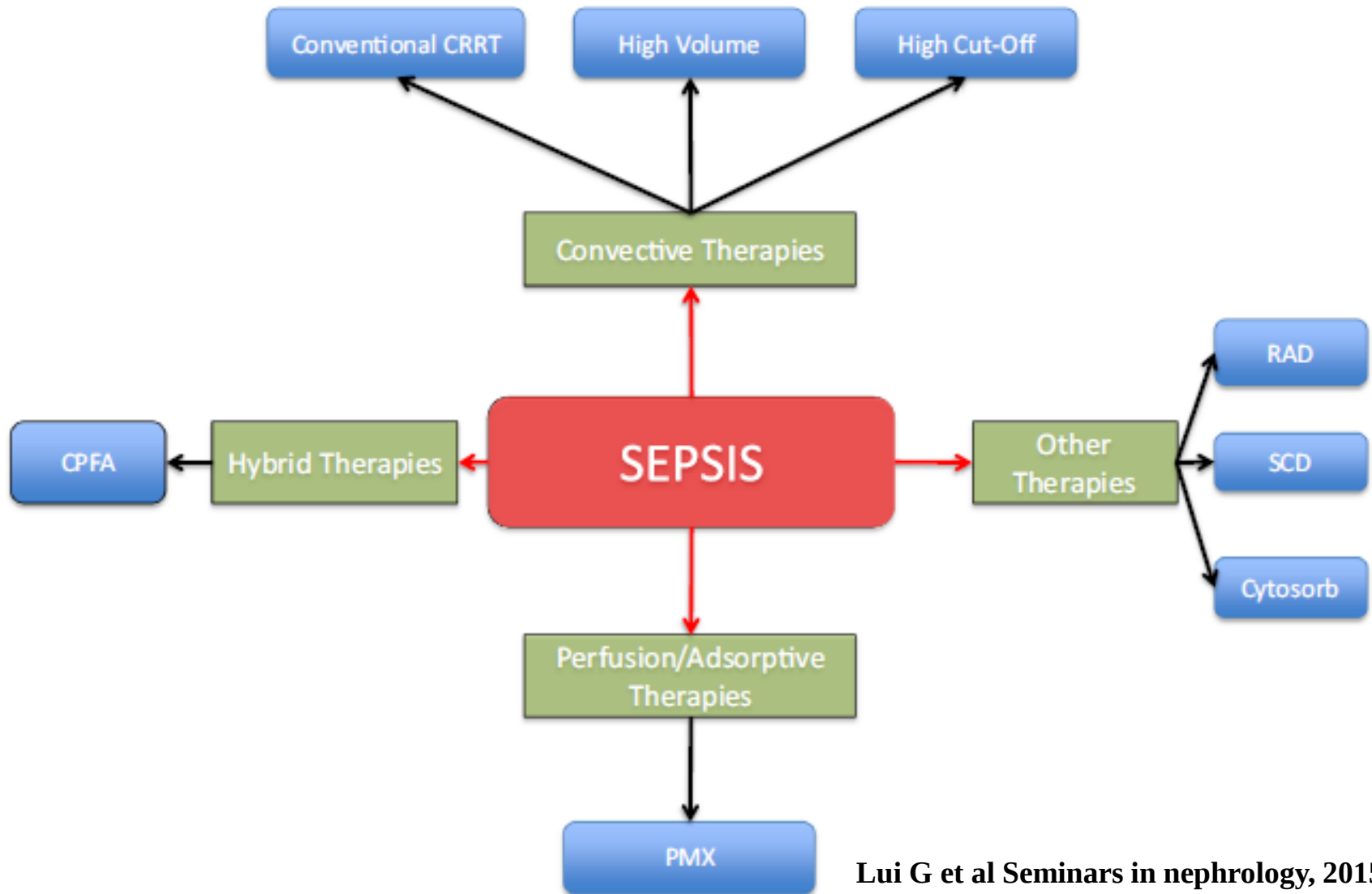
Accord fort

**EER ET PURIFICATION
SANGUINE AU COURS DU CHOC
SEPTIQUE
AU DELÀ DE LA SUPPLÉANCE
RÉNALE !**

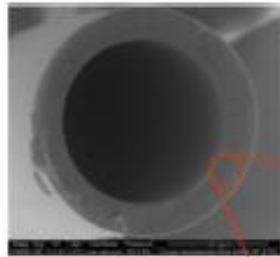
RRT FOR BLOOD PURIFICATION



RRT FOR BLOOD PURIFICATION



RRT FOR BLOOD PURIFICATION



Endotoxin
Polyethyleneimine coating

Fig. 4. High adsorptive hemofilter polarity is modified by the adsorption coating, a positively charged polymer surface adsorption.

Rimmelé et al Anesthesiology, 2004

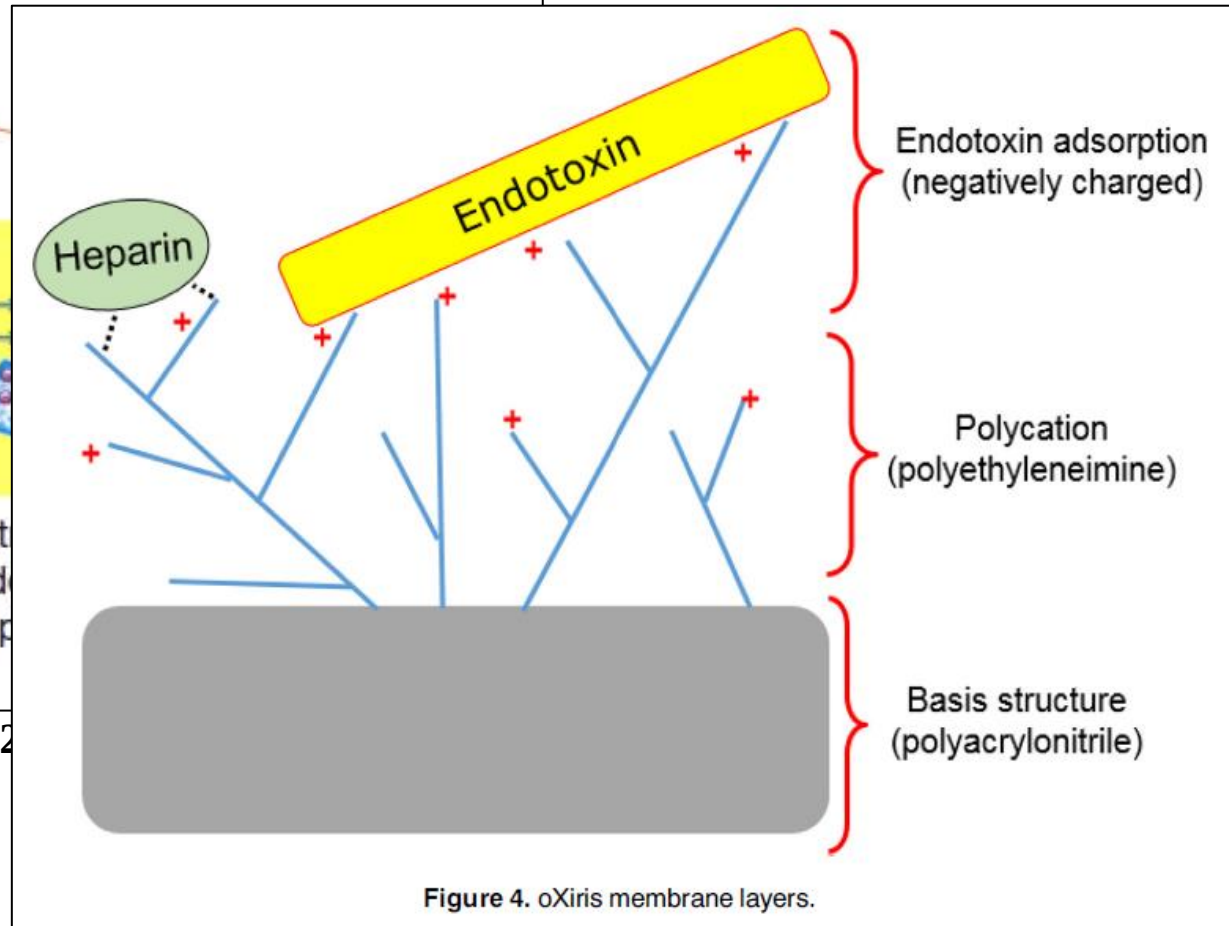


Figure 4. oXiris membrane layers.

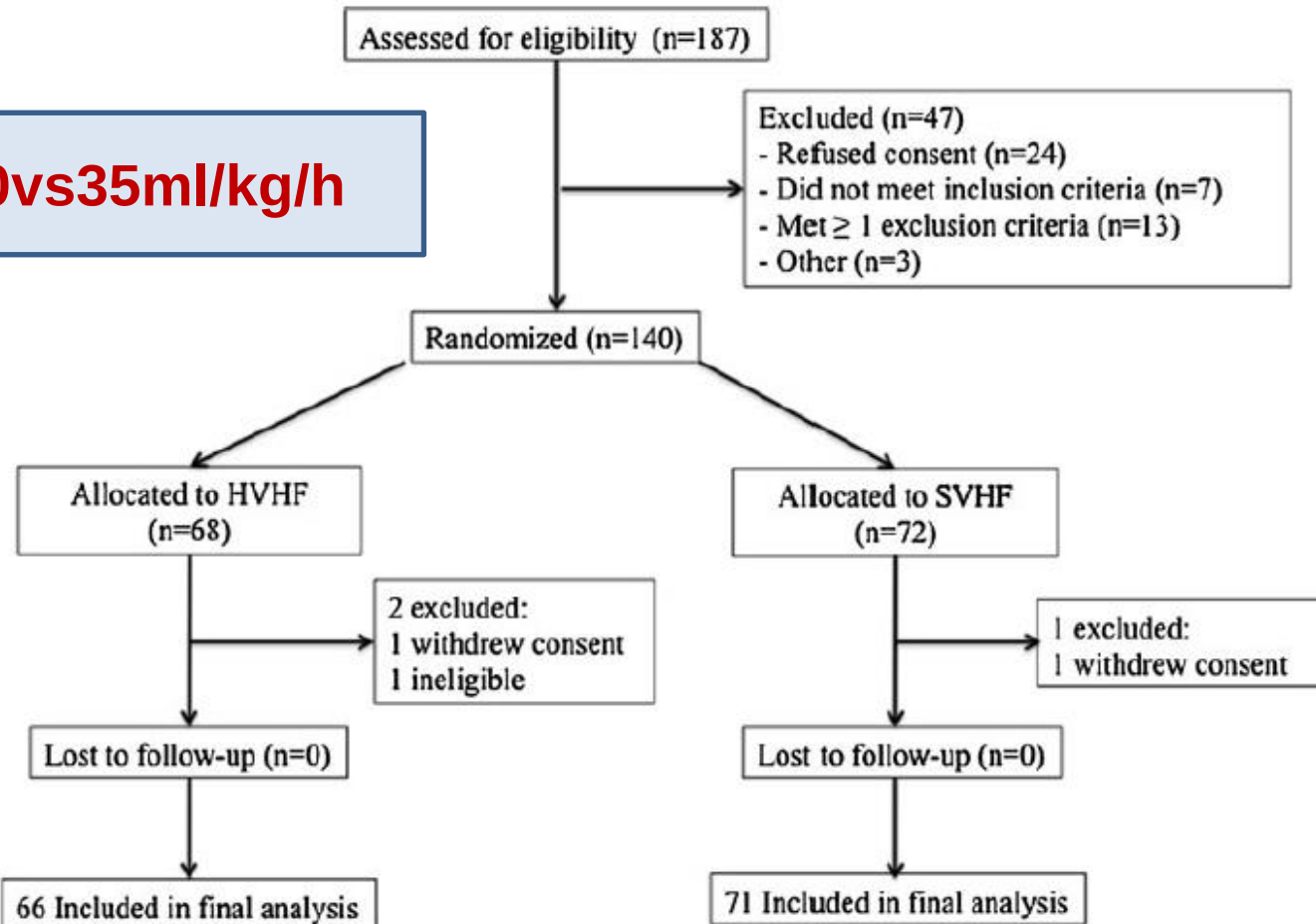
Thibaut Girardot et al, Seminars in Nephrology 2019

IVOIRE STUDY
Intensive Care Med 2013

Olivier Joannes-Boyau
Patrick M. Honoré
Paul Perez
Sean M. Bagshaw
Hubert Gran
Jean-Luc Car
Antoine Dewi
Claire Flan
Wilfried P
Anne-Soph
Catherine
Rita Jacobs
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Hervé Floch
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Stephane Fra
Hadrien Rozé
Vincent Colli
Willem Boer
Joachim Cald
Bernard Gau
Herbert D. S
Gérard Janvi
Alexandre Ou

**High-volume versus standard-volume
haemofiltration for septic shock patients**

70vs35ml/kg/h



IVOIRE STUDY
Intensive Care Med 2013

Olivier Joannes-Boyau
Patrick M. Honoré

High-volume versus standard-volume

Paul
Sean
Hub
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Will
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Cath
Rita
Chri
Herv
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Step
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Vinc
Will
Joac
Berr
Herl
Géra
Alex

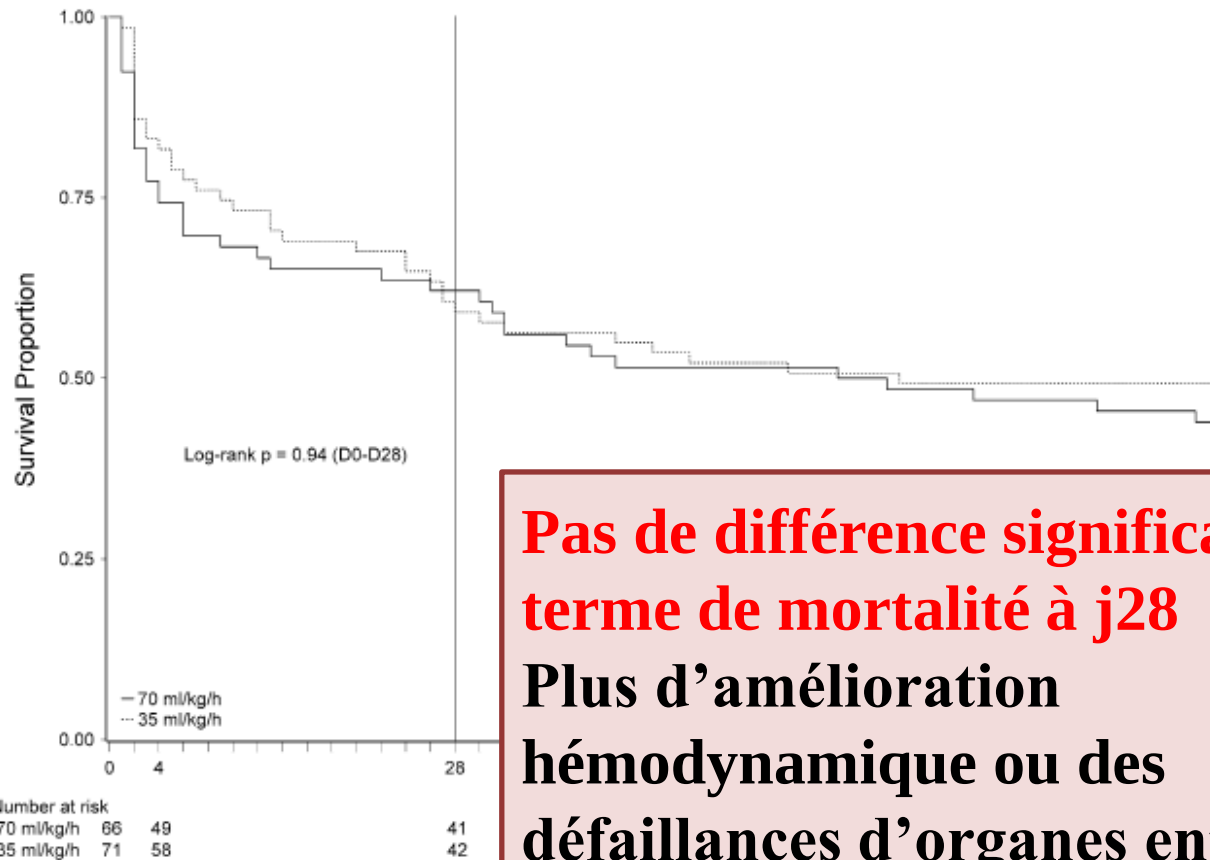


Fig. 2 Survival of participants in the IVOIRE (hIgh VO) underwent haemofiltration at 70 mL/kg/h and 35 mL/kg/h

Pas de différence significative en terme de mortalité à j28
Plus d'amélioration hémodynamique ou des défaillances d'organes entre groupe HVHF vs SVHF

RESEARCH

Open Access

2014

High-volume hemofiltration for septic acute kidney injury: a systematic review and meta-analysis

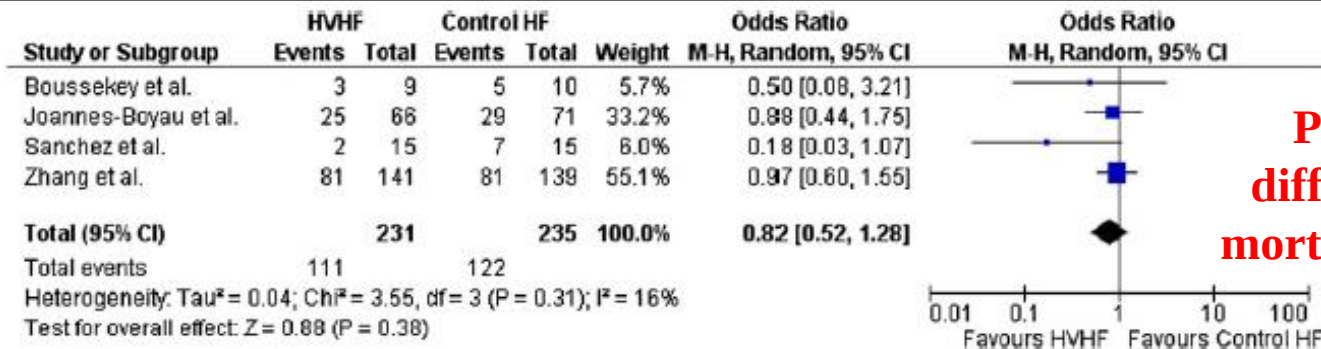


Figure 2 Forest plot for odds of

**Plus d'effets indésirables:
hypophosphatémie et
hypokaliémie groupe HVHF**

The therapeutic effect of high-volume hemofiltration on sepsis: a systematic review and meta-analysis

2020

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Contributions: (I) Conception and design: B Ning, S Liu; (II) Administrative support: None; (III) Provision of study materials or patients: F Yin, F Zhang; (IV) Collection and assembly of data: F Yin, S Liu; (V) Data analysis and interpretation: F Yin; (VI) Manuscript writing: All authors; (VII) Final approval of manuscript: All authors.

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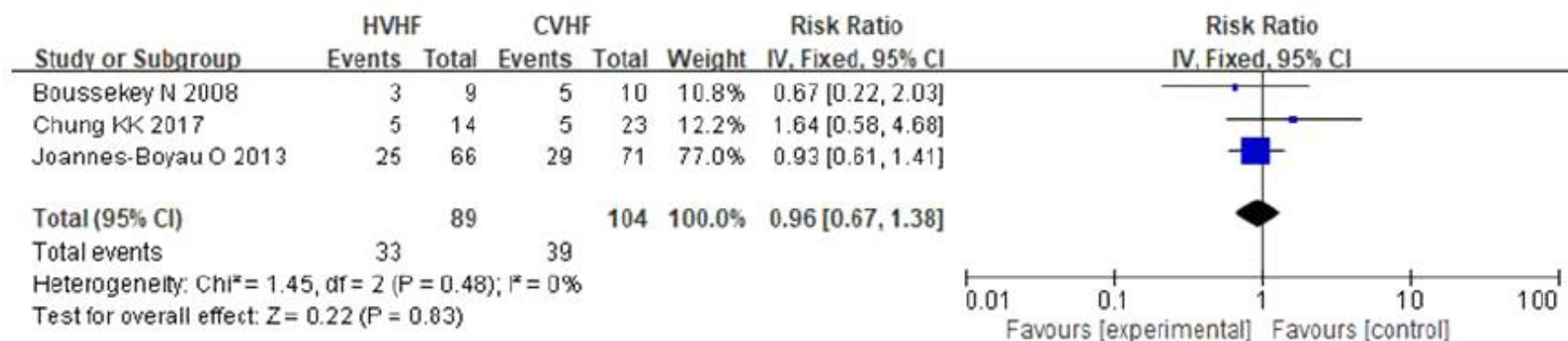


Figure 4 Forest plot of comparison in relative risk of mortality.

Pas de différence significative en terme de mortalité



2015

Jean-Pierre Quenot
Christine Biquet
Christophe Vinsonneau
Saber-David Barbar
Sandrine Vinault
Valérie Degeilh

Very high volume hemofiltration with the Cascade system in septic shock patients

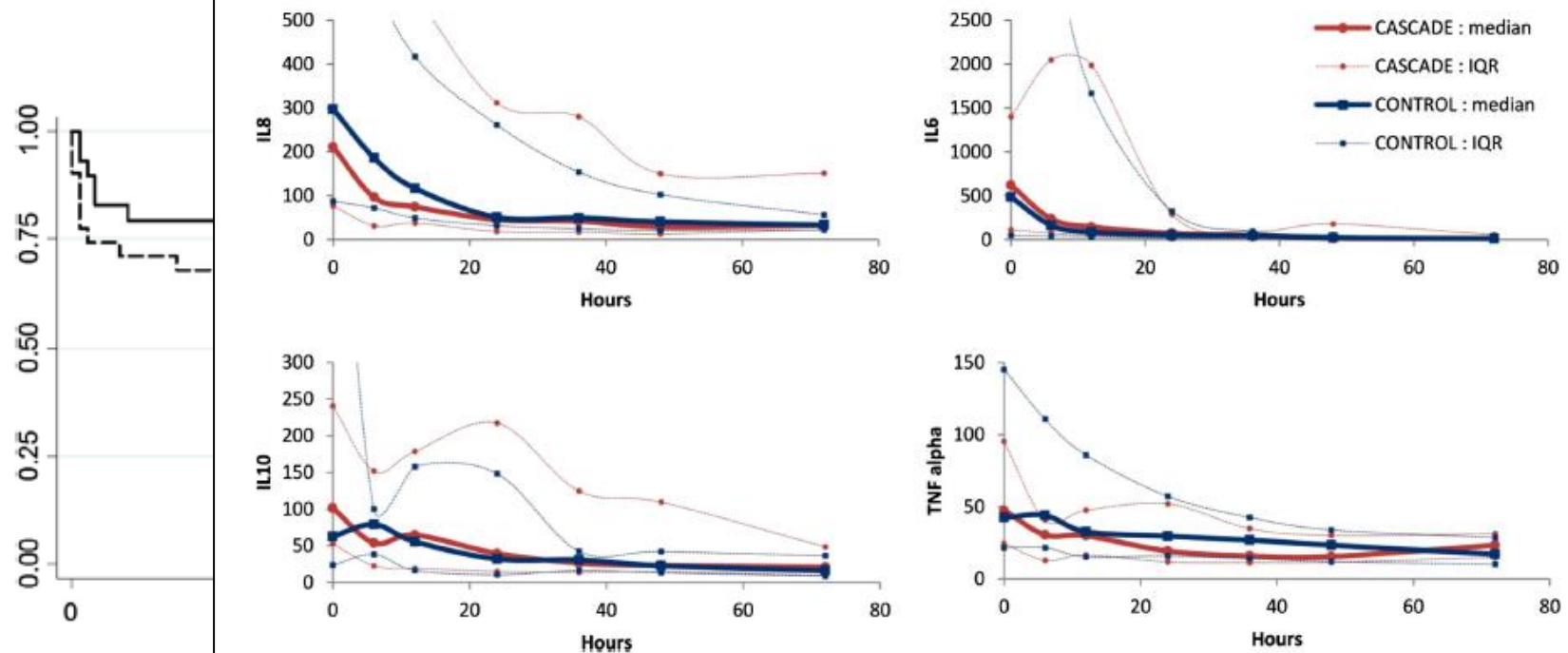
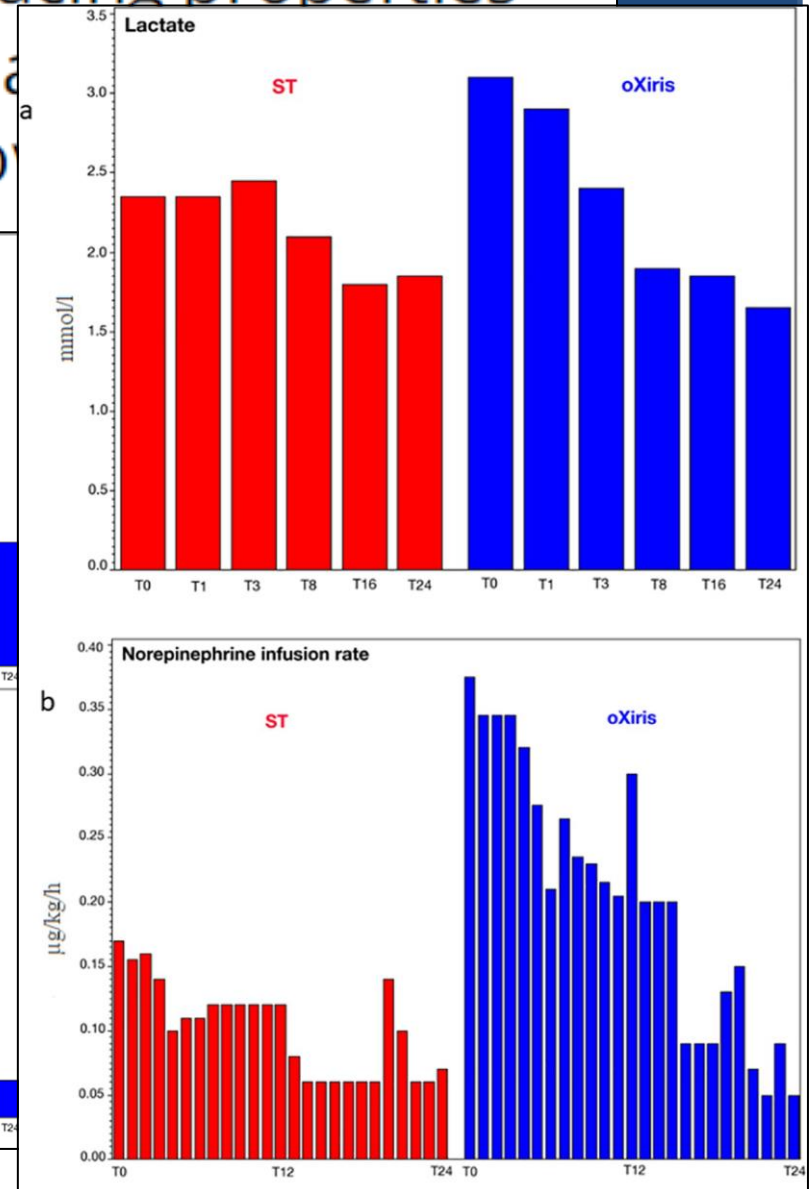
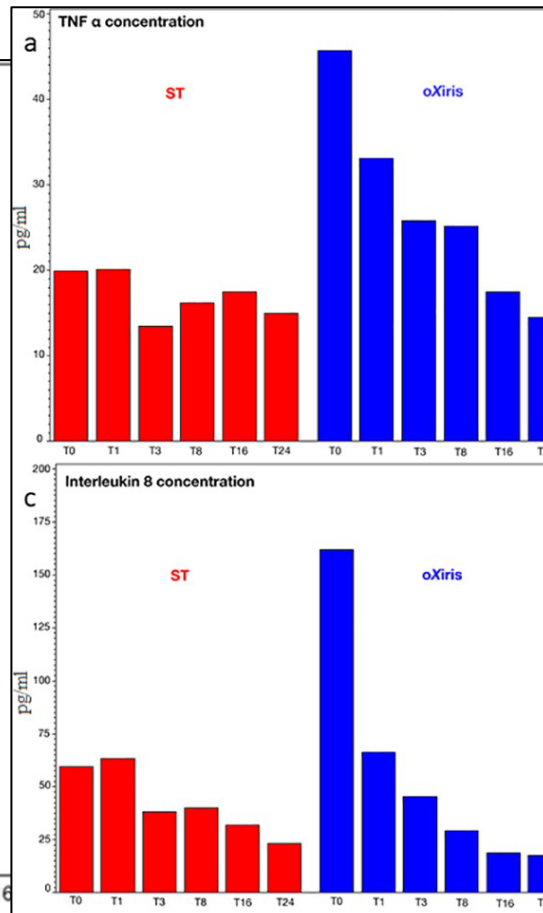
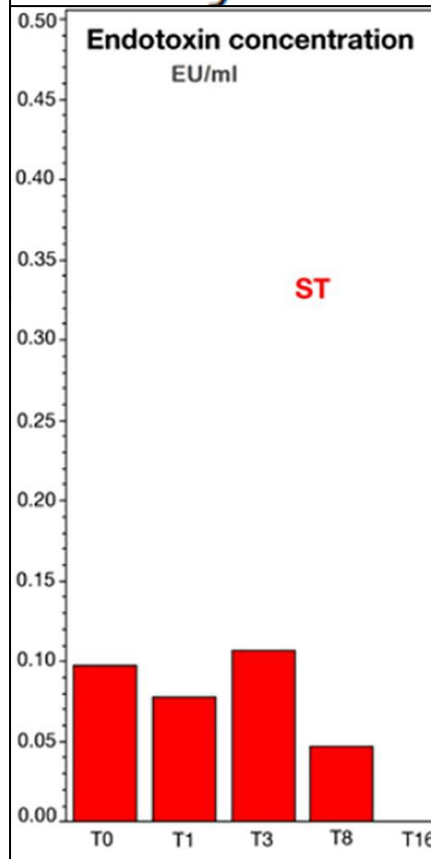


Fig. 3 Circulating cytokine levels (pg/mL) in the control and Cascade groups at 0, 12, 18, 24, 36, 48, and 72 h

Fig. 2 Cumulative survival probabilities in each group at 90 days

Endotoxin and cytokine reducing properties of the oXiris membrane in pa shock: A randomized crossover study



Enrollment

Assessed for eligibility (n = 437)

Excluded (n = 399)

Table 2 Reduction in endotoxin activity, Interleukin-6, procalcitonin and HMGB-1 over time

Intervention arm	Filter intervention				p—over groups	p—over timepoints
	0 h	24 h	48 h	72 h		
A. Reduction in endotoxin activity [%]						
Standard of care	0	16 [10–28]	2 [–18–21]	12 [1–42]	0.82	<0.01
oXiris	0	35 [18–53]	33 [5–53]	31 [18–53]		
Toraymyxin	0	35 [18–53]	33 [5–53]	31 [18–53]		
B. Reduction in Interleukin-6 [ng/l]						
Standard of care	0	16 [10–28]	2 [–18–21]	12 [1–42]	0.58	<0.001
oXiris	0	35 [18–53]	33 [5–53]	31 [18–53]		
Toraymyxin	0	35 [18–53]	33 [5–53]	31 [18–53]		
C. Reduction in Procalcitonin [pg/ml]						
Standard of care	0	16 [10–28]	2 [–18–21]	12 [1–42]	0.16	<0.01
oXiris	0	35 [18–53]	33 [5–53]	31 [18–53]		
Toraymyxin	0	35 [18–53]	33 [5–53]	31 [18–53]		
D. Reduction in HMGB-1 [pg/ml]						
Standard of care	0	16 [10–28]	2 [–18–21]	12 [1–42]	0.68	0.46
oXiris	0	35 [18–53]	33 [5–53]	31 [18–53]		
Toraymyxin	0	35 [18–53]	33 [5–53]	31 [18–53]		

A

B

Endotoxin Activity

Time [hours]

Standard of Care oXiris Toraymyxin

Interleukin-6 [ng/l]

Time [hours]

Screening 0 24 48 72

- Use of 3 filters (n=10)
- Death less than 24h (n=10)
- Death less than 24h randomization (n=10)

g. 1 Screening, ran

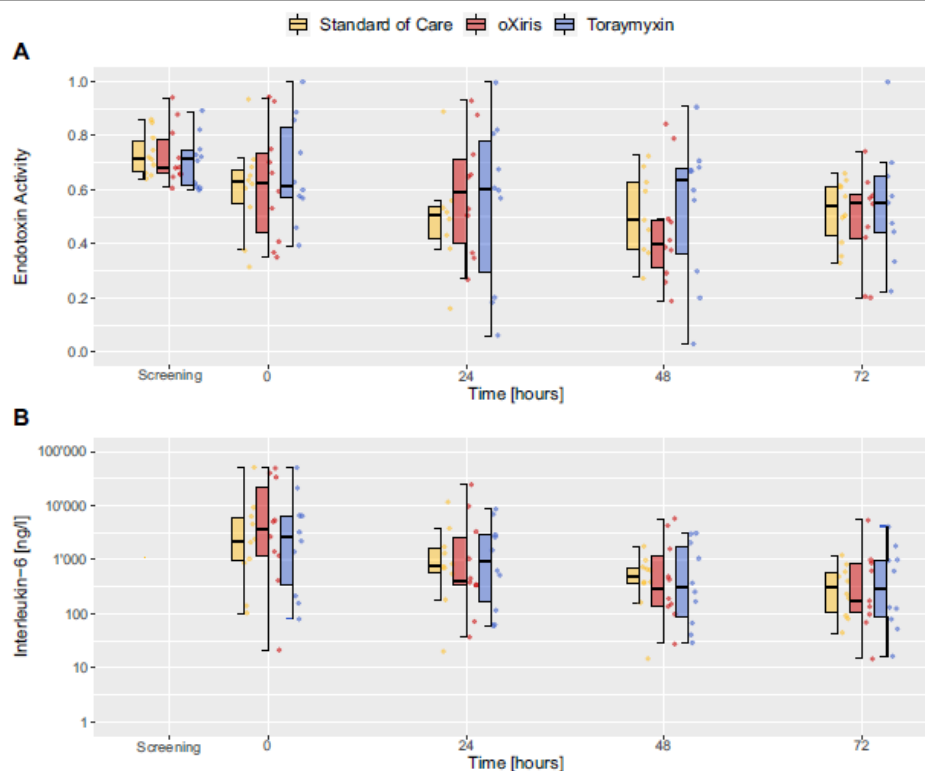


Fig. 1 Screening, randomization, and follow-up flowchart

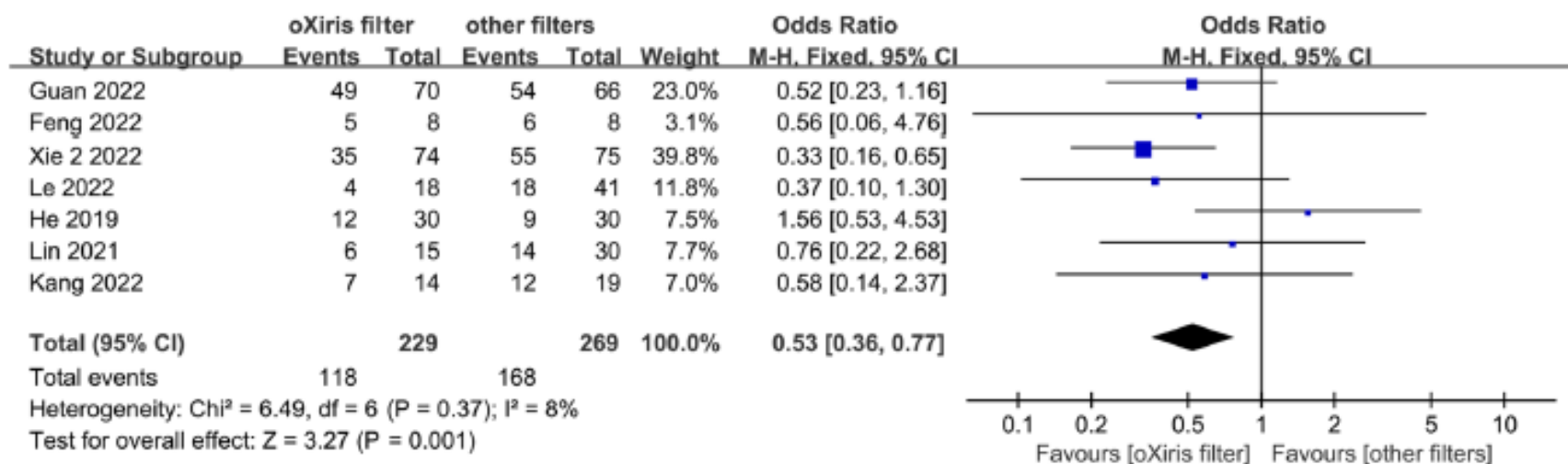


Fig. 3. 28-day mortality (adopting Xie et al's data after IPTW)

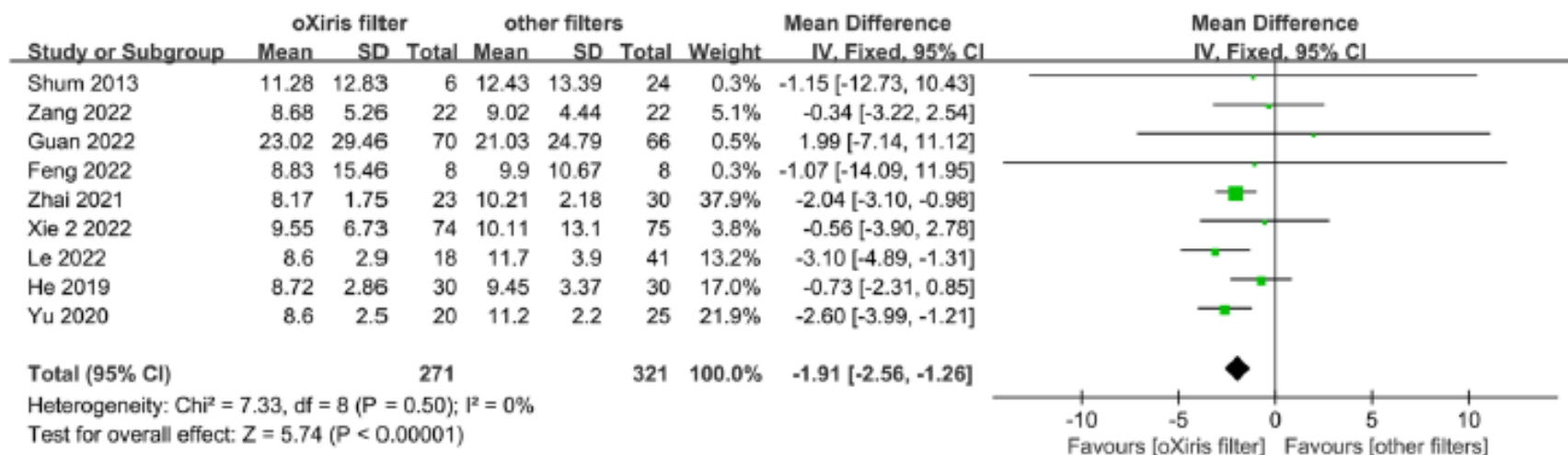


Fig. 4 The length of ICU stay (adopting Xie et al's data after IPTW)

LES RECOMMANDATIONS / BLOOD PURIFICATION

Réanimation

DOI 10.1007/s13546-014-0917-6

RÉFÉRENTIEL / *GUIDELINES*

2014

**Épuration extrarénale en réanimation adulte et pédiatrique.
Recommandations formalisées d'experts sous l'égide**

En épuration extrarénale continue, *il ne faut pas, sur la seule présence d'un sepsis, intensifier la dose d'épuration.*
Accord fort

Renal replacement therapy Adult and Children Intensive Care Unit.

Société de réanimation de langue française. Experts Recommendations

Christophe Vinsonneau, Emma Allain-Launay, Clarisse Blayau, Michael Darmon, Damien du Cheyron, Théophile Gaillot, Patrick Honoré, Étienne Javouhey, Thierry Krummel, Annie Lahoche, Matthieu Legrand, Serge Le Tacon, Mehran Monchi, Christophe Ridel, René Robert, Frédérique Schortgen, Bertrand Souweine, Patrick Vaillant, Lionel Velly, David Osman, Ly Van Vong

LES RECOMMANDATIONS / BLOOD PURIFICATION

Intensive Care Med (2021) 47:1181–1247
<https://doi.org/10.1007/s00134-021-06506-y>

Intensive Care Med 2021

GUIDELINES

Surviving sepsis campaign: international guidelines for management of sepsis and septic shock 2021



- There is insufficient evidence to make a recommendation on the use of other blood purification techniques

Theodore Iwashyna³³, Shevin Jacob³⁴, Ruth Kleinpell³⁵, Michael Klompas^{36,37}, Younsuck Koh³⁸, Anand Kumar³⁹, Arthur Kwizera⁴⁰, Suzana Lobo⁴¹, Henry Masur⁴², Steven McGloughlin⁴³, Sangeeta Mehta⁴⁴, Yatin Mehta⁴⁵,

Conclusion

Les données de la littérature sur **l'EER comme moyen de suppléance de la défaillance rénale au cours du choc septique (SA-AKI)** sont **insuffisante** pour recommander l'une ou l'autre modalité et définir le timing optimal

- **Continue vs intermittente ?????**
- **Précoce vs tardive ?????**

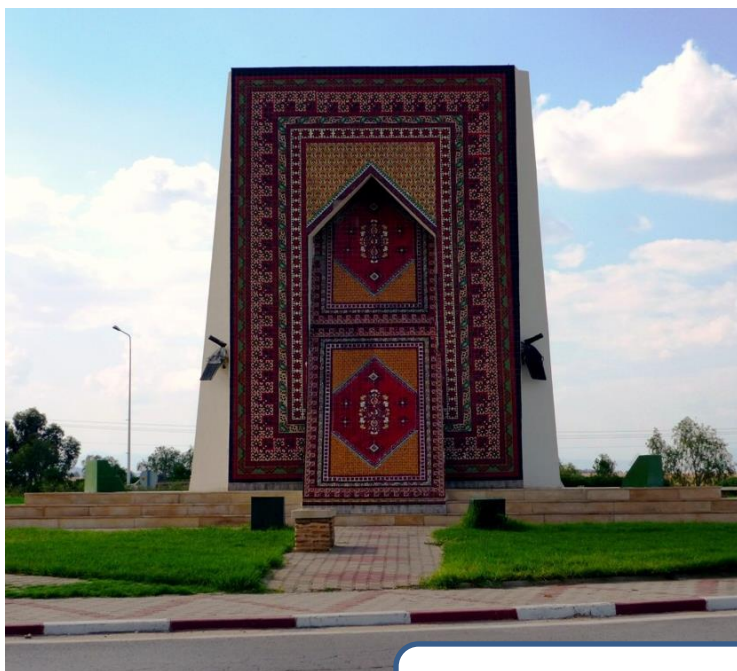
Il faut tenir compte de **la disponibilité de la technique** et de **l'expérience de l'équipe.**

Conclusion

Les arguments en **faveur ou contre l'EER comme moyen de purification sanguine** pour la modulation de l'inflammation ne sont pas suffisamment importants

Pas de recommandations claires a ce jour !!

Il faut continuer la recherche...



Merci pour votre attention

