



XXXI Corso Nazionale ANTE - Dialisi e Tecnologia

**“Evoluzione tecnologica nei trattamenti dialitici cronici e acuti:
dalla teoria alla pratica”**

**15-16-17 Aprile 2024 Sala Congressi Hotel CORALLO Viale
Antonio Gramsci, 113, 47838 Riccione RN**

Membrane adsorbenti tra vecchie e nuove indicazioni cliniche

**dott. P. Fabbrini
ASST NORD MILANO
OSPEDALE BASSINI**

Tossine Uremiche e correlazioni cliniche

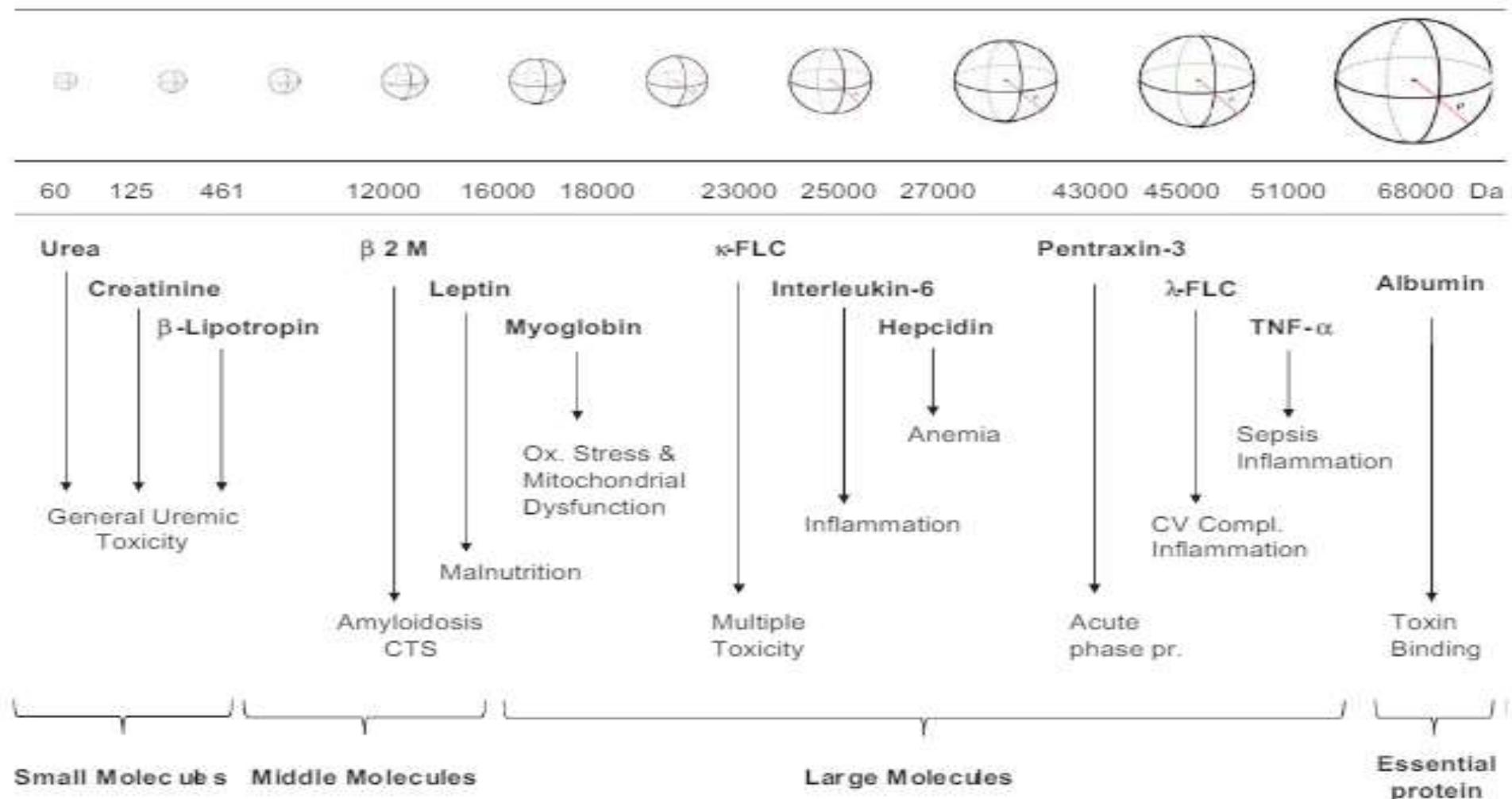
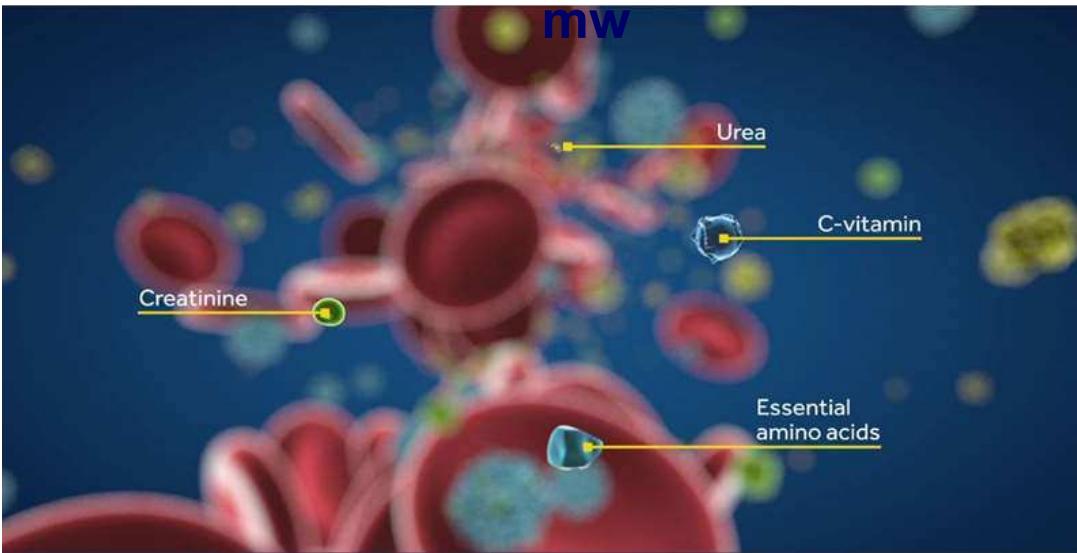
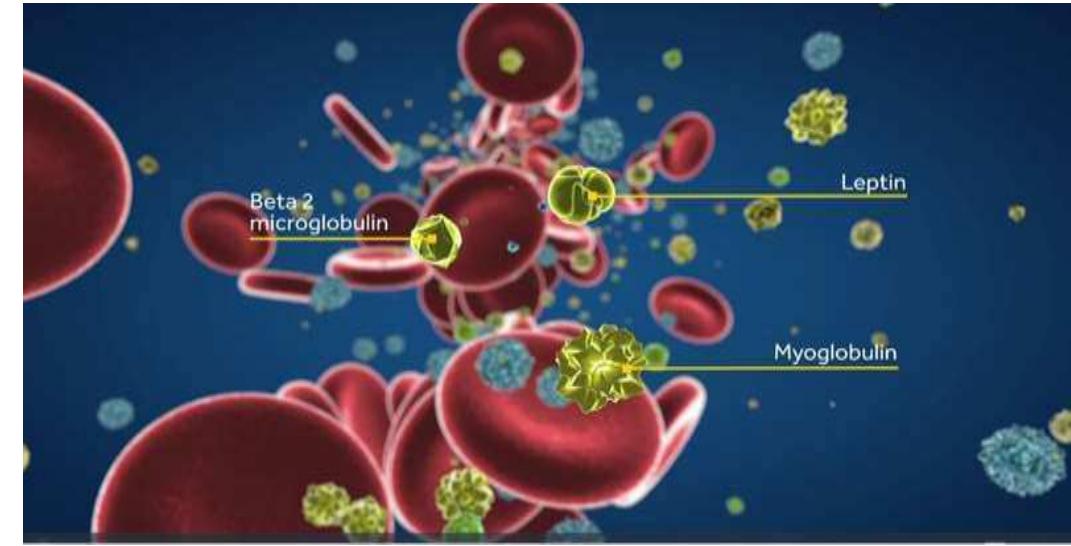


FIGURE 1: Schematic representation of different classes of uremic toxins with their molecular size and relevant clinical effects.

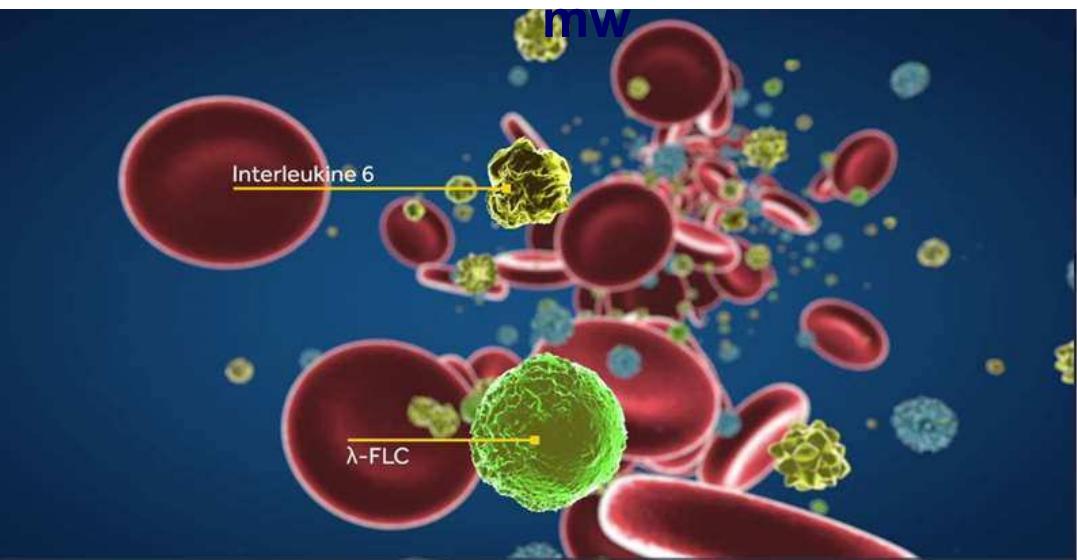
Low



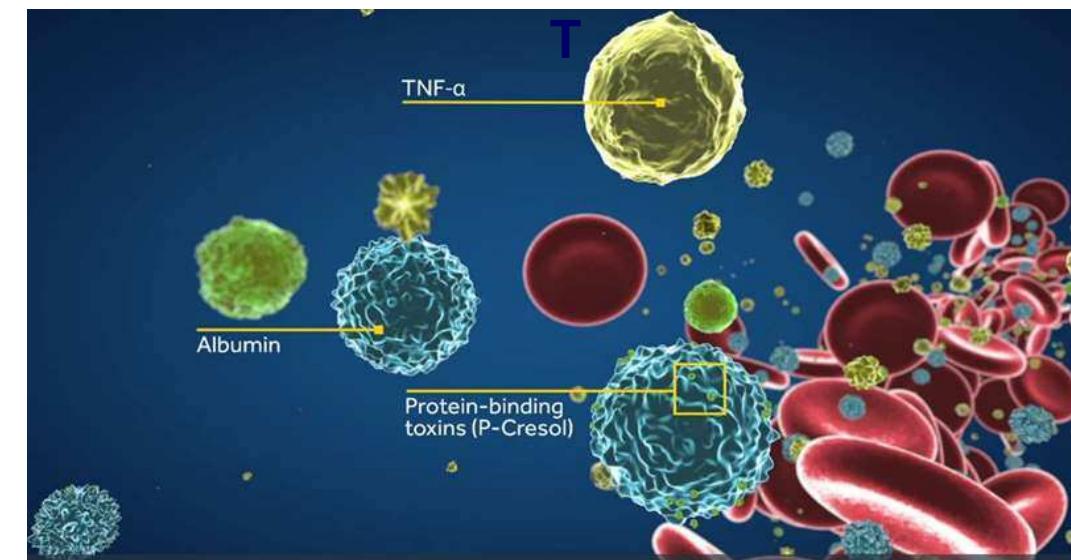
Medium mw



Large



PBU



DIFFUSIONE E CONVEZIONE: CONTRAST STUDY

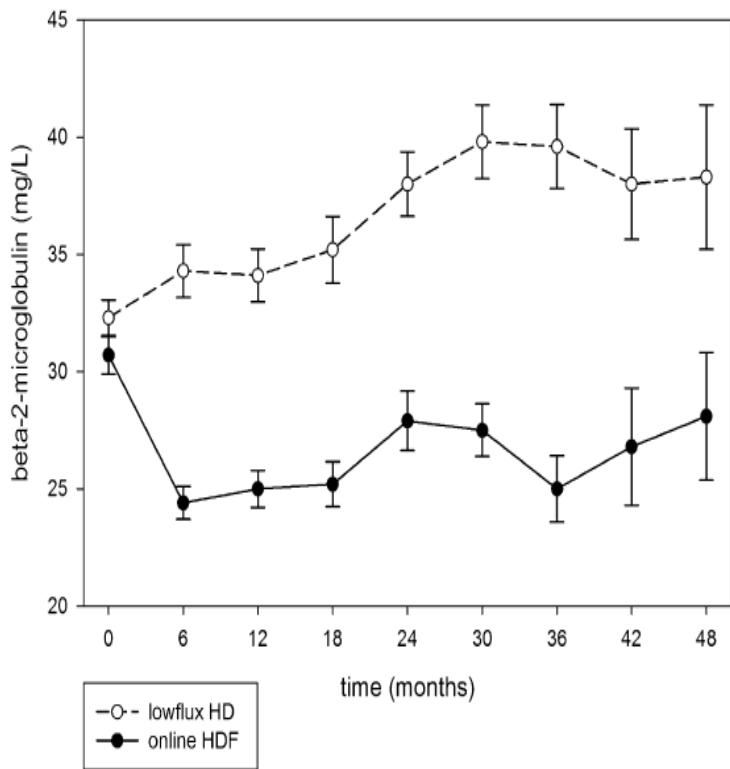
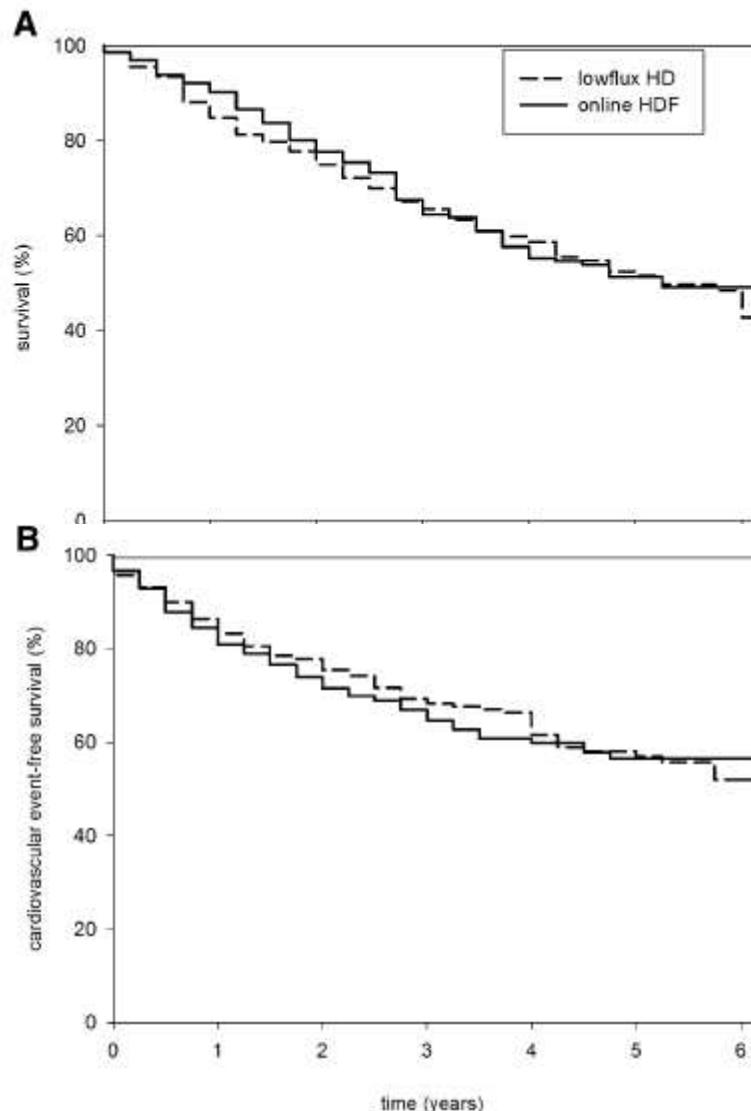


Figure 2. Predialysis β -2-microglobulin levels in patients treated with online hemodiafiltration and low-flux hemodialysis (mean \pm SEM) using measurements of individuals at those time points. The difference between β -2-microglobulin levels for both treatments was significant ($P<0.001$).



Effect of Online Hemodiafiltration on All-Cause Mortality and Cardiovascular Outcomes

DIFFUSIONE E CONVEZIONE: CONTRAST STUDY

Table 4. Risk of all-cause mortality and fatal and nonfatal cardiovascular events by achieved convection volume in liters per treatment

| | Hemodialysis | Online Hemodiafiltration Convection Volume Tertiles | | | P for Trend |
|--|--------------|---|------------------|------------------|-------------|
| | | <18.17 L | 18.18–21.95 L | >21.95 L | |
| Total mortality | | | | | |
| crude | 1.0 | 0.95 (0.66–1.38) | 0.83 (0.57–1.22) | 0.62 (0.41–0.93) | 0.010 |
| adjusted ^a | 1.0 | 0.79 (0.53–1.14) | 0.77 (0.51–1.14) | 0.65 (0.42–0.99) | 0.012 |
| adjusted ^b | 1.0 | 0.80 (0.52–1.24) | 0.84 (0.54–1.29) | 0.61 (0.38–0.98) | 0.015 |
| Fatal and nonfatal cardiovascular events | | | | | |
| crude | 1.0 | 1.37 (0.94–1.98) | 1.06 (0.72–1.56) | 0.76 (0.50–1.16) | 0.473 |
| adjusted ^a | 1.0 | 1.41 (0.92–2.11) | 0.93 (0.62–1.40) | 0.77 (0.48–1.21) | 0.369 |
| adjusted ^b | 1.0 | 1.35 (0.86–2.11) | 1.04 (0.66–1.62) | 0.72 (0.44–1.19) | 0.475 |

THE MORE THE BETTER?

RESEARCH SUMMARY

Effect of Hemodiafiltration or Hemodialysis on Mortality in Kidney Failure

Blankestijn PJ et al. DOI: 10.1056/NEJMoa2304820

CLINICAL PROBLEM

Hemodiafiltration and hemodialysis are two methods of treating kidney failure. Data comparing survival outcomes with these approaches have been largely inconclusive; results of one randomized, controlled trial indicating a lower risk of death with hemodiafiltration may have reflected confounding bias.

CLINICAL TRIAL

Design: A multinational, pragmatic, open-label, randomized, controlled trial assessed the benefits and harms of high-dose hemodiafiltration as compared with conventional high-flux hemodialysis in patients with kidney failure.

Intervention: 1360 adults with stage V kidney failure who had received hemodialysis for ≥3 months were assigned to receive high-dose hemodiafiltration (a convection volume of ≥23 liters in postdilution mode per session) or continuation of high-flux hemodialysis. The primary outcome was death from any cause.

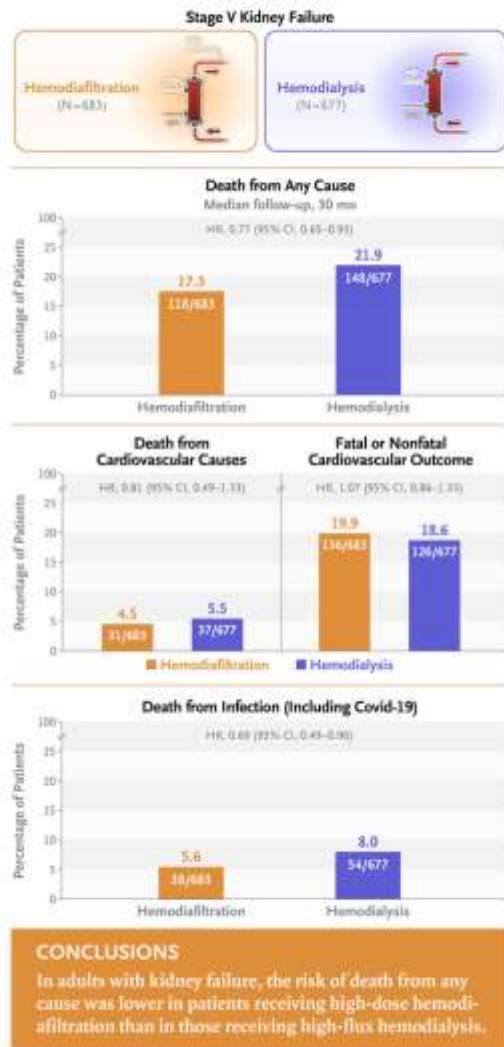
RESULTS

Outcomes: During a median follow-up of 30 months, the risk of death from any cause was lower in patients receiving high-dose hemodiafiltration than in those receiving hemodialysis. The survival effect of hemodiafiltration appeared to be limited to those without a history of cardiovascular disease or diabetes.

LIMITATIONS AND REMAINING QUESTIONS

- The achieved sample size was lower than originally planned because of difficulty recruiting patients during the Covid-19 pandemic.
- Inclusion criteria may have resulted in a trial population that was healthier than the general population of patients who receive hemodialysis in Europe and the United States.
- Findings may not be generalizable beyond White European populations.

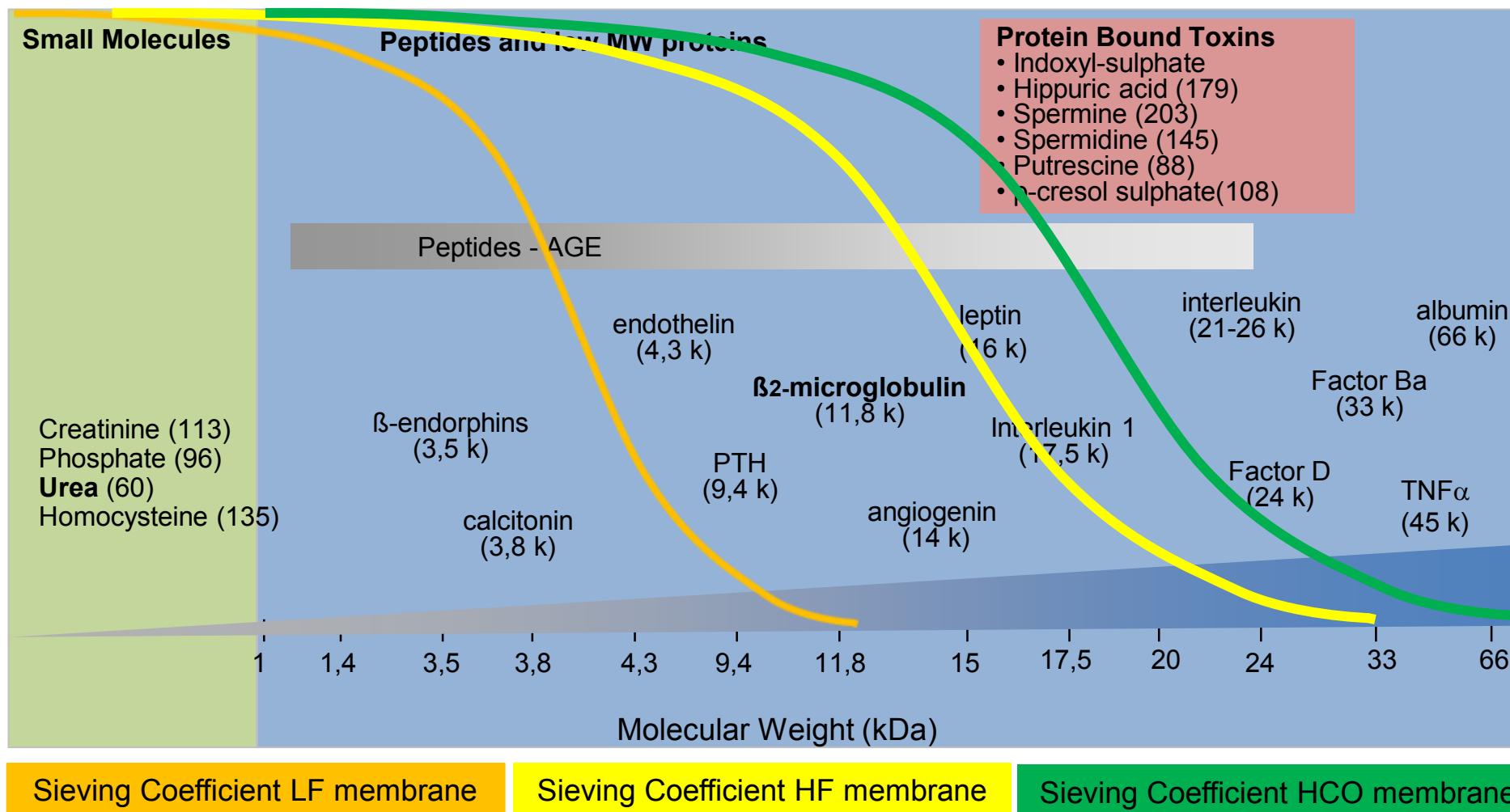
Links: Full Article | NEJM Quick Take | Editorial



ALLORA HA SENSO ESTENDERE ANCHE LO SPETTRO DELLE MOLECOLE RIMOSSE? COME INTRODURRE L'ADSORBIMENTO?

Tossine Uremiche

overcome Membranes limitations



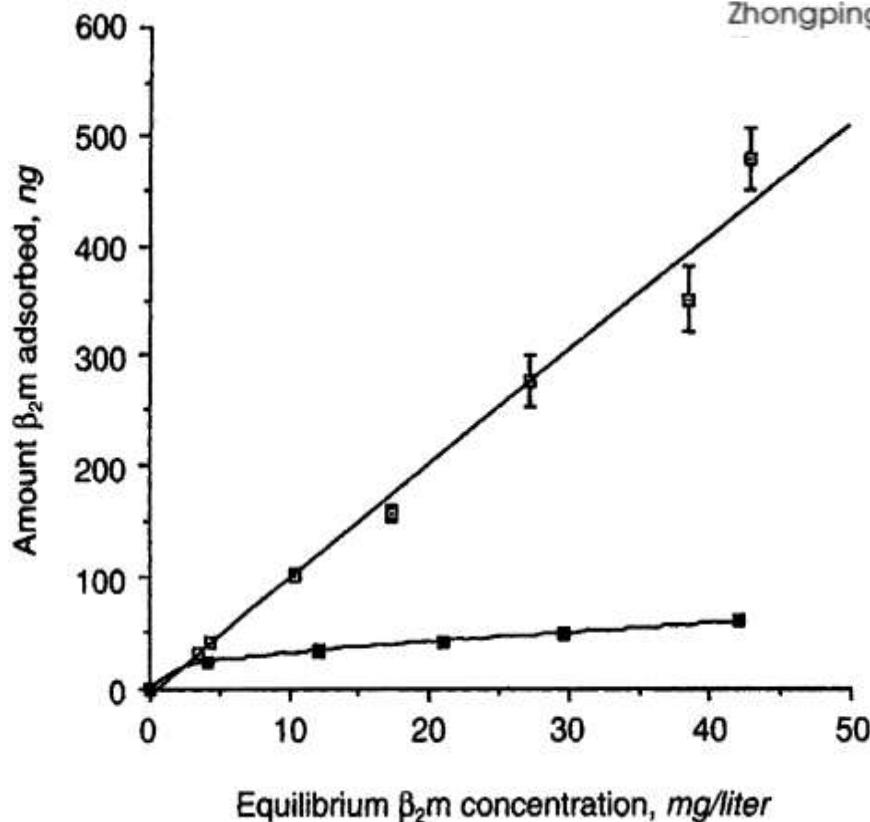
DIALISI CON ADSORBIMENTO



ADSORBIMENTO

Blood-Membrane Interactions During Dialysis

Zhongping Huang,* Dayong Gao,† Jeffrey J. Letteri,† and William R. Clark†§

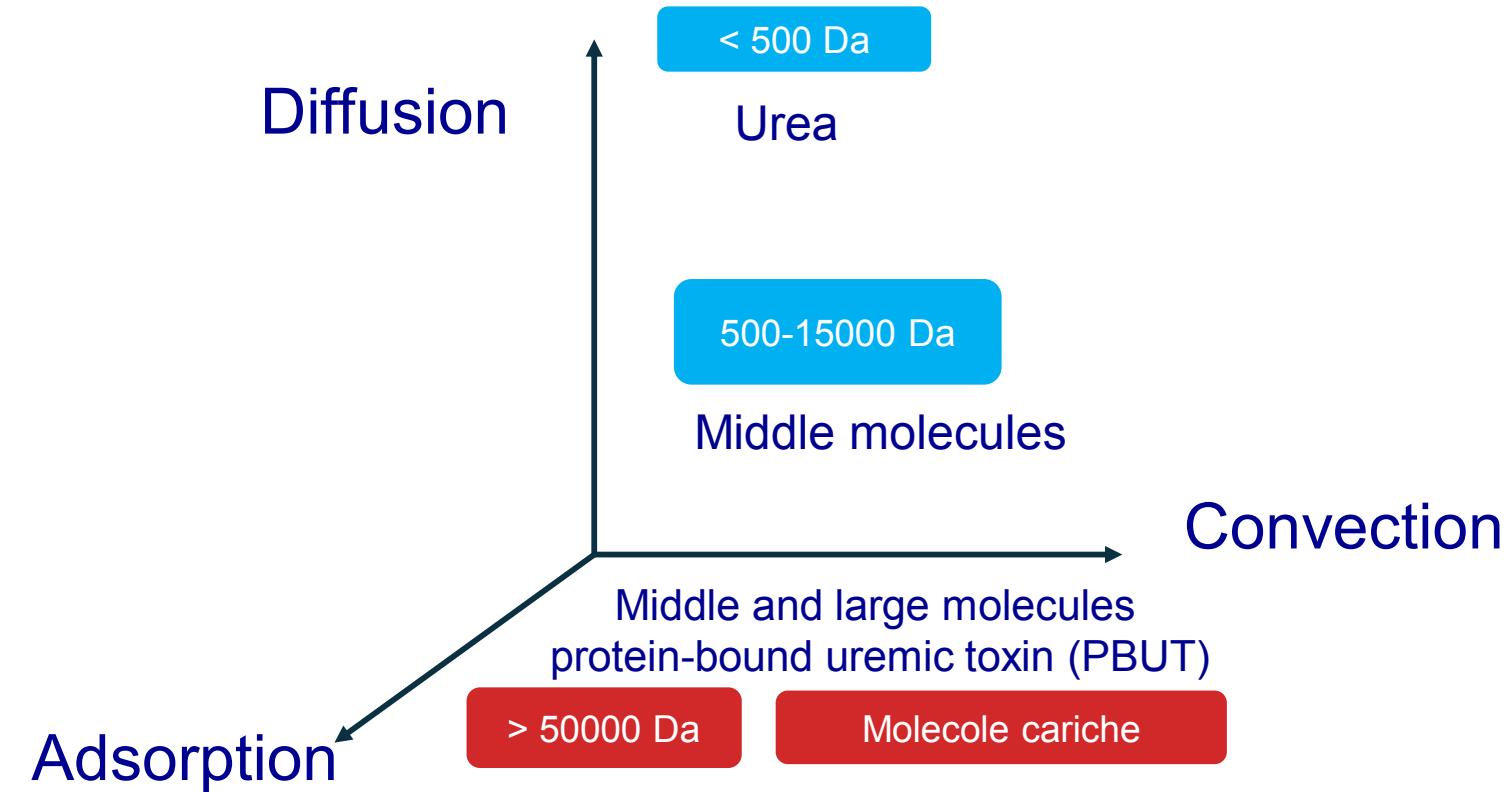


FORZE COINVOLTE

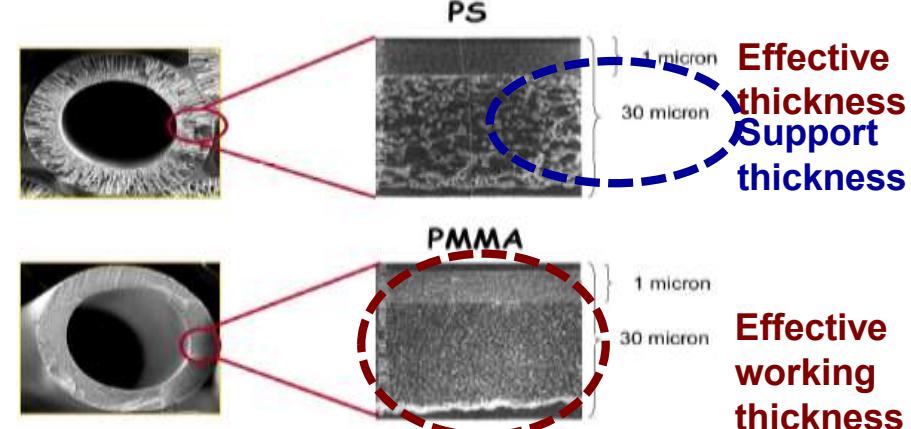
- INTERAZIONI IDROFOBICHE
- FORZE IONICHE-ELETTROSTATICHE
- FORZA DI VAN DER WAALS

A **Langmuir isotherm** is characterized by a linear relationship between the solid and solution phases at low values of the latter. As solution phase increases, the solid phase concentration reaches a plateau, corresponding to monolayer protein deposition. The slope of the initial linear portion is proportional to adsorption affinity while the plateau surface concentration represents the surface's capacity

Membrane adsorbenti: PMMA, PEPA, AN69....



CARATTERISTICHE DEI FILTRI IN POLIMETILMETACRILATO (PMMA)



ABSENCE OF OH GROUPS



NO COMPLEMENT ACTIVATION

HYDROFOBICITY AND HIGH THICKNESS OF MEMBRANE



ENDOTOXIN NOT PASSING FROM DIALYSATE SIDE TO BLOOD SIDE

ADSORPTION

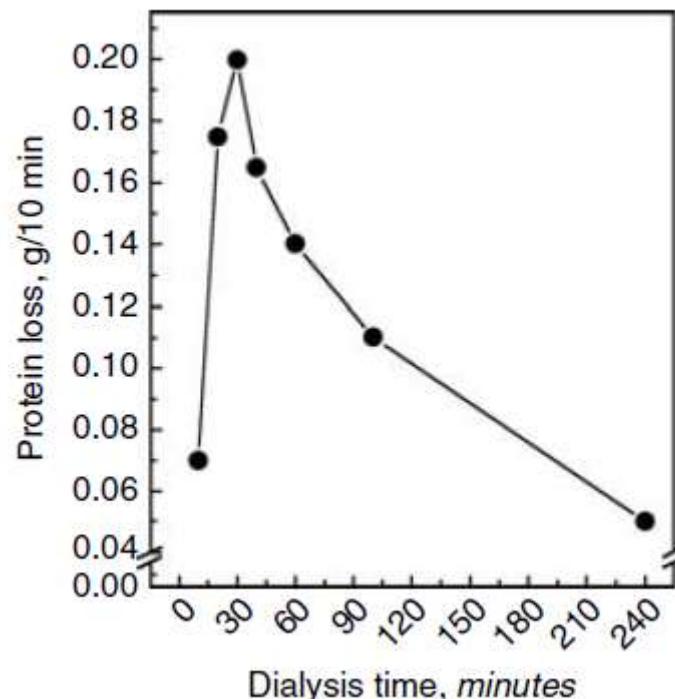
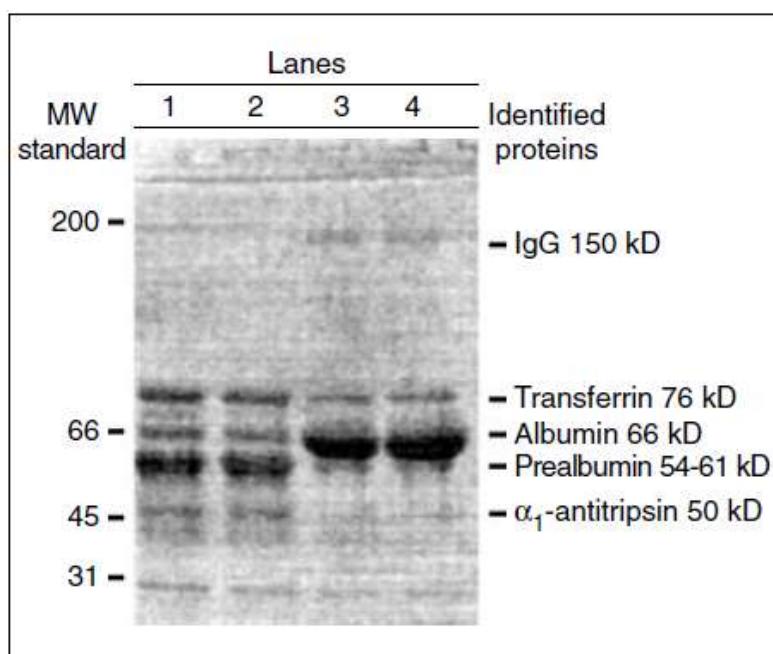


DETENTION OF CYTOKINES PRODUCED BY ACTIVATED MACROPHAGES

PMMA ADSORPTION DIALYSIS

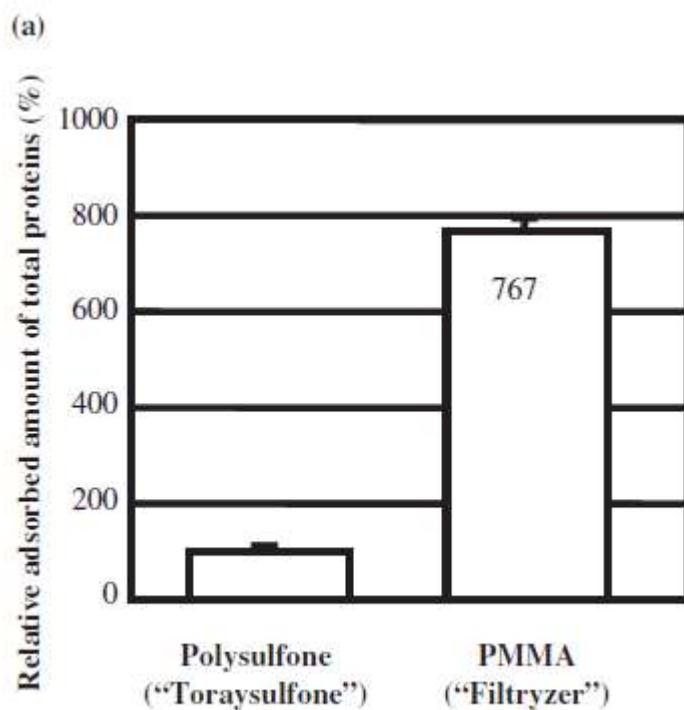
Galli F. et al KI 2005

| | NPLD | PLD |
|-----------------------------------|-----------------------------------|------------------|
| Total proteins g/dialysis session | 0.26 (0.20–0.32) | 2.86 (1.28–4.13) |
| Proteins subfractions MW | Densitometry (% of total O.D.) | |
| α_1 -antitrypsin (50) | 15.5 | 1.0 |
| Prealbumin (54–61) | 31.0 | 7.8 |
| Albumin (66) | 20.7 | 75.6 |
| Transferrin (76) | 17.4 | 10.9 |
| IgG (150) | N.D. | 2.8 |

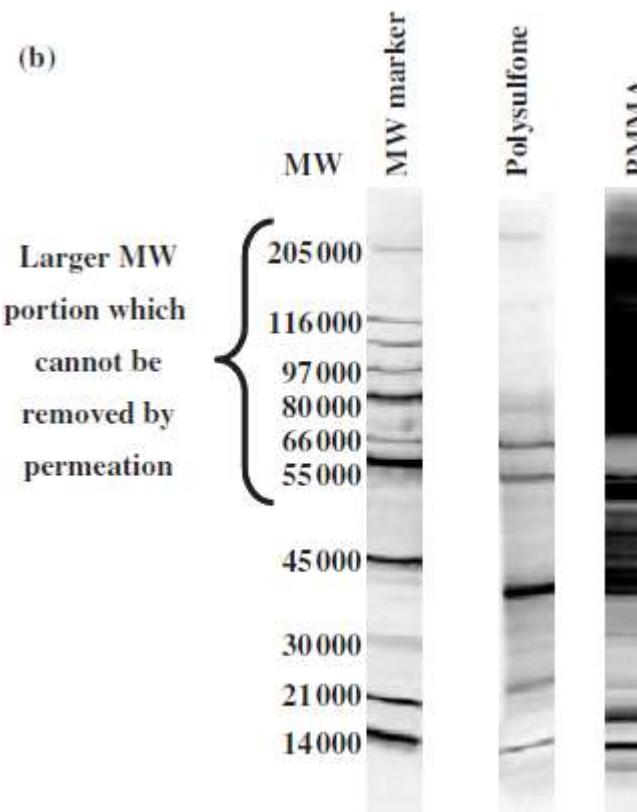


PMMA ADSORPTION DIALYSIS

a) QUANTITA' TOTALE DI PROTEINE ADSORBITE



b) PATTERN ELETTROFORETICO DELLE PROTEINE ASSORBITE



PMMA ADSORPTION DIALYSIS

Table 2. Pre-HD levels of proteins, protein glycation and oxidation indices, cytokines and CRP, adhesion molecules, and MCP-1 in the plasma of patients at baseline and after 6 months (end of the study) of treatment with PLD (study group) and NPLD (control group)

| | Healthy subjects | Control group on NPLD | | Study group on PLD | |
|--|----------------------------|-----------------------|--------------------------|--------------------|----------------------------|
| | | Baseline | 6 months | Baseline | 6 months |
| Plasma proteins and glycoxidation markers | | | | | |
| Plasma proteins g/dL | 7.6 ± 1.3 | 6.7 ± 0.4 | 6.7 ± 0.9 | 6.9 ± 0.5 | 6.8 ± 0.6 |
| Albumin g/dL | 4.5 ± 0.9 | 4.1 ± 0.3 | 3.9 ± 0.8 | 4.0 ± 0.4 | 3.8 ± 0.4 |
| Transferrin mg/dL | 272.4 ± 98.2 | 260.0 ± 81.9 | 252.9 ± 67.7 | 256.4 ± 57.1 | 264.3 ± 68.5 |
| f-pentosidine pmol/mL plasma | N.D. | 110.4 ± 30.2 | 112.5 ± 27.6 | 106.1 ± 24.0 | 60.2 ± 18.9 ^b |
| Total pentosidine pmol/mg protein | 1.4 ± 0.3 ^d | 26.5 ± 8.7 | 25.0 ± 11.1 | 23.5 ± 9.3 | 15.4 ± 5.2 ^b |
| Albumin-pentosidine pmol/mg albumin | 5.4 ± 4.6 ^d | 38.1 ± 16.6 | 37.9 ± 18.0 | 36.3 ± 14.2 | 23.8 ± 8.9 ^b |
| Protein carbonyls arbitrary units | 39.1 ± 30.2 ^d | 260.5 ± 165.6 | 245.2 ± 177.0 | 229.8 ± 156.8 | 132.2 ± 117.5 ^b |
| AOPP arbitrary units | 12.7 ± 14.0 ^d | 100.3 ± 68.6 | 114.1 ± 78.3 | 104.7 ± 60.2 | 64.6 ± 56.6 ^b |
| Cytokines and CRP | | | | | |
| IL-1β pg/mL | 2.7 ± 1.1 ^d | 4.4 ± 3.0 | 4.8 ± 2.2 | 4.1 ± 3.3 | 2.1 ± 1.9 ^{a,e} |
| TNF-α pg/mL | 9.1 ± 3.2 ^d | 19.0 ± 4.0 | 21.0 ± 5.5 | 18.7 ± 4.3 | 15.1 ± 3.1 ^b |
| IL-6 pg/mL | 2.4 ± 0.9 ^d | 5.3 ± 2.1 | 5.8 ± 2.3 | 5.0 ± 1.9 | 3.1 ± 0.6 ^{b,f} |
| IL-10 pg/mL | 6.0 ± 2.5 ^d | 3.3 ± 0.4 | 3.0 ± 0.5 | 3.0 ± 0.6 | 5.0 ± 4.4 ^{b,e} |
| INF-γ pg/mL | 12.7 ± 3.2 ^c | 9.9 ± 1.5 | 10.9 ± 1.9 | 9.2 ± 1.6 | 12.6 ± 1.7 ^b |
| CRP mg/L | 7.9 ± 3.6 ^d | 25.8 ± 28.6 | 27.4 ± 24.0 | 22.7 ± 33.9 | 12.1 ± 9.1 |
| Adhesion molecules and MCP-1 | | | | | |
| ICAM-1 ng/mL | 232.5 ± 89.6 | 283.4 ± 65.3 | 282.9 ± 96.5 | 270.7 ± 77.8 | 240.8 ± 67.6 |
| VCAM-1 ng/mL | 589.0 ± 171.4 ^d | 973.6 ± 435.9 | 938.5 ± 259.9 | 941.0 ± 458.2 | 772.3 ± 285.4 |
| Selectin-E ng/mL | 48.4 ± 16.5 | 63.5 ± 30.5 | 68.2 ± 26.9 ^e | 58.2 ± 27.3 | 63.2 ± 36.7 |
| MCP-1 pg/mL | 207.3 ± 91.5 ^d | 420.1 ± 154.9 | 468.9 ± 182.5 | 408.8 ± 170.2 | 381.5 ± 141.5 |

N.D., not detectable. Data are mean ± 1 SD.

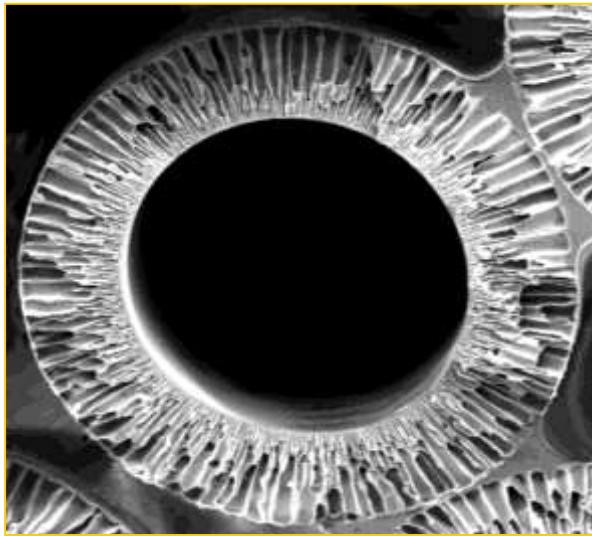
^aP < 0.05 and ^bP < 0.01 vs. baseline values.

^cP < 0.05 and ^dP < 0.01 vs. patients' data (when not otherwise specified).

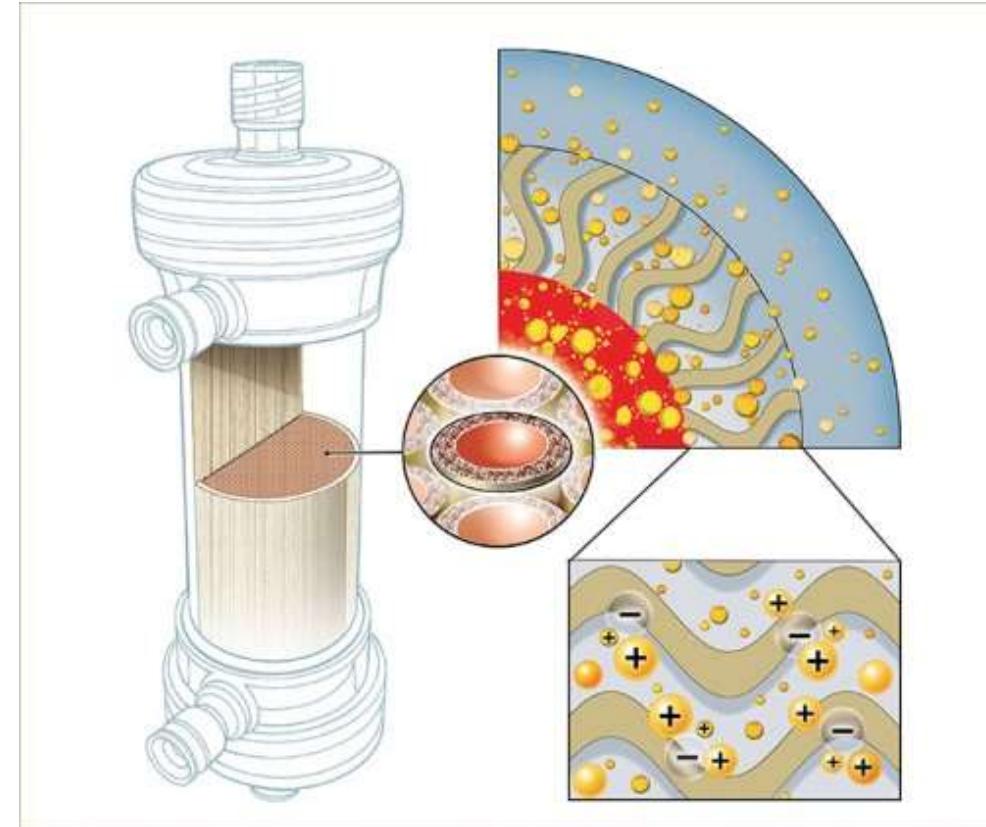
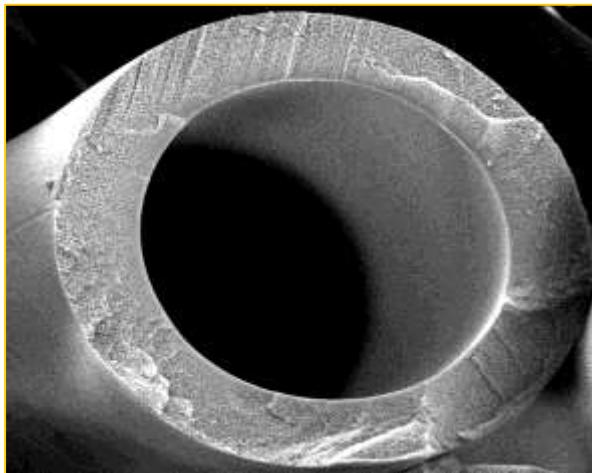
^eP < 0.05 and ^fnot significant vs. healthy control subject data.

POLIMETILMETACRILATO (PMMA)

POLISULFONE



PMMA



Membrane e sopravvivenza

RESEARCH ARTICLE

Effect of dialyzer membrane materials on survival in chronic hemodialysis patients: Results from the annual survey of the Japanese Nationwide Dialysis Registry

Masanori Abe^{1*}, Takayuki Hamano², Atsushi Wada³, Shigeru Nakai⁴, Ikuto Masakane⁵, on behalf of the Renal Data Registry Committee, Japanese Society for Dialysis Therapy¹

142,412 patients on maintenance hemodialysis considered (from 2008 to 2009).

Patients treated with:

cellulose triacetate (**CTA**), ethylene vinyl alcohol (**EVAL**), polyacrylonitrile (**PAN**), polyester polymer alloy (**PEPA**), polyethersulfone (**PES**), polymethylmethacrylate (**PMMA**), and polysulfone (**PS**).

>>> 1 YEAR All Cause Mortality analysis (Cox regression) to estimate the possible association with baseline membrane dialyzers use.

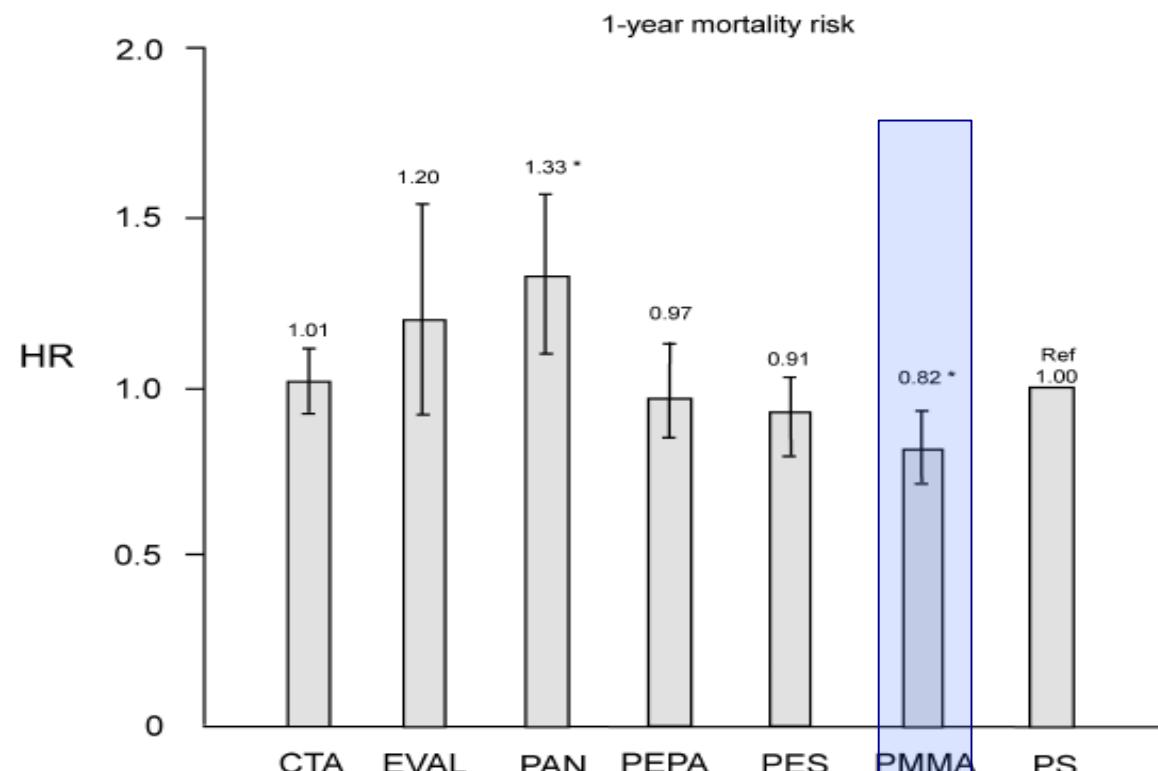


Fig 3. HRs of all-cause mortality after propensity score matching for six types of dialyzer groups compared to the PS group using Cox proportional hazards regression. *P < 0.01 vs. PS. CTA, cellulose triacetate; EVAL, ethylene vinyl alcohol; HR, hazard ratio; PAN, polyacrylonitrile; PEPA, polyester polymer alloy; PES, polyethersulfone; PMMA, polymethylmethacrylate; PS, polysulfone.

Infiammazione in uremia e CVD

Review Article

Biomarkers of Chronic Inflammatory State in Uremia and Cardiovascular Disease

International Journal of Inflammation

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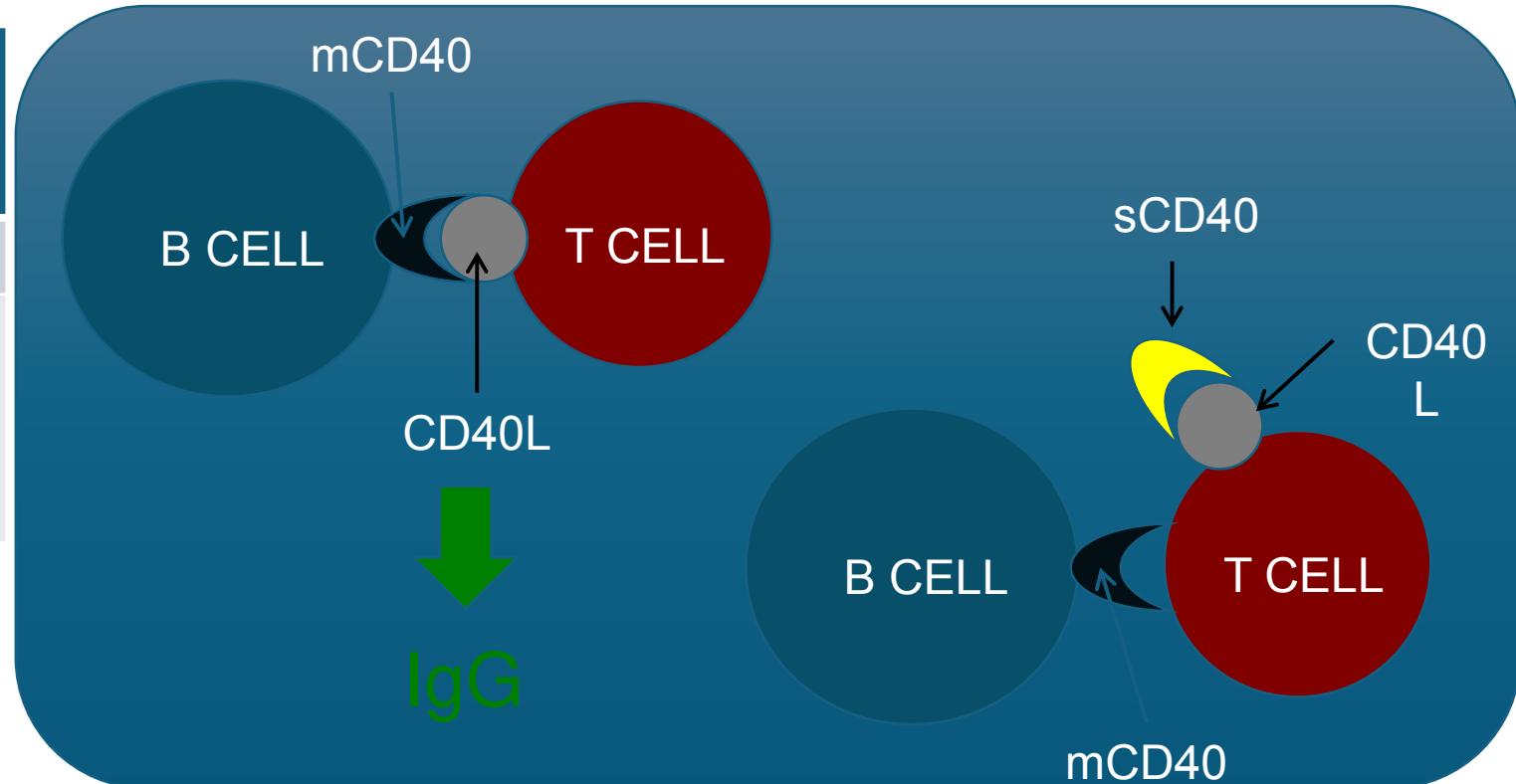
Vincenzo Panichi, Alessia Scatena, Massimiliano Migliori,
Valentina Marchetti, Sabrina Paoletti, and Sara Beati

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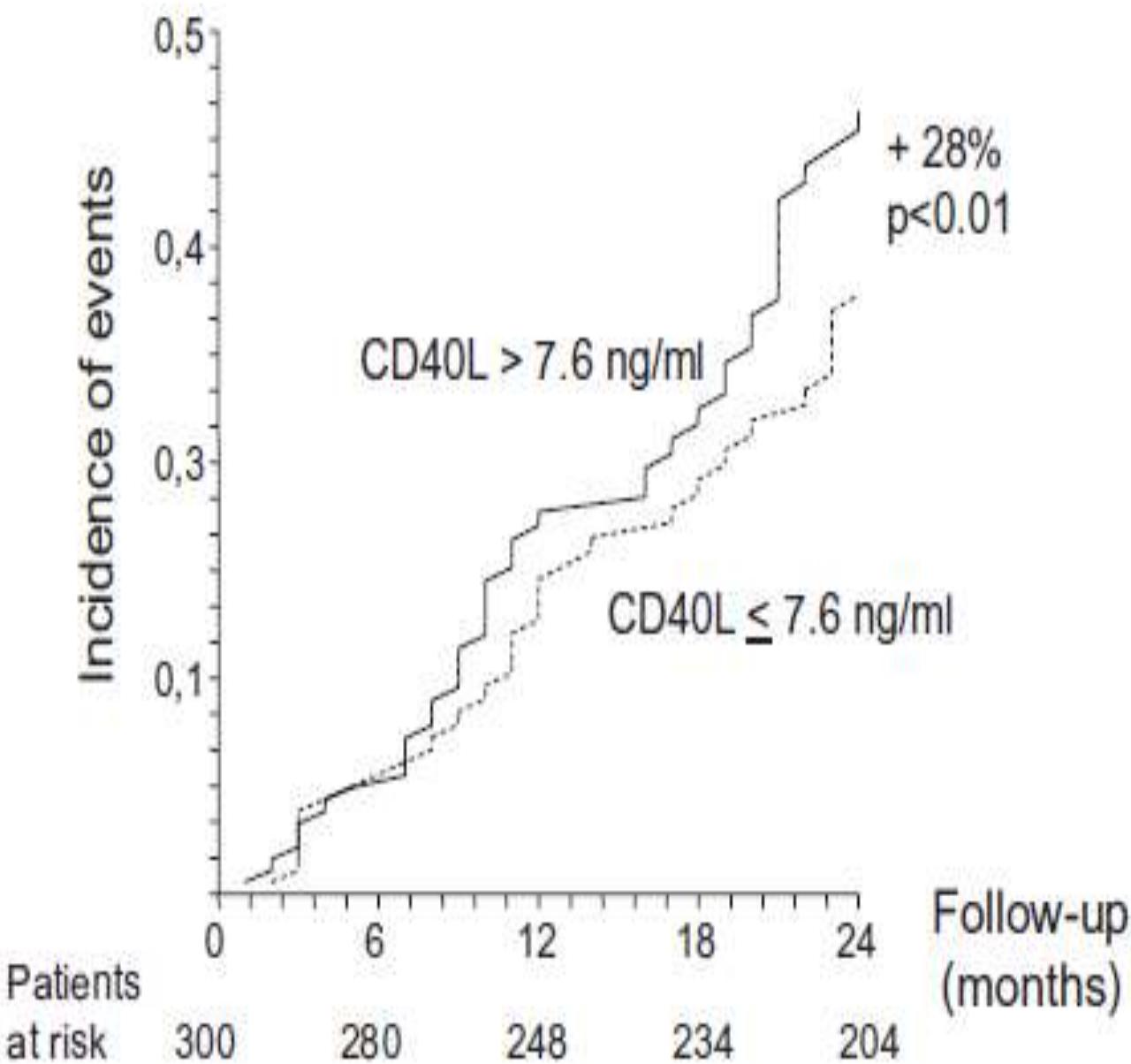
Correspondence should be addressed to Vincenzo Panichi, vpanichi@usl12.toscana.it

Received 6 March 2012; Revised 5 April 2012; Accepted 17 April 2012

| TRADITIONAL MARKERS | IL-6 | CRP |
|-------------------------|---------------------|-------|
| | | |
| NON TRADITIONAL MARKERS | CD40/CD40 L complex | PTX-3 |

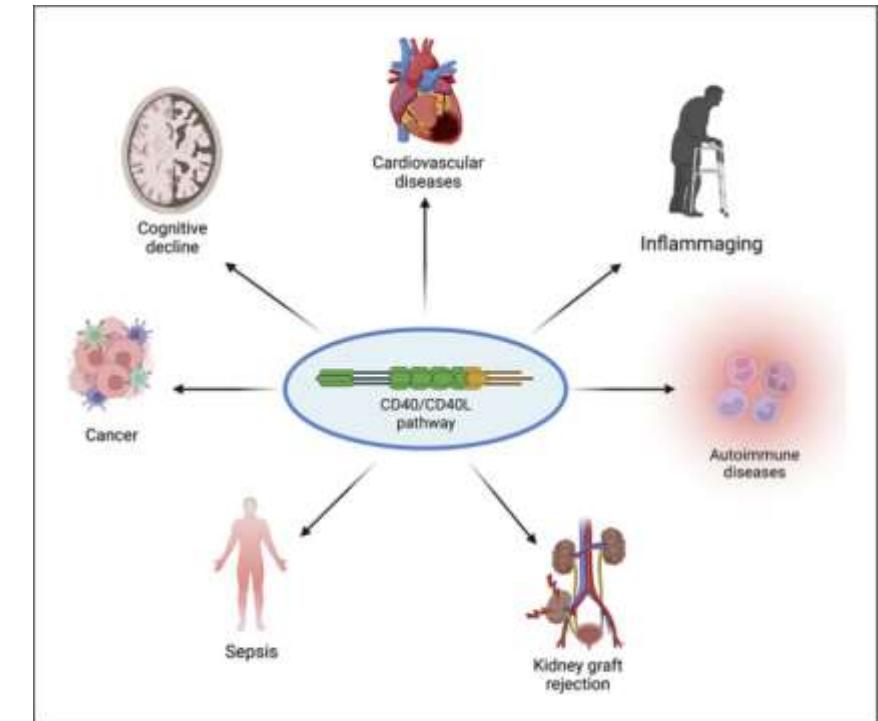
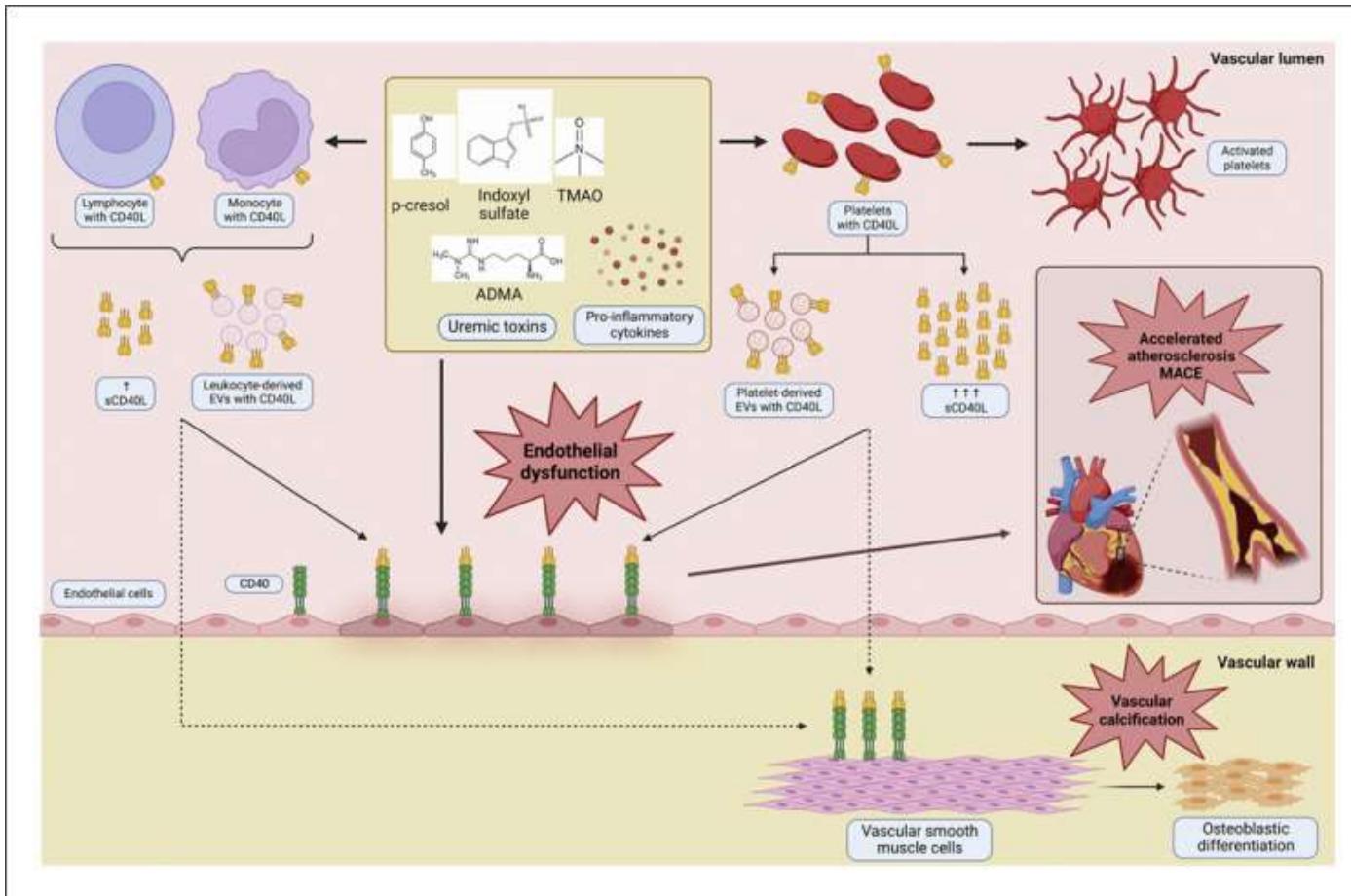


Il fattore Solubile CD40 Ligando predittivo di mortalità e morbidità cardio vascolare



the prognostic value of sCD40L as factor of cardiovascular Morbidity and Mortality is evident also in over 200 chronic HD patients from the RISCAVID population at 24-month follow-up

Role of the CD40-CD40 Ligand Pathway in Cardiovascular Events, Neurological Alterations, and Other Clinical Complications of Chronic Hemodialysis Patients: Protective Role of Adsorptive Membranes



HBV and sCD40

Immunology 2003 110 131–140

Potential role of soluble CD40 in the humoral immune response impairment of uraemic patients

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JEAN-FRANÇOIS MOREAU,* PIERRE MERVILLE*† & JULIE DÉCHANET-MERVILLE* *UMR-CNRS 5540,
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Bordeaux, France and ‡INSERM U404, Lyon, France

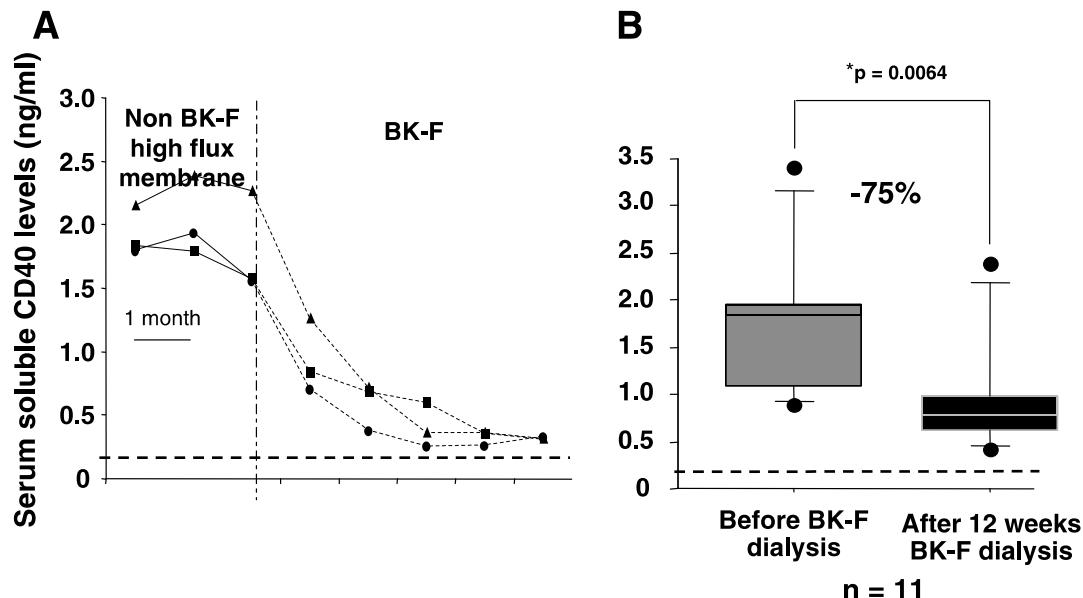


Fig. 3. Longitudinal follow-up of sCD40 levels in patients dialysed on non-PMMA high-flux membranes or BK-2.1F membranes. (A) Soluble CD40 concentrations were measured monthly by ELISA in the serum of three patients dialysed on non-PMMA high-flux membrane and who were then switched to BK-2.1F membrane. Blood samples were taken after the dialysis session. Dotted line represents mean level of sCD40 in healthy subjects. (B) Levels of sCD40 in the serum of 11 patients before and after 12 weeks of dialysis on BK-2.1F membrane. *Non-parametric Wilcoxon *U*-test.

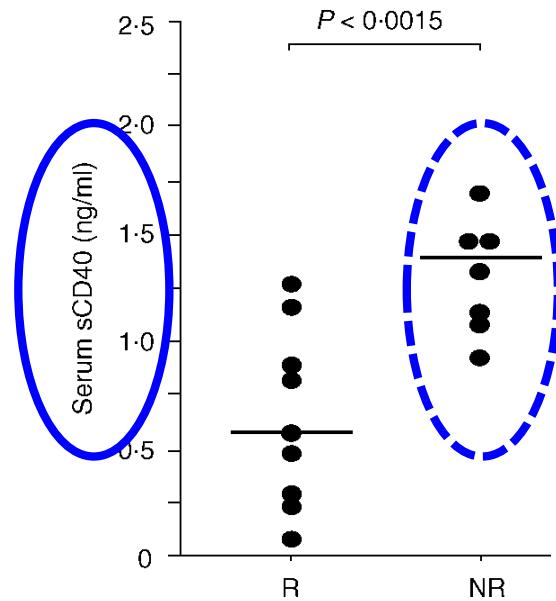


Figure 6. Serum sCD40 levels in chronic haemodialysed patients in the course of Hepatitis B vaccination and correlation to their vaccinal response status. Two groups of haemodialysed patients were set up according to their response to hepatitis B vaccination. Responsive patients (R, $n = 9$) presented >10 IU/l of anti-HBs IgG one month after last vaccine injection whereas Non-responsive patients (NR, $n = 8$) presented <10 IU/l. sCD40 values of each patients correspond to the mean of the four sCD40 values obtained during the vaccination. Lines represent the median values of sCD40 for all the patients of each group. $*P < 0.0015$ determined using a Mann-Whitney test.

Ralli Chiara¹, Imperiali Patrizio¹, Gabbrilelli Claudio², Conti Paolo², Lombardi Marco³, Sidoti Antonino⁴, Capitanini Alessandro⁵, Piluso Adriano⁶, Tekle Kiro Seble⁶, Duranti Diletta⁷ and Duranti Ennio^{1*}

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Keywords: Hemodialysis; Immune dysfunctions;

Dates: Received: 10 December, 2015; Accepted: 20 January, 2016; Published: 21 January, 2016

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Research Article

Hemodialysis with Polymethylmethacrylate Restores the Response to Hepatitis B Vaccination in Chronic Dialysis Patients: Hypothesized Mechanism of Action

Abstract

Patients undergoing hemodialysis often present with a reduced response to anti-hepatitis B virus (anti-HBV) vaccination. The soluble form of CD40 (sCD40) is elevated in hemodialysis patients and this has been shown to correlate with lack of response to anti-HBV vaccination. Due to its high molecular weight, conventional dialyzers cannot clear sCD40. Previous studies have demonstrated, that dialysis membranes in polymethylmethacrylate (PMMA) can reduce the levels of sCD40. We have studied the effect of dialysis with PMMA membranes in patients who were non-responders to anti-HBV vaccination after a complete cycle of vaccinations. Interestingly, we found that significantly more patients in the PMMA group were able to mount a response to vaccination, compared to the control group ($P = 0.04$).



Journal of Chronic Diseases and Management

Short Notes

Adsorptive Hemodialysis by Polymethylmethacrylate (PMMA): an update on Hepatitis B Vaccination Immunoresponce

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OPEN ACCESS



Potential role of the soluble form of CD40 in deficient immunological function of dialysis patients: new findings of its amelioration using polymethylmethacrylate (PMMA) membrane

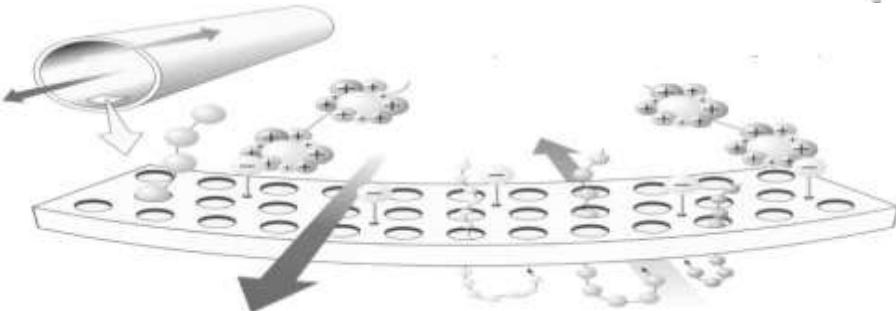
Cécile Contin-Bordes^{1,2}, Adeline Lacraz³ and Valérie de Précigout³

| | Total | Controls | PMMA |
|----------------------------------|-----------------------|-----------------------|------------------|
| Patients | 32 | 15 | 17 |
| Average age | 73 ± 12 | 78 ± 9 | 67 ± 15 |
| Gender (M/F) | 18/14 | 8/7 | 10/7 |
| Dry weight (kg) | 74 ± 21 | 75 ± 17 | 73 ± 26 |
| Dialysis vintage (months) | 75 ± 58 | 97 ± 67 | 54 ± 48 |
| Type of dialysis | Bicarbonate | Bicarbonate | Bicarbonate |
| Membrane | Polysulfone-polyamide | Polysulfone-polyamide | PMMA series BK-F |
| HBsAb (UI/L) | < 10 | < 10 | < 10 |

| | Controls | PMMA | X ² test |
|------------------------------------|-------------|-------------|---------------------|
| Patients | 15 | 17 | |
| Patients with HBsAb > 10 | 2 | 8 | |
| Percentage of responders | 13 % | 47 % | 0.04 |

Uraemic itching: do polymethylmethacrylate dialysis membranes play a role?

Filippo Aucella^{1,2}, Mimmo Vigilante¹, Antonio Gesuete¹, Gianfranco Maruccio¹, Angelo Specchio¹ and Loreto Gesualdo⁵



F. Aucella *et al.*

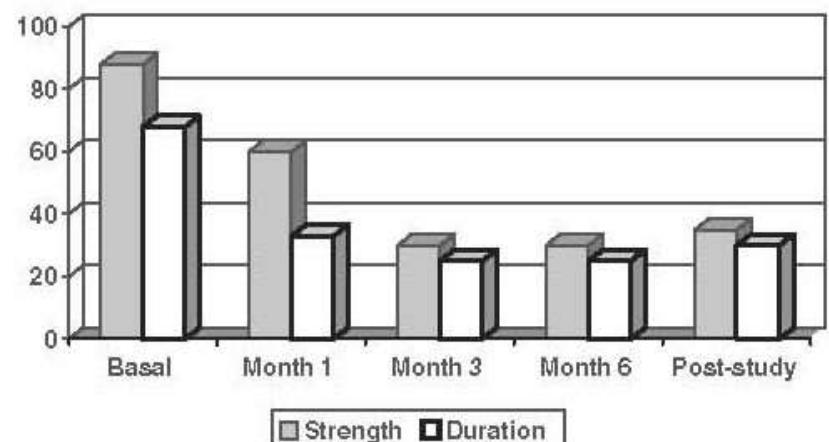


Fig. 3. Effect of BG-U (PMMA membrane) on the subjective assessment of uraemic itch: average values of eight patients who had previously been dialysed using low-flux membranes.



IL PRURITO UREMICO: UNA SFIDA NON RISOLTA

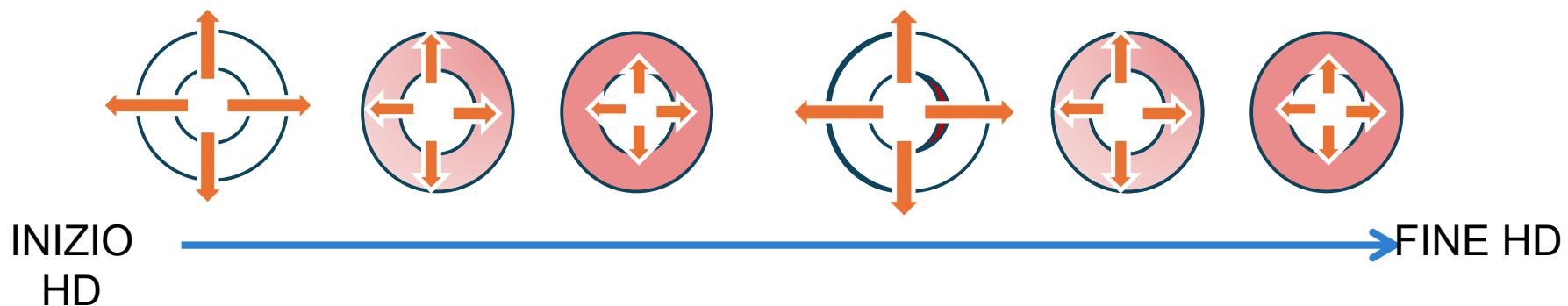
F. Aucella, A. Gesuete

Struttura Complessa di Nefrologia e Dialisi, Ospedale "Casa Sollievo della Sofferenza, Istituto di Ricovero e Cura a Carattere Scientifico, San Giovanni Rotondo [FG]

TABELLA III - ALGORITMO DI TRATTAMENTO DI UP

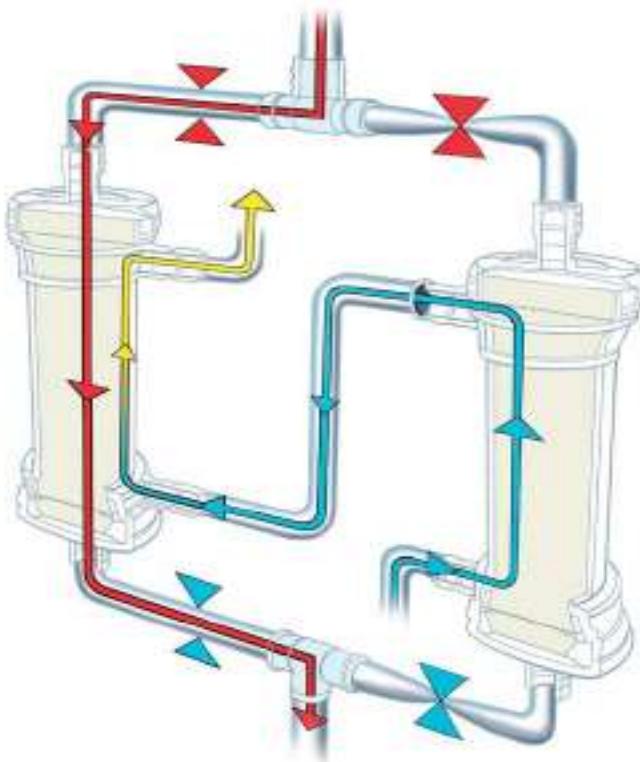
| Steppe | Possibili opzioni terapeutiche | Livello di evidenza | Nomi commerciali e dose |
|--------|--|---------------------|--|
| STEP 1 | <ul style="list-style-type: none"> Ottimizzare la dialisi: efficienza, biocompatibilità, uso di membrane in PMMA Controllo del metabolismo calcio-fosforo Controllo iperparatiroidismo secondario | 3 | filtri serie BG-U, Estor |
| STEP 2 | <ul style="list-style-type: none"> Emollienti ad alto contenuto di acqua | 3 | Lenoxiol fluido, Leviax lozione, Cetafil crema; 2 applicazioni al dì. |
| STEP 3 | <ul style="list-style-type: none"> Creme alla Capsaicina allo 0,025% (off-label) Capsule o creme a base di acido γ-linolenico | 1 | Cerotto Bertelli, Dolpyc, Capsolin; 2 volte al dì. Primrose oil, 2 cps/die o Eucerin crema, 2 volte al dì |
| STEP 4 | <ul style="list-style-type: none"> Ciclo di terapia fisica con UVB (a banda ristretta) | 1 | 3 sedute a dì alterni, indi 1 a settimana, sino a n. 9 sedute totali |
| STEP 5 | <ul style="list-style-type: none"> Gabapentin post-dialisi (off-label) | 1 | Neurontin, Gabapentin Teva, Gabapentin Sigma Tau; 100 mg post dialisi e successiva titolazione |
| STEP 6 | <ul style="list-style-type: none"> Carbone attivo per os | 1 | 6 gr/die per os |
| STEP 7 | <ul style="list-style-type: none"> Nalfurafina ev o Butorfanolo intranasale | 2 | non ancora in commercio in Italia |

ENHANCED ADSORPTION DIALYSIS

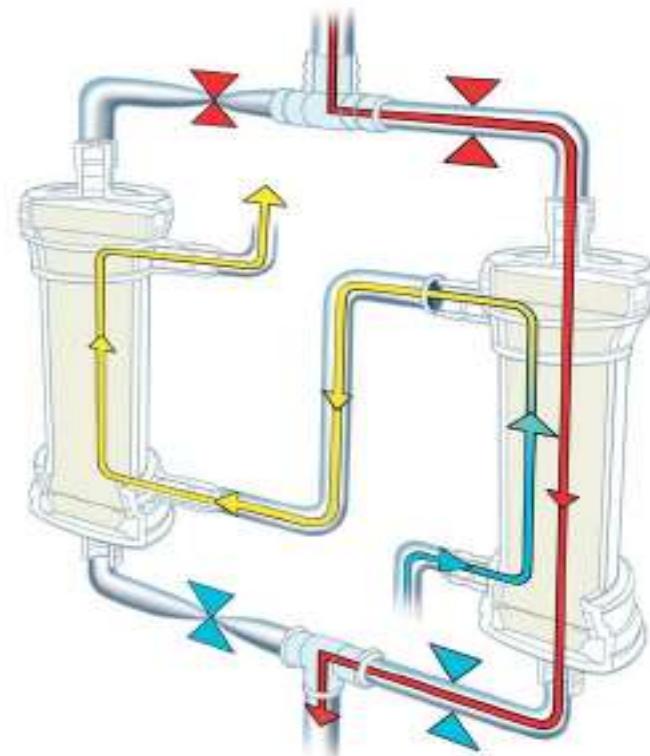


ENHANCED ADSORPTION DIALYSIS

PRIME DUE ORE



ULTIME DUE ORE



Role of light chain clearance in the recovery of renal function in multiple myeloma: another point of view

Acute kidney injury (AKI) in patients with multiple myeloma (MM) requiring renal replacement treatment (RRT) is associated with high morbidity and mortality. Early reduction of serum free light chains (FLC) with adjuvant intensive hemodialysis (IHD) may improve renal outcomes. We aim to evaluate the impact on kidney function of IHD (PMMA dialysis membranes or HFR) vs. standard dialysis.

Methods



Multicentric retrospective study



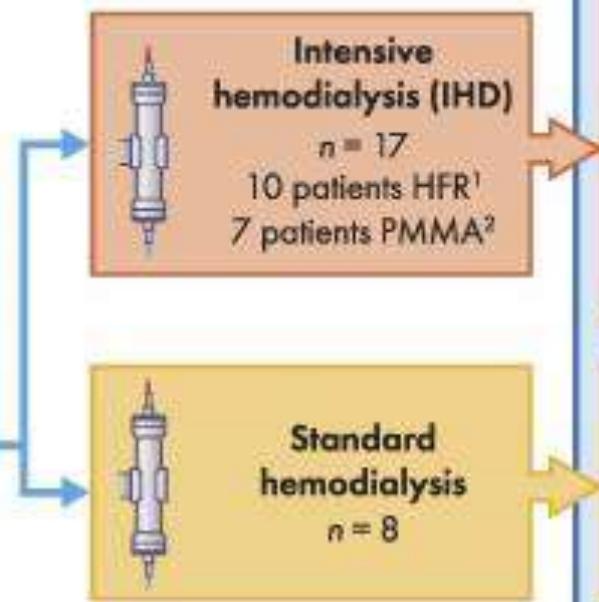
25 patients (2011–2018)



AKI related to de novo or first relapsed MM
Need of RRT



Same targeted MM therapy
(84% bortezomib-based therapy)



¹HFR: hemodiafiltration with endogenous reinfusion

²PMMA: polymethylmethacrylate membranes

Results

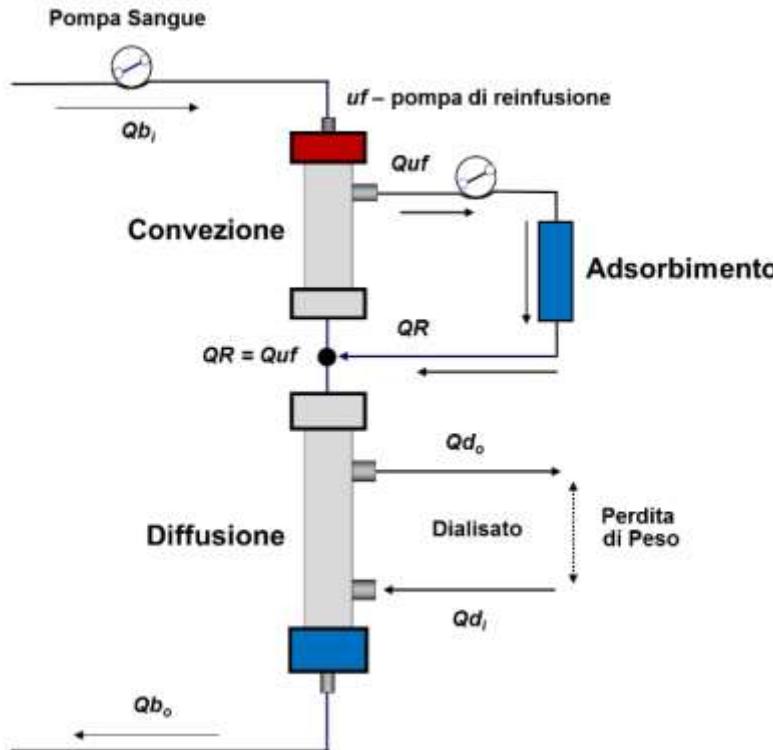
76.5% patients dialysis-free at 6 months
13 patients recovered kidney function
Median time to be dialysis-free: 21 days (IQR 14–55)
Average creatinine: 1.75 mg/dL

12.5% patients dialysis-free at 6 months
Only 1 patient recovered kidney function

Conclusion: Early reduction of FLC with IHD as an adjuvant treatment along with MM targeted therapy may exert a positive impact on renal recovery.

Ramos, N.
Clinical Kidney Journal (2023)
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[@CKJsocial](https://twitter.com/CKJsocial)

HFR – HFR SUPRA



Evidence That p-Cresol and IL-6 Are Adsorbed by the HFR Cartridge: Towards a New Strategy to Decrease Systemic Inflammation in Dialyzed Patients?

Eleonora Riccio¹, Mauro Cataldi^{2,,}, Lucia Grumetto³, Loredana Postiglione⁴, Bruna Guida⁵, Bruno Maristella Minco¹, Gennaro Argentino¹, Roberta Russo¹, Stefania Brancaccio¹, Andrea Memoli¹Memoli^{1*}

PLOS ONE. April 2014 | Volume 9 | Issue 4

| | Before HFR | After HFR | Reduction Ratio (%) | Before HD | After HD | Reduction Ratio (%) |
|----------------------------|------------|-----------|---------------------|-----------|-----------------------|------------------------|
| Albumin (g/dl) | 3.6±0.2 | 4.2±0.3 | - | 3.8±0.3 | 4.2±0.2 | - |
| Total Cresol (mg/l) | 11.6±6.3 | 5.8±2.7* | 53.6±12.5 | 8.6±5.3 | 5.4±3.0 ^{II} | 37.1±20.2 ^I |
| IL-6 (pg/ml) | 57.6±58.3 | 48.4±53.1 | 4.3±34.5 | 48.7±33.6 | 53.9±48.3 | - |

La mortalità della popolazione uremica in trattamento sostitutivo permane inaccettabilmente alta principalmente per cause cardiovascolari. È noto infatti che oltre i fattori di rischio tradizionali il paziente uremico presenta fattori non tradizionali come l'accumulo di tossine uremiche e una condizione di aumentata infiammazione



PICCOLO PESO MOLECOLARE
GRANDE PESO MOLECOLARE

MEDIO PESO MOLECOLARE.
TOSSICI LEGATI ALLA PROTEINE

Criteri di inclusione

- Età > 18 anni
- Età dialitica > 6 mesi
- Portatore di FAV con Q_B di almeno 300-350 ml/min
- Consenso informato

Cross over

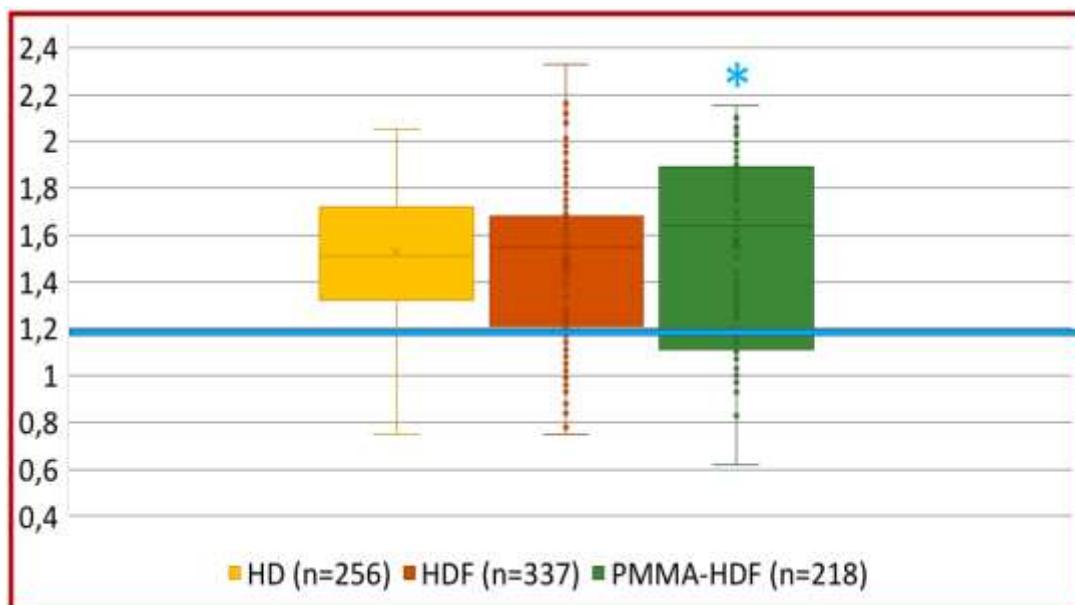
| | |
|-------|---------------------|
| Gr. A | HD → HDF → PMMA-HDF |
| Gr. B | HDF → PMMA-HDF → HD |
| Gr. C | PMMA-HDF → HD → HDF |

Prelievi

| | |
|-------------|---|
| A seduta | Kt/V; Totale quota convettiva; durata metodica |
| Mensili | Hb; Ferritina; PCR; TnT; P; Urea; ALB |
| Trimestrali | Tossine: sFLC; β 2m; p-Cresol; p-Indoxyl Marker di flogosi: Epcidina; IL-6; TNF α ; CD40L; NGAL |

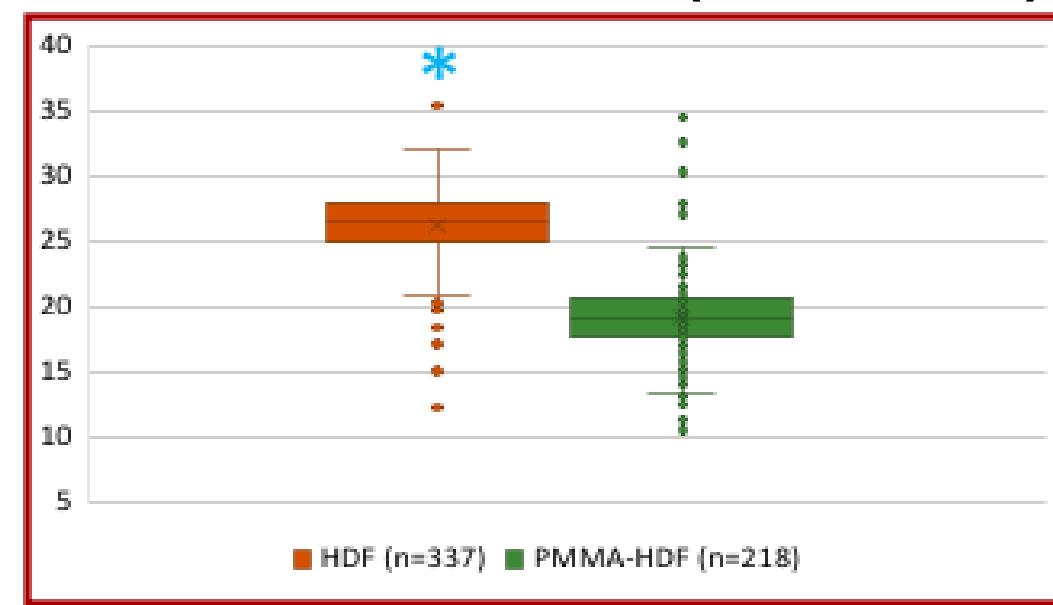
| | HD (n=256) | HDF (n=337) | PMMA-HDF (n=218) |
|--------------------------------|-------------------|--------------------|-----------------------------|
| Durata seduta (min) | 224 | 234 | 222 |

Kt/V



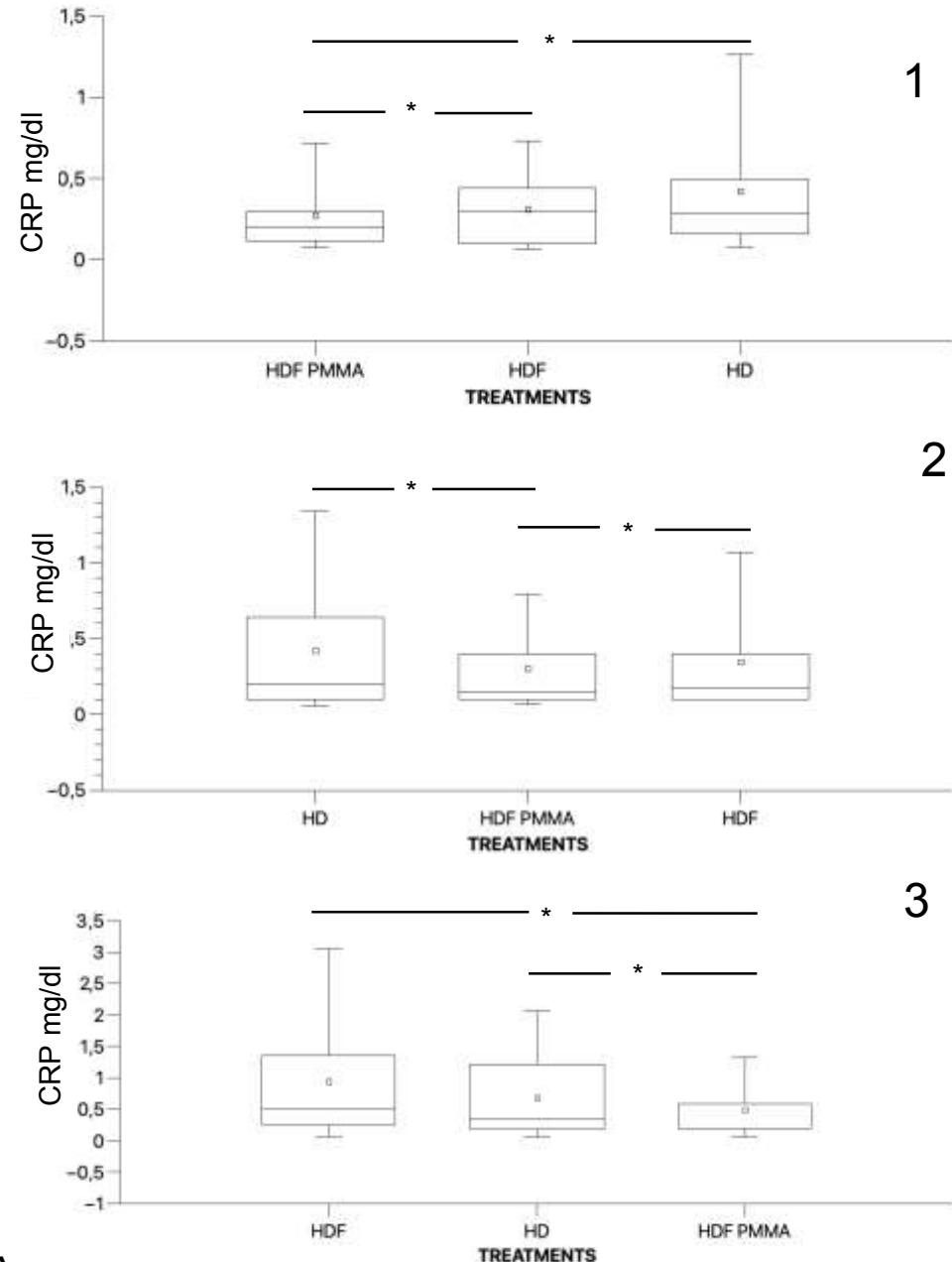
* $p < 0.001$ vs HDF

Volume convettivo (L / seduta)



* $p < 0.05$ vs PMMA-HDF

A

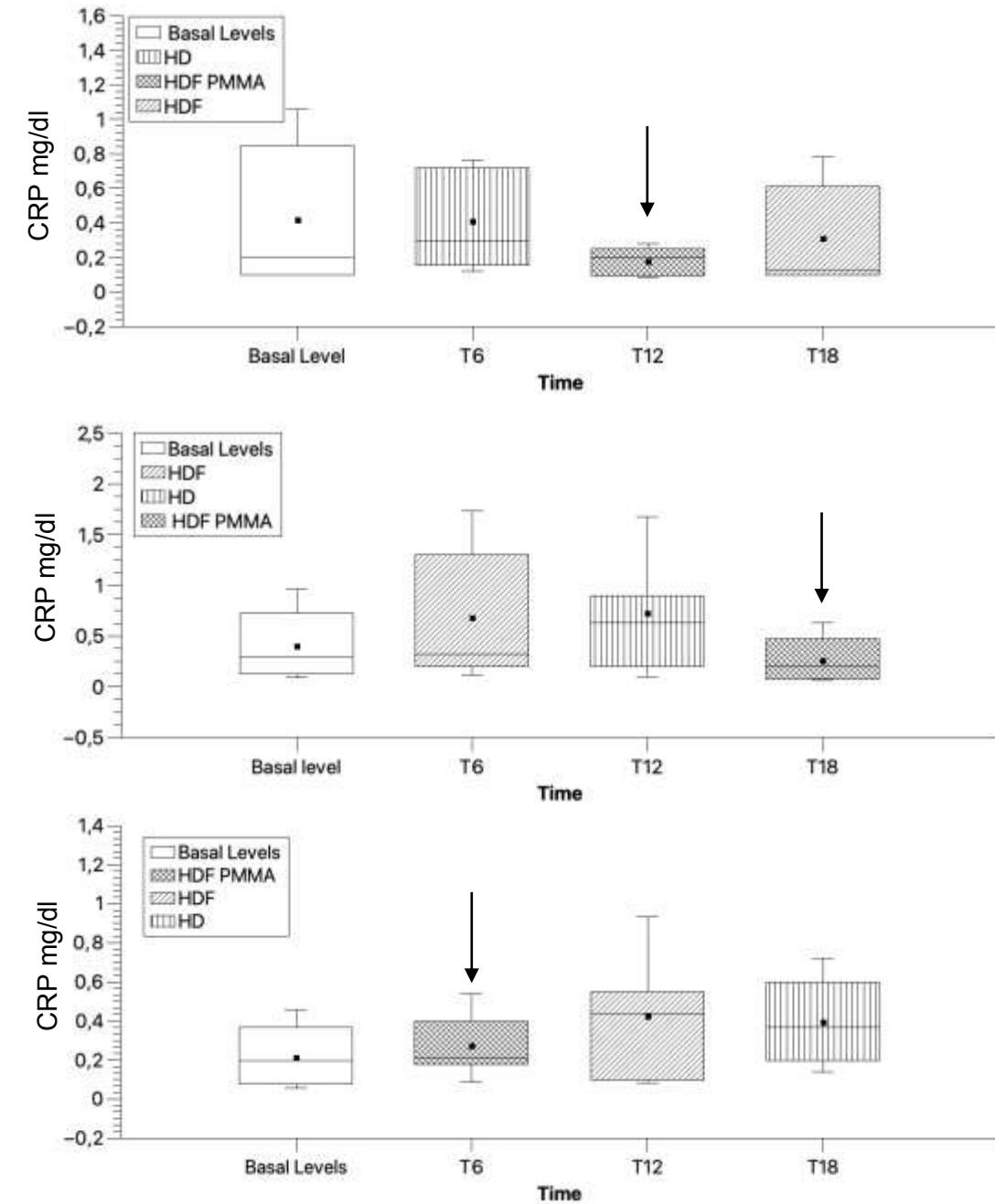


1

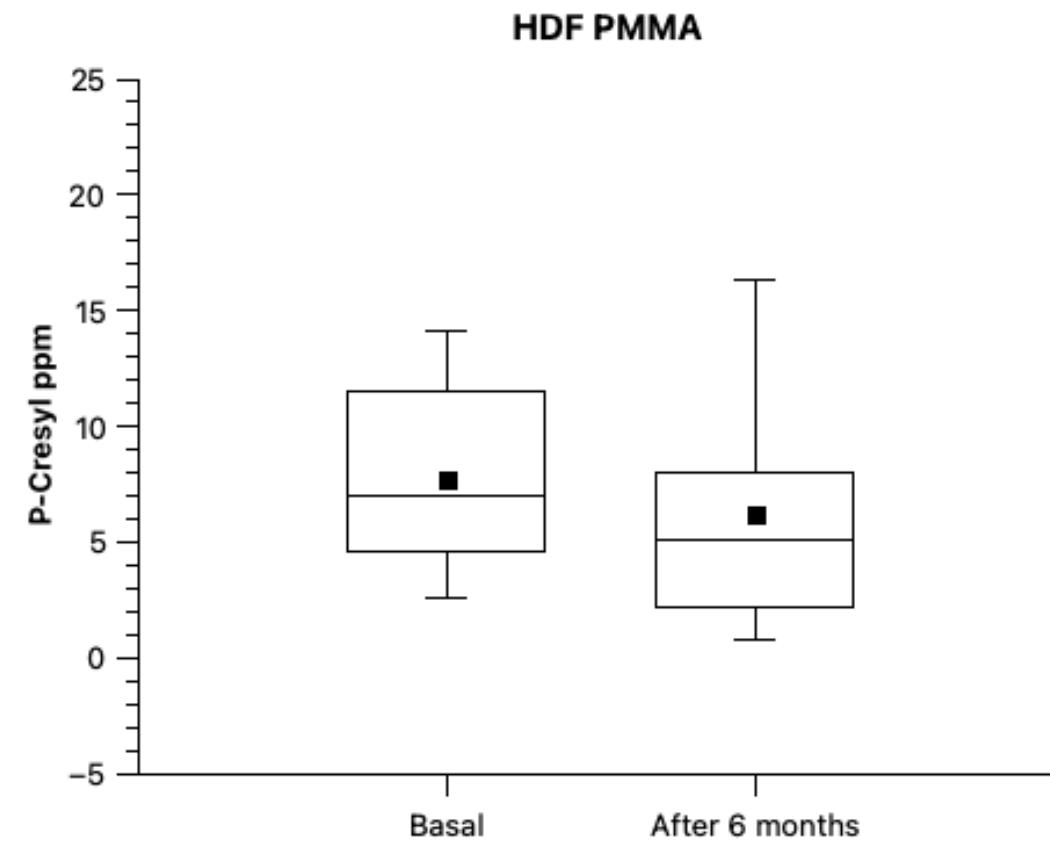
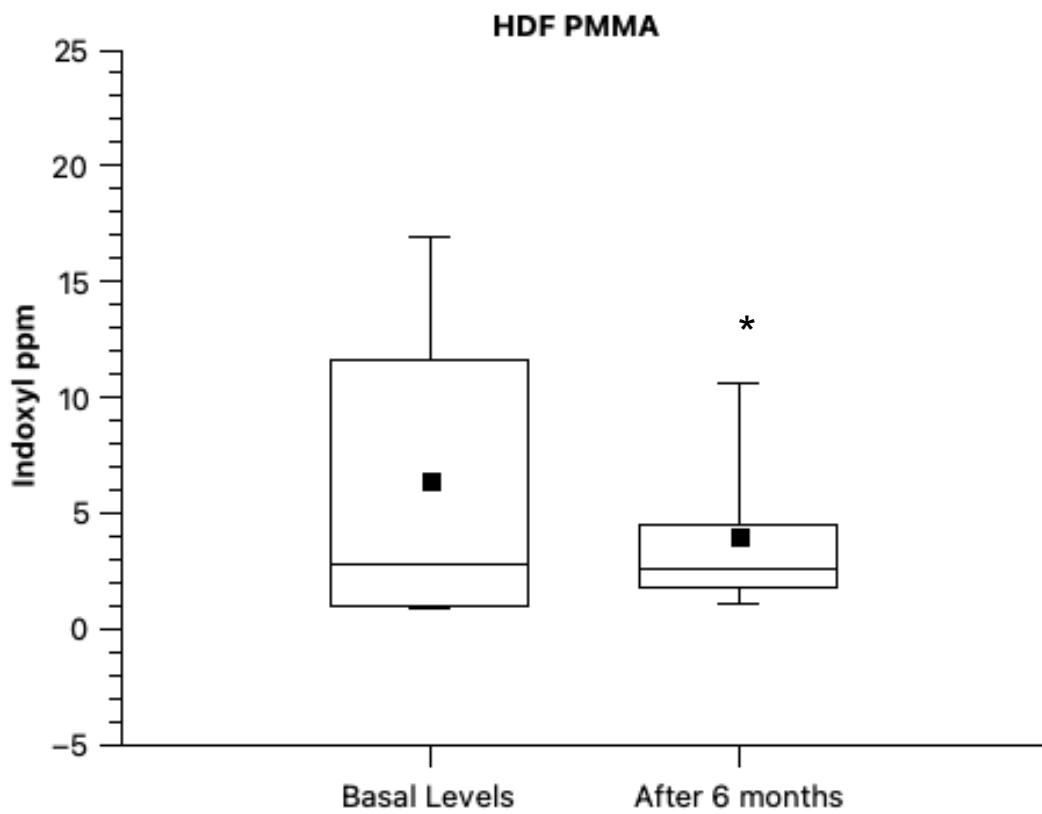
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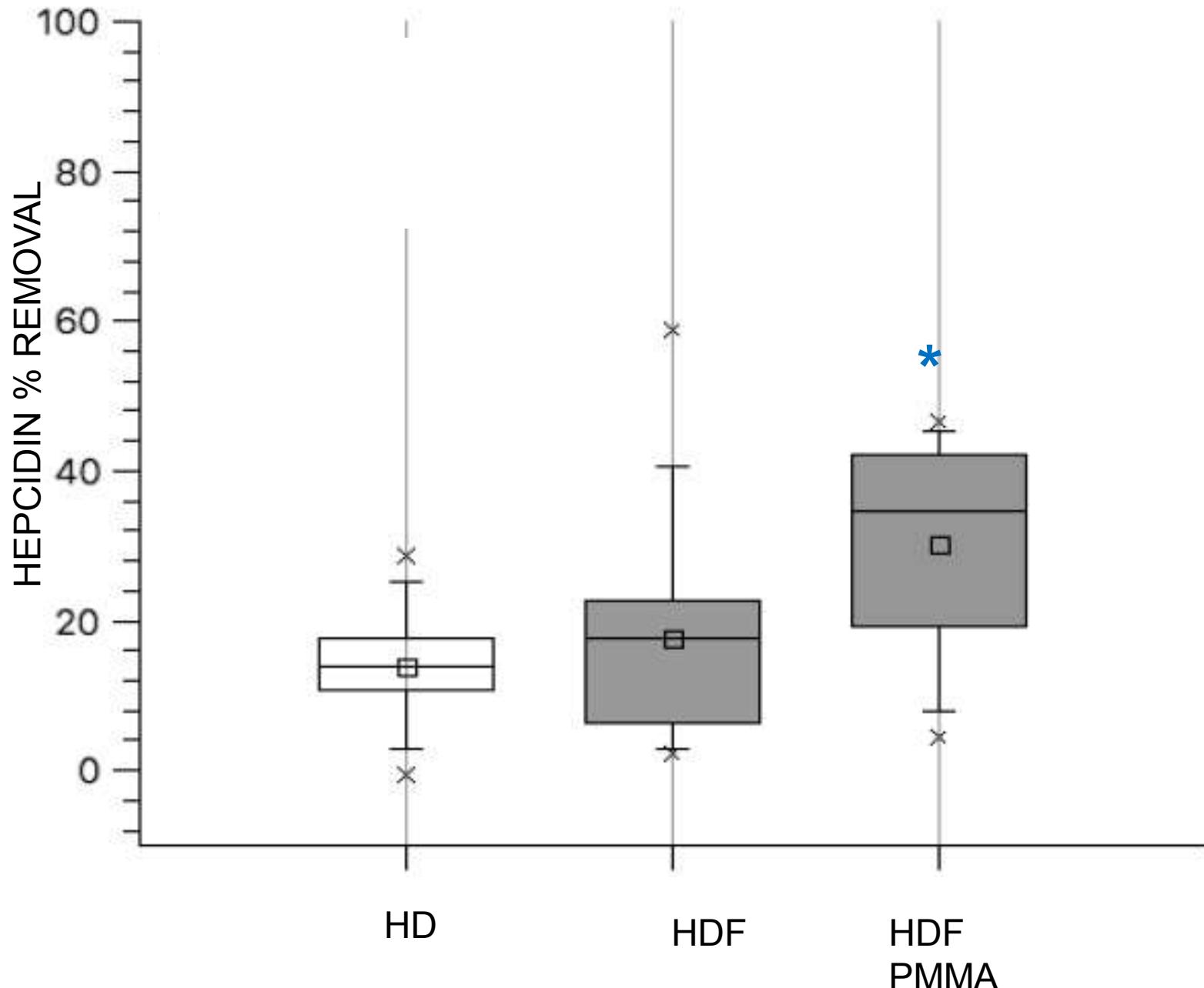
3

B



EFFETTO SULLE PBUT





LA HDF PMMA
ERA LA
METODICA PIÙ
EFFICEINTE NEL
RIMUOVERE
HEPCIDINA NELLA
SINGOLA SEDUTA

NUOVE TIPOLOGIE DI PMMA: RIDOTTA L'ATTIVAZIONE PIASTRINICA

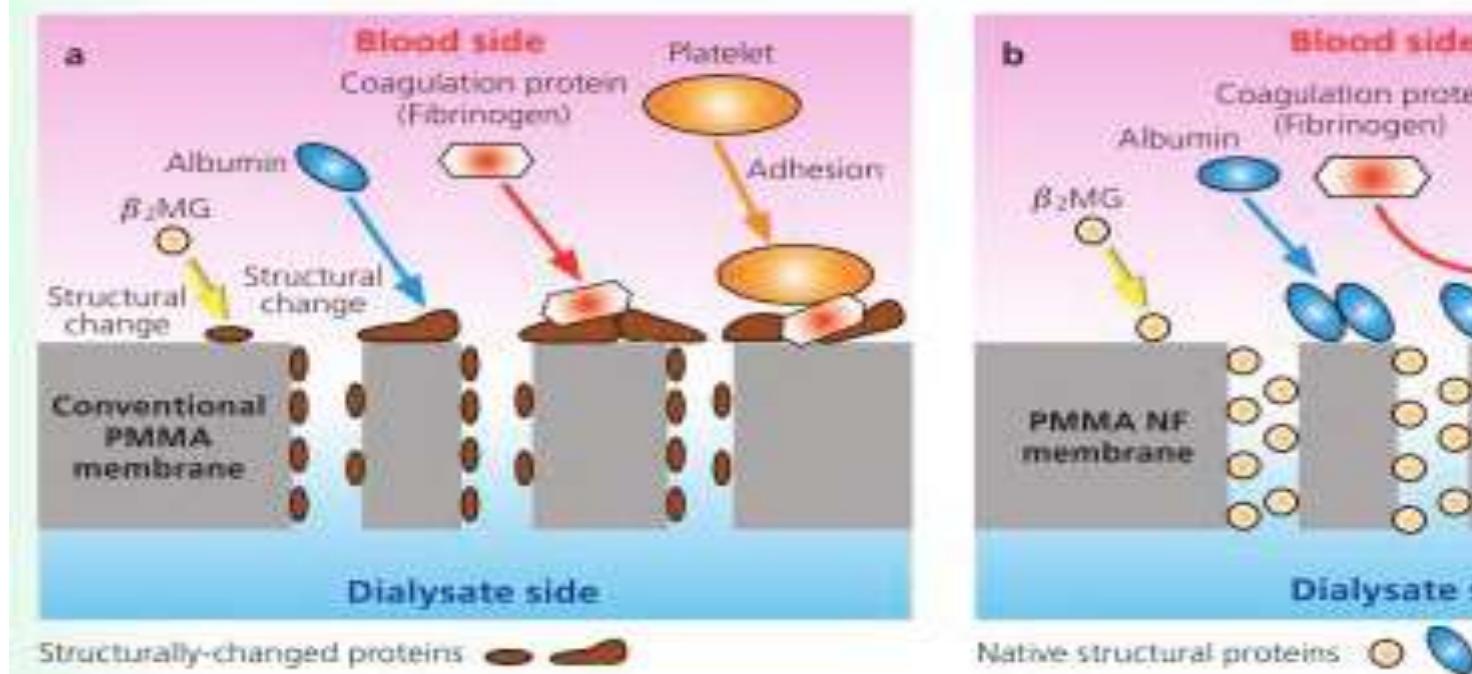


Fig.1 Schema of the protein adsorption mechanism on the PMMA membrane¹⁾

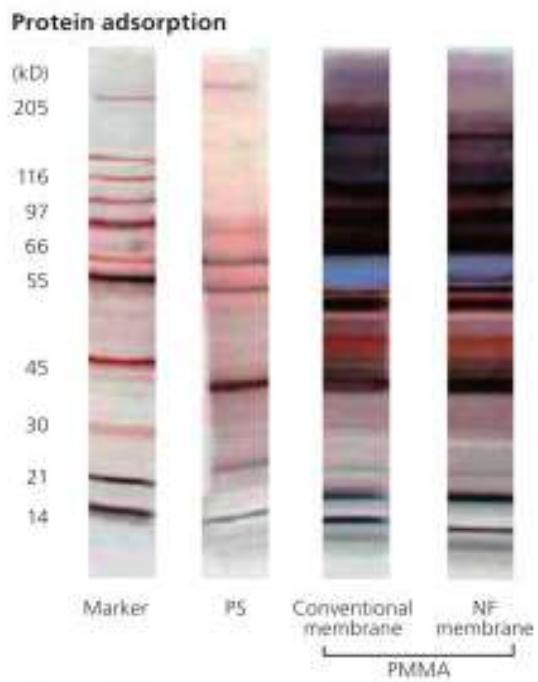


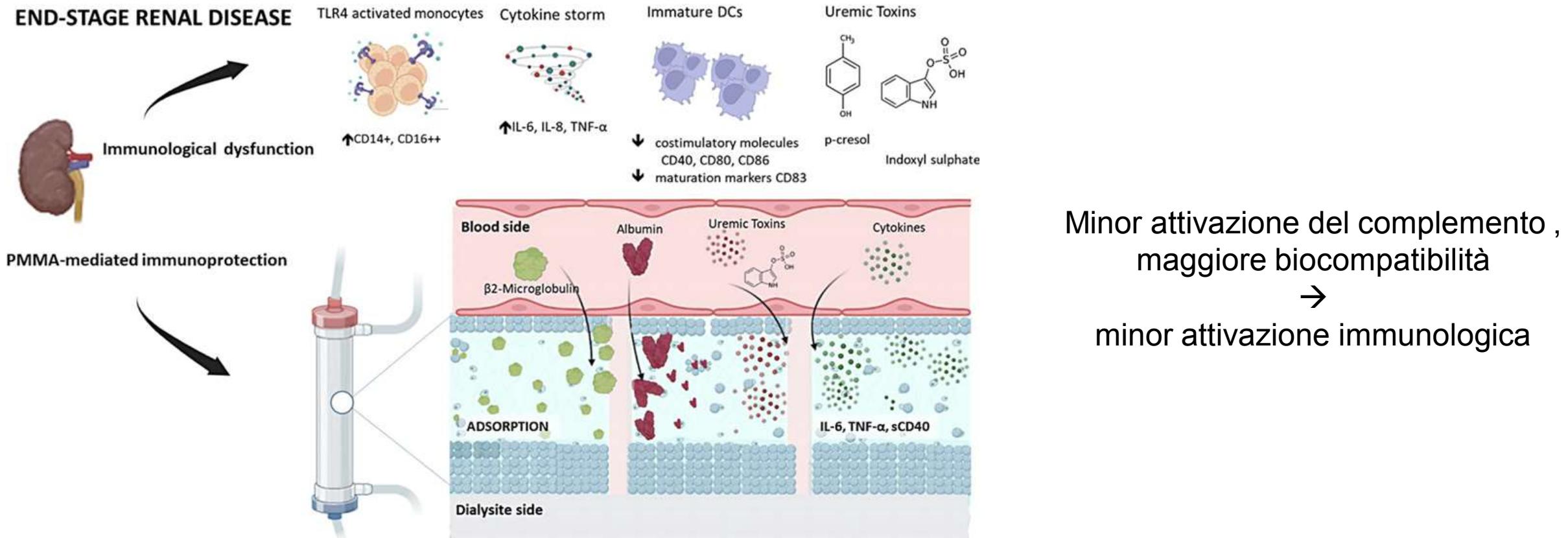
Fig.7 Electrophoretic patterns of proteins adsorbed by membrane^{2) 3)}

| Specifications | | NF-U Series | | | |
|------------------------------|--|-------------|---------|---------|---------|
| Type | | NF-1.3U | NF-1.6U | NF-1.8U | NF-2.1U |
| Fibers | Effective surface area (m ²) | 1.3 | 1.6 | 1.8 | 2.1 |
| | Effectie length (mm) | | | 195 | |
| | Inside diameter (μm) | | | 200 | |
| | Membrane thickness (μm) | | | 30 | |
| Blood volume (mL) | | 83 | 103 | 118 | 135 |
| Clearance (mL/min)* | Urea | 233 | 246 | 254 | 260 |
| | Creatinine | 200 | 217 | 225 | 231 |
| | Phosphate | 182 | 198 | 208 | 217 |
| | Vitamin B ₁₂ | 110 | 128 | 140 | 149 |
| | Inulin | 62 | 72 | 77 | 85 |
| UFR in vitro (mL/hr/mmHg) ** | | 32 | 38 | 45 | 48 |

* Clearance are typical data with aqueous solution. (Q_a: 300 mL, Q_d: 500±10 mL/min, Q_f: 10±2 mL/min, Temp.: 37±1 °C)

Enhancing Immune Protection in Hemodialysis Patients: Role of the Polymethyl Methacrylate Membrane

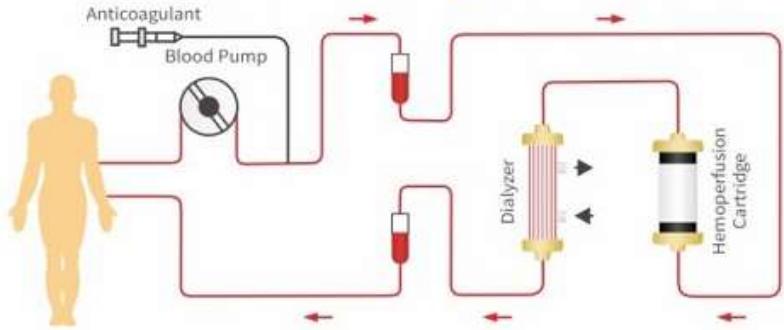
Rossana Franzina, Alessandra Stasi, Gianvito Caggiano, Elena Squicciarino, Vincenzo Losappio, Marco Fiorentino, Carlo Alfieri, Giovanni Stallone, Loreto Gesualdo, Giuseppe Castellano



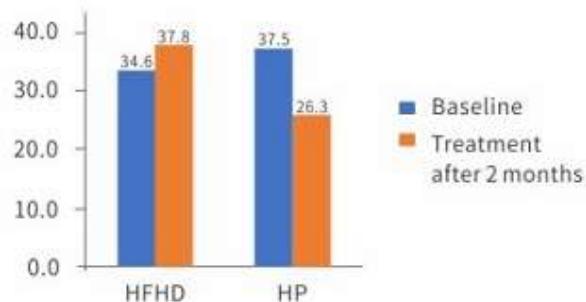
HEMOPERFUSION

N

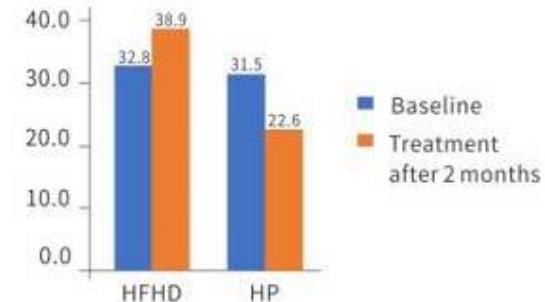
Advanced Technology · Simple Operation



Indoxyl Sulphate*

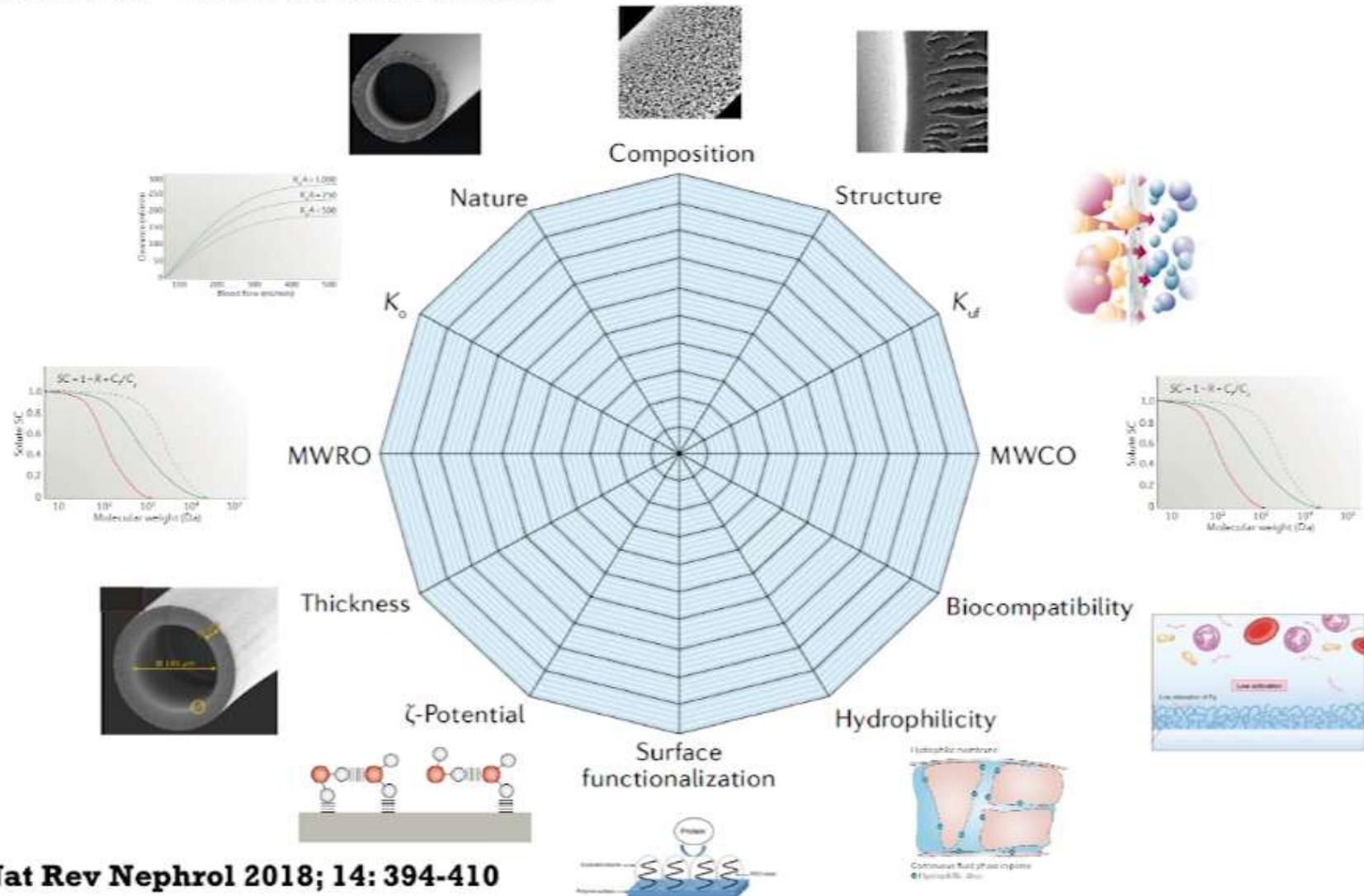


P-cresyl Sulphate*



*Unpublished data

MULTIDIMENSIONAL CLASSIFICATION OF DIALYSIS MEMBRANES



Ronco C, Clark WR. Nat Rev Nephrol 2018; 14: 394-410