

Biomarcadors en el FRA

Esteban Poch Hospital Clinic

Clasificación de la IRA (ADQI)

	GFR Criteria		Urine Output Crite	ria	
Risk	Increased SCreat x1.5 or GFR decrease > 25%		UO < 0.5mL/kg/hour for 6 hours	High	
Injury		ncreased SCreat x2 SFR decrease > 50%	UO < 0.5mL/kg/hour for 12 hours	High Sensitivity	
Failur		Increase SCreat x3 GFR decrease >75% or SCreat >4 mg/dL	UO < 0.3mL/kg/hr .g for 24hours or Anuria x 12 hrs	High	
Loss		Persistent ARF = complete loss of kidney function > 4 weeks		Specificity	
ESKD			End Stage Kidn ey Di sease (> 3 months)		

En ausencia de creatinina basal, se puede inferir tomando un MDRD de 75 ml/min/1,73m²

Critical Care 2004; 8:R204-212

Preguntas ante el FRA

- ¿Hay lesión renal (AKI)?
- ¿Qué tipo de lesión es?
- ¿Cuál es la gravedad y duración del FRA?
- ¿Existe reparación-recuperación?
- ¿Cuál es el pronóstico?

Biomarcador

'cualquier característica que es medida de forma objetiva y

evaluada como indicador de un proceso biológico normal,

un proceso patológico, o una respuesta farmacológica a una intervención terapéutica'

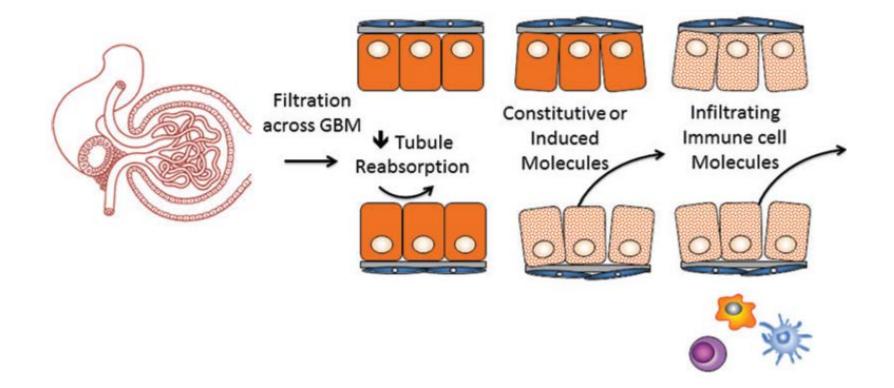
Características de un biomarcador

- Medición rápida y fácil en un margen de tiempo adecuado
- Sensibilidad alta para facilitar su detección temprana, con un rango amplio de valores para poder estratificar el riesgo (contexto clínico)
- Especificidad alta (contexto clínico)
- Accesible a partir de muestras nada (orina) o mínimamente invasivas (suero)
- Coste efectivo
- Generalizable

Características de un biomarcador de IRA

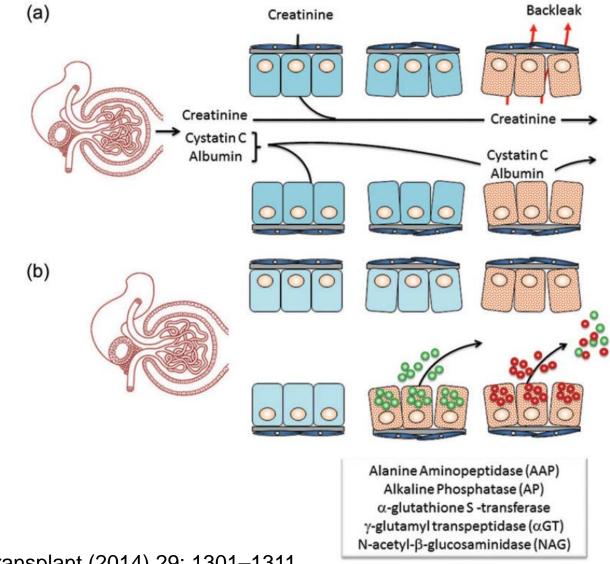
- Localización de la lesión (túbulo, glomérulo)
- Dar idea del tiempo (agudo, crónico, agudo sobre crónico)
- Mecanismo: isquemia, toxicidad, sepsis
- Predecir evolución (pronóstico): necesidad TRS, mortalidad, recuperación
- Predecir respuesta a tratamiento (si lo hubiere)





Nephrol Dial Transplant (2014) 29: 1301–1311

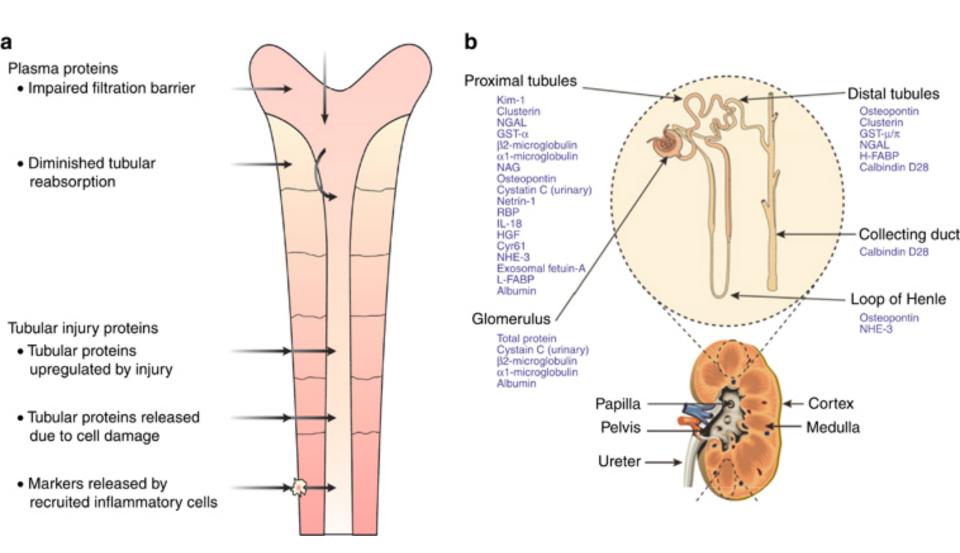




Nephrol Dial Transplant (2014) 29: 1301–1311



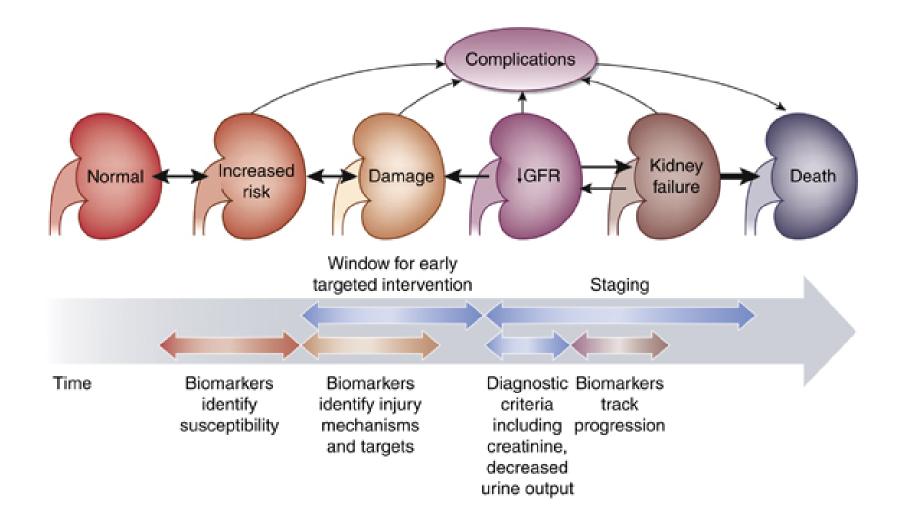
Especificidad y localización de biomarcadores urinarios



Murray PT et al, Kidney Int, 2014



Uso potencial de biomarcadores en la IRA



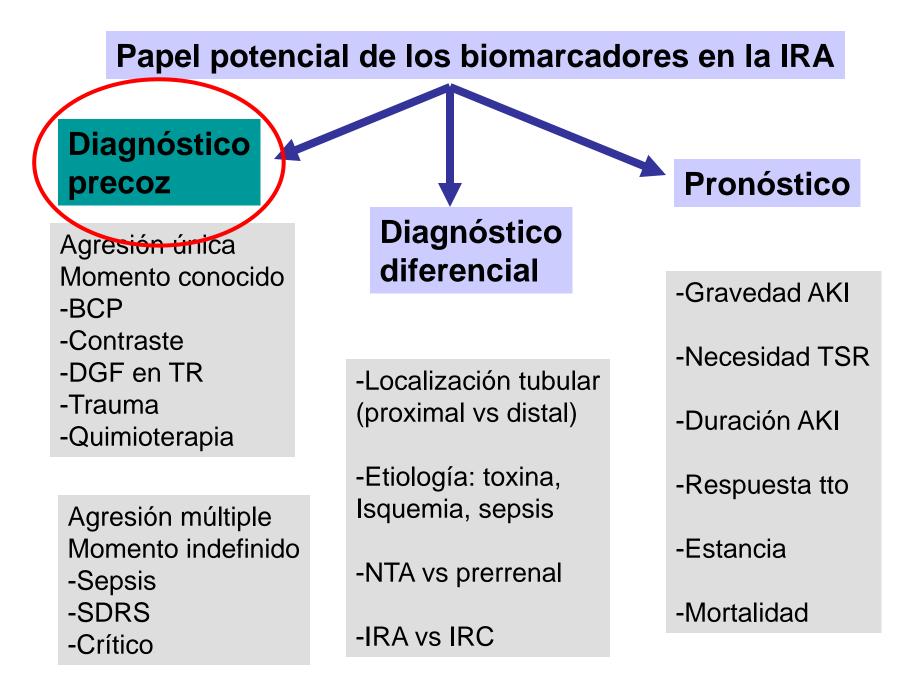
Murray PT et al, Kidney Int, 2014



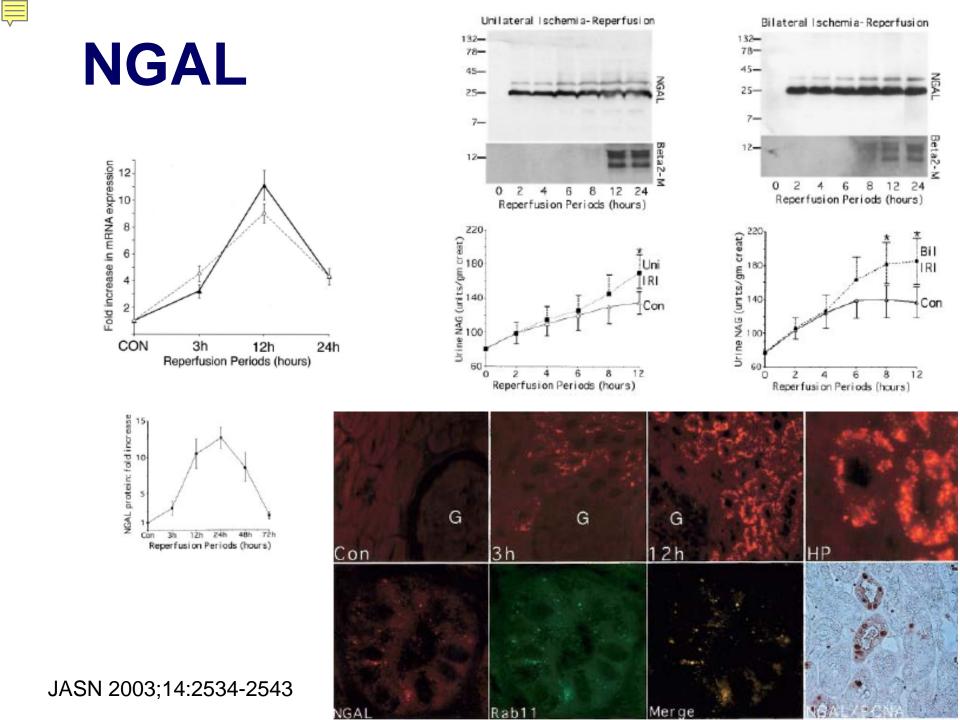
Fases en la evaluación de biomarcadores

PHASES	#1 Proof of Concept (AKI vs. no AKI)	#2 Prospective Validation (Hard Outcomes)	#3 Incremental Value to Known Predictors	#4 Does it Change Management (Clinical Utility)	#5 Improve Clinical Outcomes?	#6 Cost-Effective?
POTENTIAL STUDY DESIGNS	Cross Sectional/ Case Control/ Prospective Cohort	Nested Case Control/ Prospective Cohort	Prospective Cohort [discrimination, calibration, reclassification]	Randor	nized Clinical Trial/Pros	pective
STUDIES	NGAL (n = 35) Cystatin C (n = 22) IL-18 (n = 17) NAG (n = 15) KIM-1 (n = 14) α/π GST (n = 9) L-FABP (n = 7) Plasma IL-6 (n = 6) GGT/AlkPhos (n = 4) Netrin-1 (n = 2)	NGAL (n = 19) Cystatin C (n = 12) IL-18 (n = 9) KIM-1 (n = 4) Plasma IL-6 (n = 4) α/π GST (n = 3) NAG (n = 3) GGT/AlkPhos (n = 3) L-FABP (n = 1)	NGAL (n = 22) Cystatin C (n = 11) IL-18 (n = 10) KIM-1 (n = 6) Plasma IL-6 (n = 5) NAG (n = 5) L-FABP (n = 3) α/π GST (n = 1) Netrin-1 (n = 1)	NGAL (n = 1) GGT/AlkPhos (n = 1)		

J Am Soc Nephrol 22: 810–820, 2011

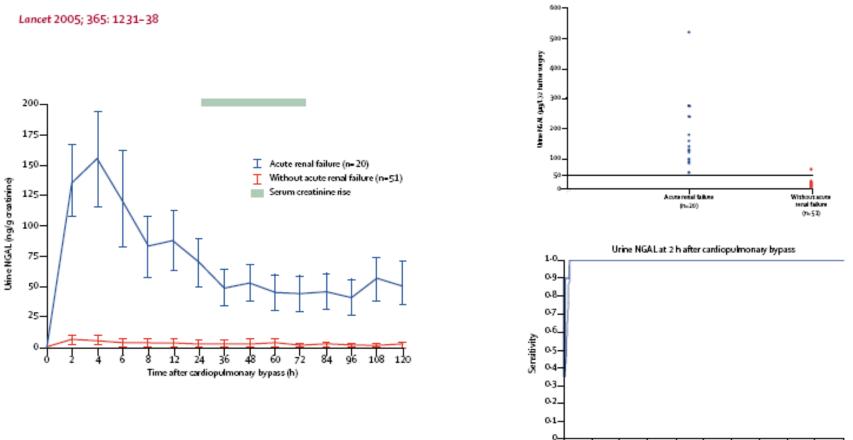


Clin Nephrol 2007; 68:269-278



Neutrophil gelatinase-associated lipocalin (NGAL) as a biomarker for acute renal injury after cardiac surgery

Jaya Mishra*, Catherine Dent*, Ridwan Tarabishi*, Mark M Mitsnefes, Qing Ma, Caitlin Kelly, Stacey M Ruff, Kamyar Zahedi, Mingyuan Shao, Judy Bean, Kiyoshi Mori, Jonathan Barasch, Prasad Devarajan



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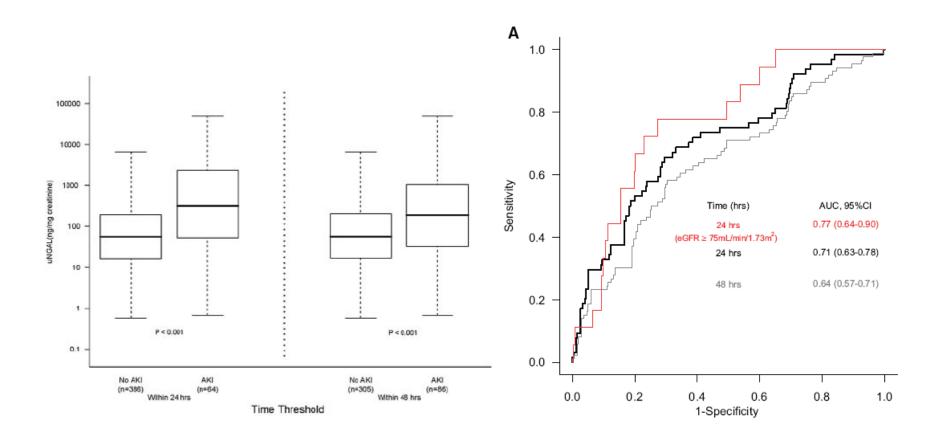
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Urine Neutrophil Gelatinase-Associated Lipocalin Moderately Predicts Acute Kidney Injury in Critically III Adults

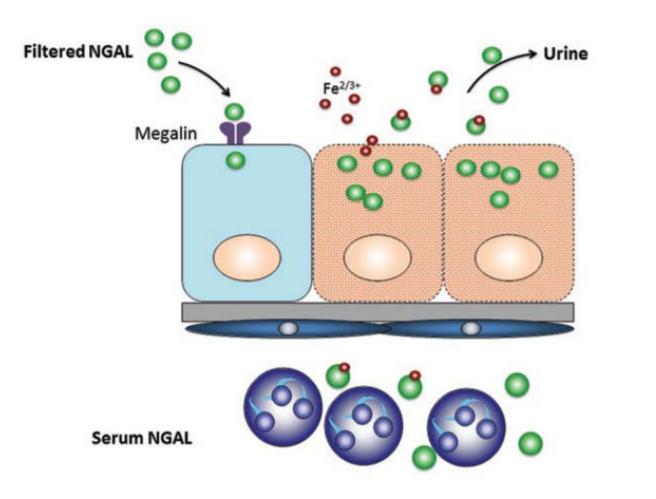
Edward D. Siew,* Lorraine B. Ware,[†] Tebeb Gebretsadik,[‡] Ayumi Shintani,[‡] Karel G. M. Moons,[§] Nancy Wickersham,[†] Frederick Bossert,[†] and T. Alp Ikizler^{*}



J Am Soc Nephrol 2009; 20:1823–1832



Función de NGAL



Monomérica

Dimérica

Nephrol Dial Transplant (2014) 29: 1301–1311

Kidney Injury Molecule-1 (KIM-1), a Putative Epithelial Cell Adhesion Molecule Containing a Novel Immunoglobulin Domain, Is Up-regulated in Renal Cells after Injury^{*}

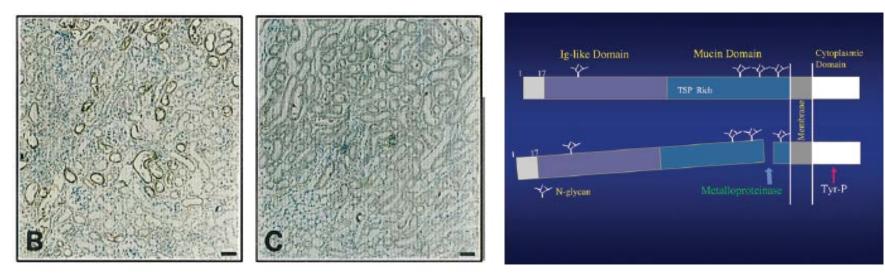
(Received for publication, September 29, 1997, and in revised form, November 18, 1997)

Takaharu Ichimura‡, Joseph V. Bonventre‡§, Véronique Bailly¶, Henry Wei¶, Catherine A. Hession¶, Richard L. Cate¶, and Michele Sanicola§¶

From the ‡Renal Unit, Medical Services, Massachusetts General Hospital East and the Department of Medicine, Harvard Medical School, Boston, Massachusetts 02129, and ¶Biogen Incorporated, Cambridge, Massachusetts 02142

Expresión de KIM-1

Estructura de KIM-1

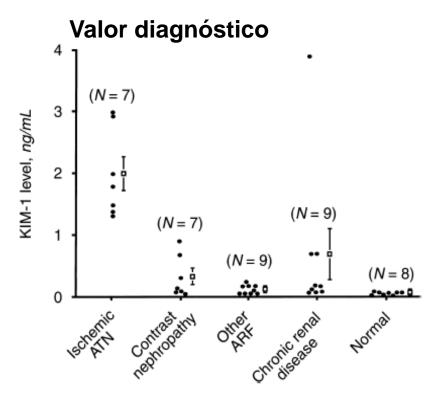


Isquemia 48h

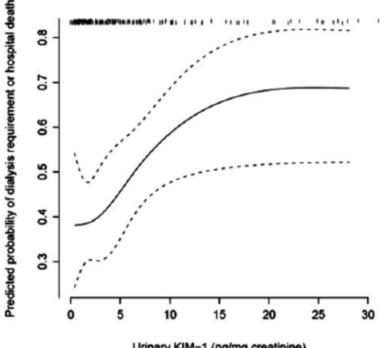




KIM-1 en la IRA







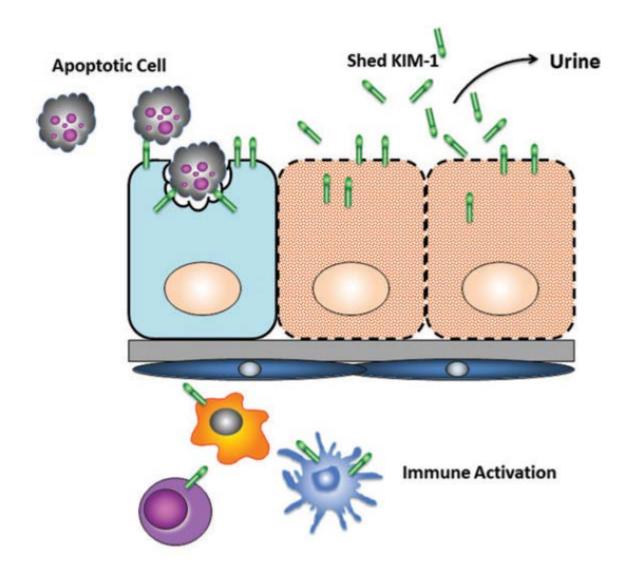
Urinary KIM-1 (ng/mg creatinine)

Kidney Int 2002;62:237-244

JASN 2007;18:904-912



Función de KIM-1



Nephrol Dial Transplant (2014) 29: 1301–1311



RESEARCH



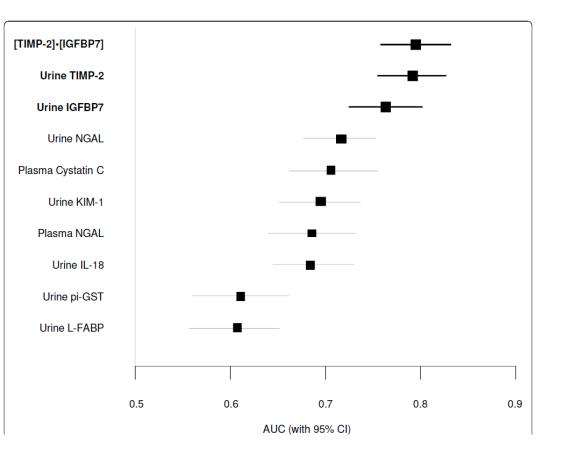
Open Access

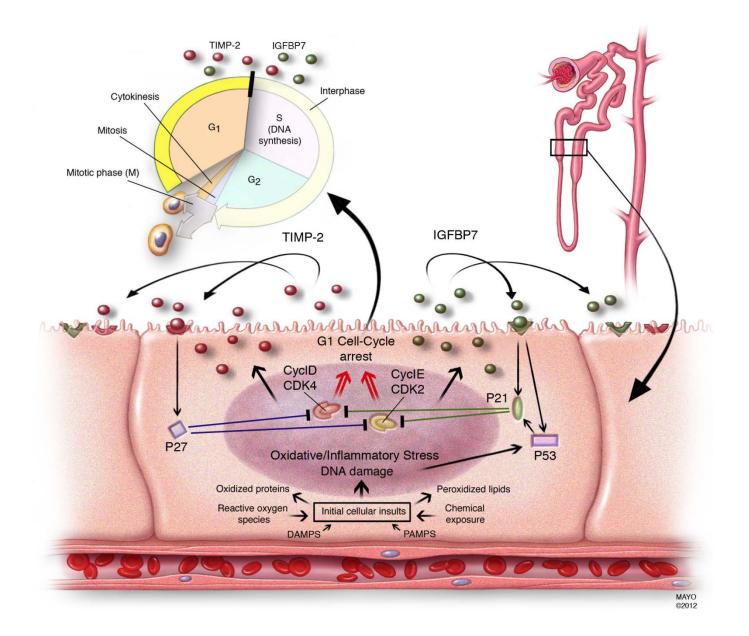
Discovery and validation of cell cycle arrest biomarkers in human acute kidney injury

Fase de prospección 500 pacientes 300 marcadores

Fase de validación 728 pacientes

Variable: AKIN 2-3 en 12

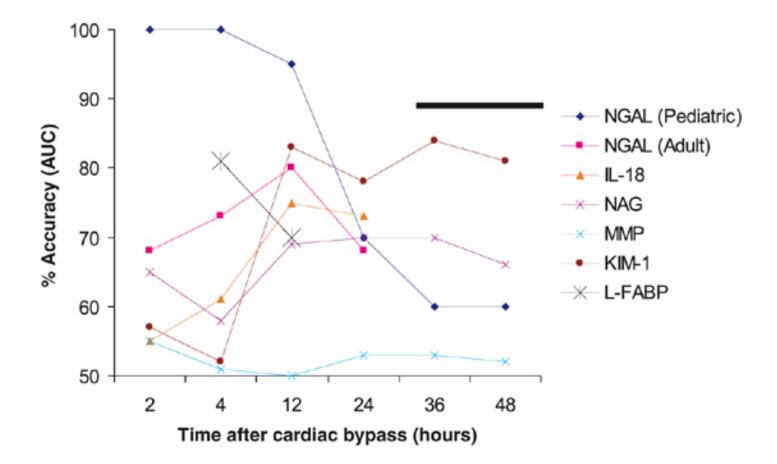




Eficacia diagnóstica de biomarcadores en orina en AKI establecida (102 AKI vs 102 no-AKI)

Biomarcador	AUC-ROC (IC 95%)	Corte	Sensibil. %	Especif. %
Cistatina C	0,85	0,12	78	83
(µ g/mg)	0,80-0,90			
IL-18 (pg/mg)	0,83	2,74	68	95
	0,77-0,88			
KIM-1 (ng/mg)	0,93	1,73	80	99
	0,88-0,96			
NAG (u/mg)	0,83	0,015	80	65
	0,77-0,88			
NGAL (ng/mg)	0,89	82,7	80	96
	0,84-0,93			
Proteína	0,91	0,46	81	87
(mg/mg) Clin Transl Sci 2008;1:2	0,87-0,95 00-2008			

Precisión de biomarcadores urinarios de IRA en post cirugía cardíaca

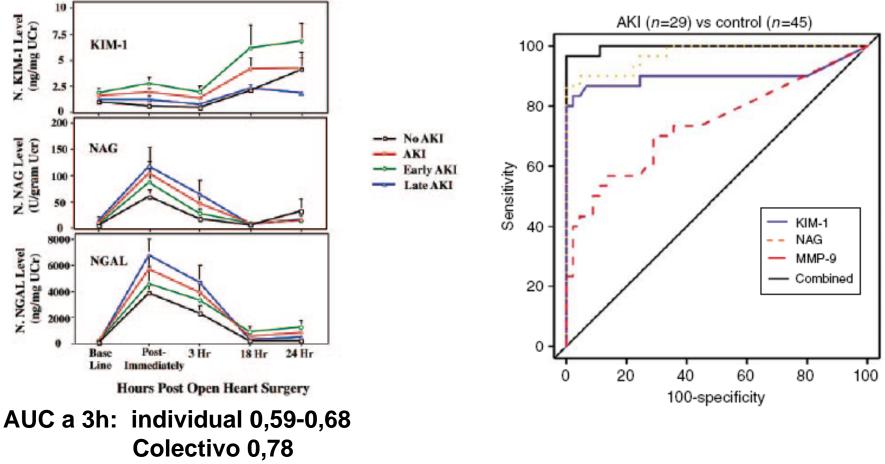




Biomarcadores combinados en orina en la IRA

CCV N=90 (36 AKI)

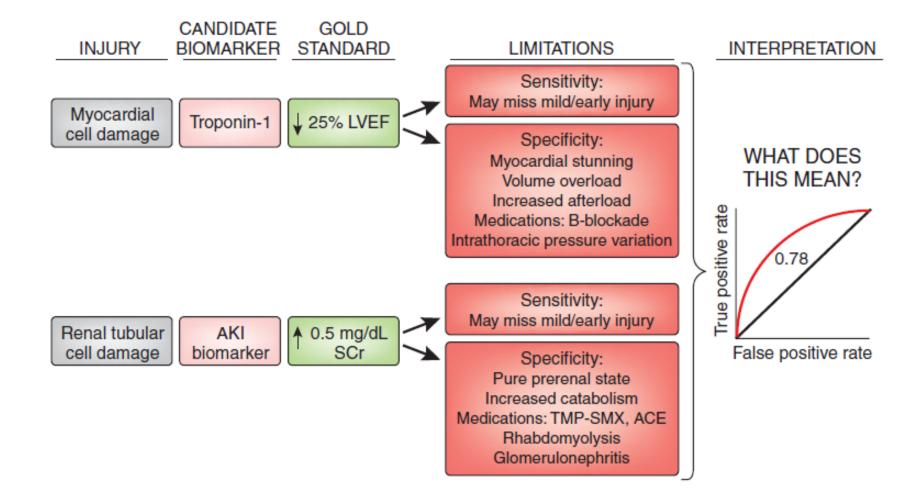
AKI varios



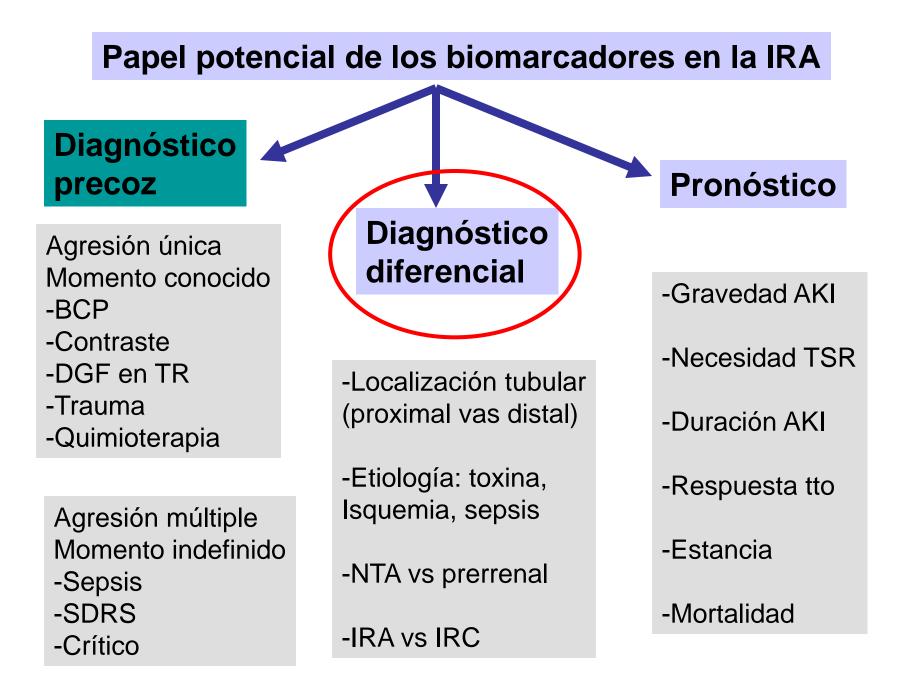
Han et al, Clin JASN 2009;4:873-882

Han et al, Kidney Int 2007





J Am Soc Nephrol 22: 810-820, 2011

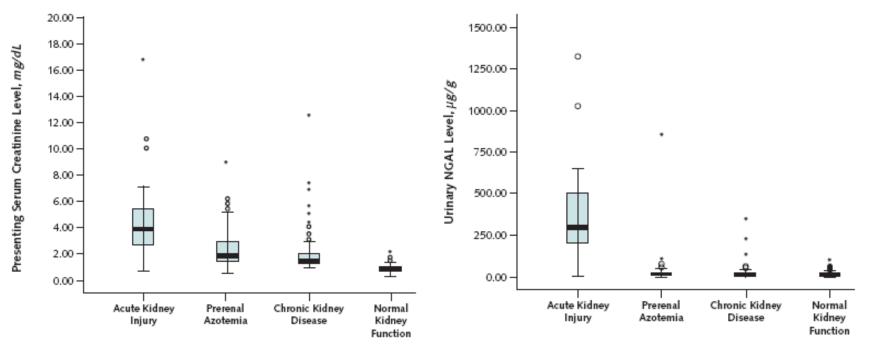


Clin Nephrol 2007; 68:269-278



Sensitivity and Specificity of a Single Emergency Department Measurement of Urinary Neutrophil Gelatinase–Associated Lipocalin for Diagnosing Acute Kidney Injury

Thomas L. Nickolas, MD, MS; Matthew J. O'Rourke, BS; Jun Yang, MD, PhD; Meghan E. Sise, BS; Pietro A. Canetta, MD; Nicholas Barasch, BS; Charles Buchen; Faris Khan, MD; Kiyoshi Mori, MD, PhD; James Giglio, MD; Prasad Devarajan, MD; and Jonathan Barasch. MD. PhD



uNGAL \geq 130 g/g creat: sensibilidad 0,90 y especificidad 0,995.

Ann Intern Med 2008;148:810-819





Urinary neutrophil gelatinase-associated lipocalin as biomarker in the differential diagnosis of impairment of kidney function in cirrhosis

Claudia Fagundes^{1,2,3,4}, Marie-Noëlle Pépin^{1,2,3,4}, Mónica Guevara^{1,2,3,4}, Rogelio Barreto^{1,2,3,4}, Gregori Casals^{2,3,5}, Elsa Solà^{1,2,3,4}, Gustavo Pereira^{1,2,3,4}, Ezequiel Rodríguez^{1,2,3,4}, Elisabet Garcia², Verónica Prado^{1,2,3,4}, Esteban Poch^{2,6}, Wladimiro Jiménez^{2,3,5}, Javier Fernández^{1,2,3,4}, Vicente Arroyo^{1,2,3,4}, Pere Ginès^{1,2,3,4,*}

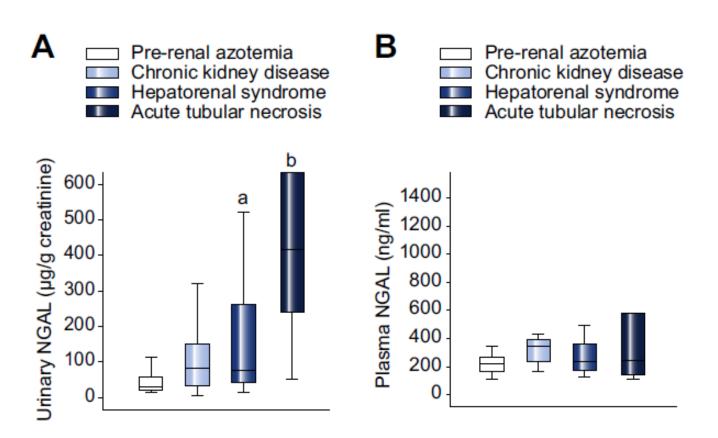


Fig. 1. Box-plot of (A) urine and (B) plasma (NGAL)

Journal of Hepatology **2012** vol. 57 | 267–273





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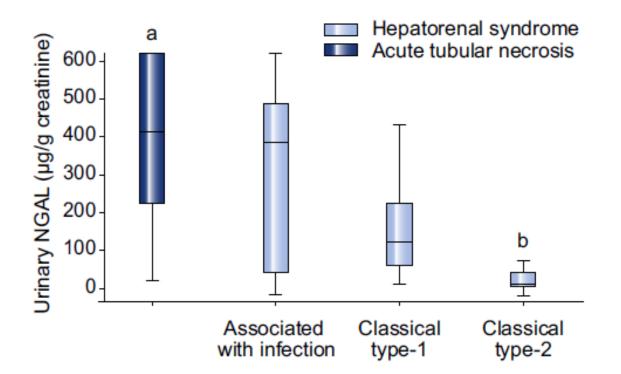
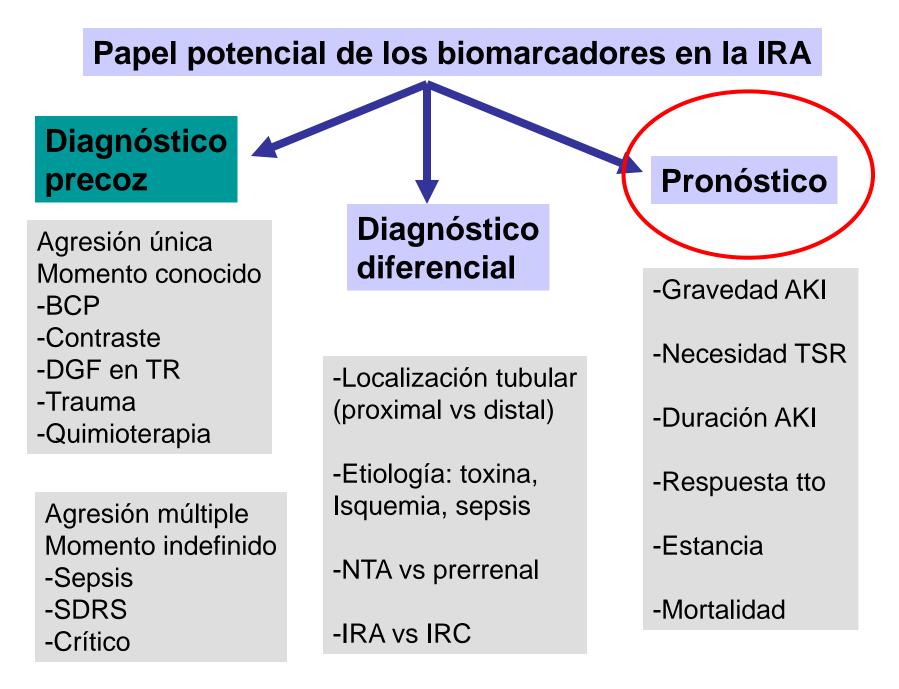
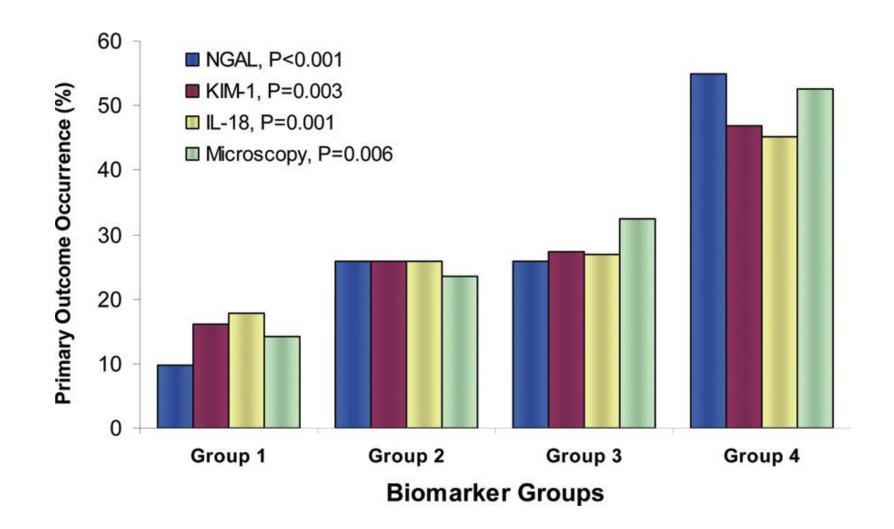


Fig. 2. Urinary neutrophil gelatinase-associated lipocalin (NGAL) levels in patients with acute tubular necrosis (ATN) and different types of hepatorenal syndrome (HRS): HRS associated with infection; classical type-1 HRS; and classical type-2 HRS. The boxes represents the 25th percentile (bottom line). Journal of Hepatology **2012** vol. 57 | 267–273



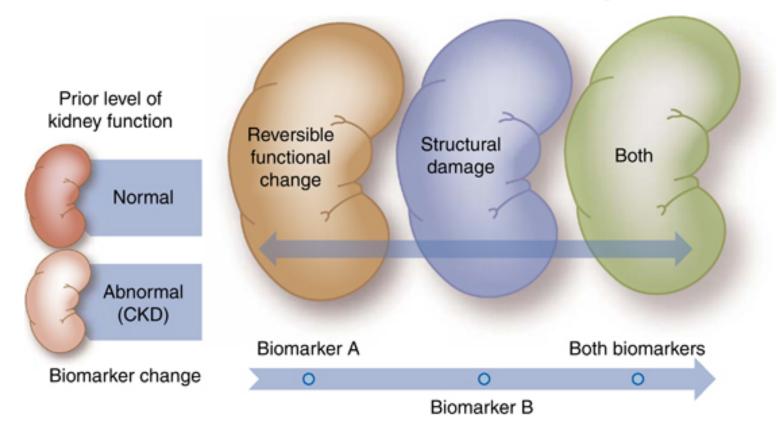
Clin Nephrol 2007; 68:269-278

Risk of Poor Outcomes with Novel and Traditional Biomarkers at Clinical AKI Diagnosis



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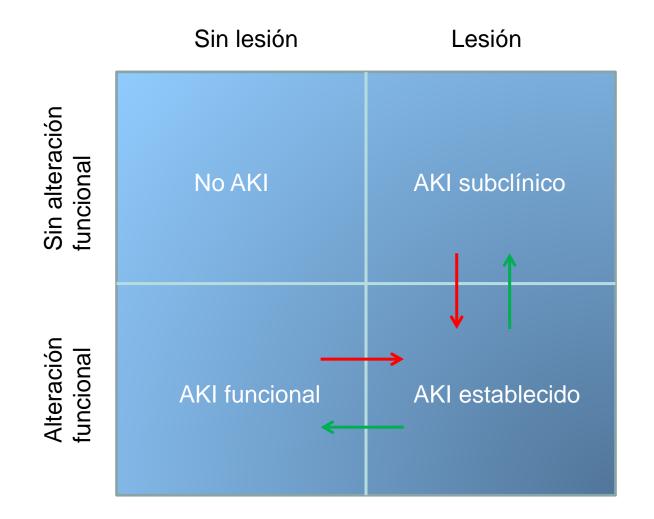




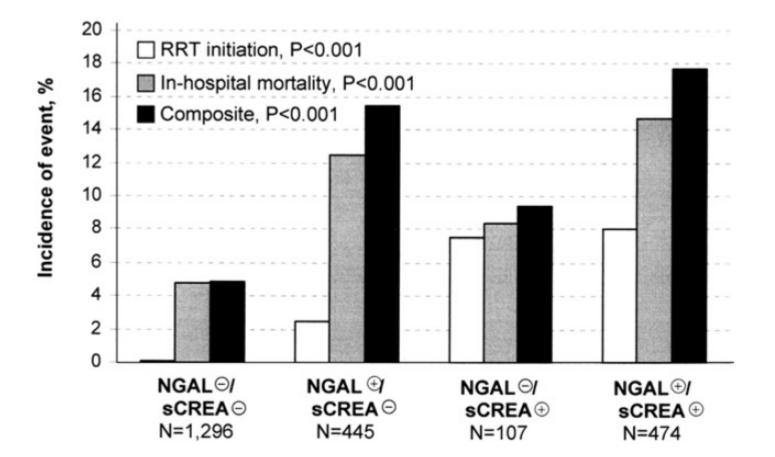
Pre-renal condition and inciting event

Murray PT et al, Kidney Int, 2014

Biomarcadores de lesión y espectro de la IRA-AKI



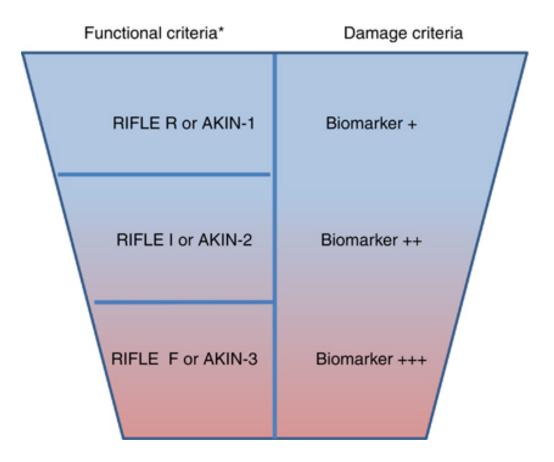
The Outcome of Neutrophil Gelatinase-Associated Lipocalin-Positive Subclinical Acute Kidney Injury



Haase M, et al. J Am Coll Cardiol 2011;57:1752–61



Propuesta de estadiaje funcional y de lesión



Murray PT et al, Kidney Int, 2014

Biomarcadores en la IRA

- La mayoría de biomarcadores nuevos rinden bien en estudios transversales de detección de IRA
 - Marcadores establecidos (diuresis, creatinina) también
- La mayoría de biomarcadores nuevos detecta AKI antes que la creatinina...
 - Pero aún no disponemos de terapia específica
- Algunos biomarcadores nuevos son pronósticos
 - Escalas clínicas (APACHE) rinden igual de bien, no cuestan, pero no se usan

Biomarcadores en la IRA

- Necesidad imperiosa en la IRA
- Poca aplicabilidad hasta ahora
 - Complejidad de la fisiopatología de la IRA
 - Heterogeneidad de pacientes
- Diagnóstico precoz de la IRA
 - No terapéutica específica
 - Medidas preventivas: ¿deberían ser universales?
- Futuro más cercano
 - Diagnóstico diferencial
 - Pronóstico: necesidad de TRS, mortalidad, duración FRA