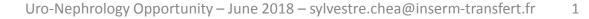




Selected opportunities in Uro-Nephrology

NGAL as a biomarker for Predicting the evolution and treatment of chronic kidney disease (BIO10870)





NGAL AS A BIOMARKER FOR PREDICTING THE EVOLUTION AND TREATMENT OF CHRONIC KIDNEY DISEASE (BIO10870)

Stage: **Product factsheet** Human POC Information: **Biomarker:** Technology: NGAL IHC, ELISA, RT-PCR Patient Stratification Prognosis ٠ Sample: Blood, Urine Scientific and Clinical Rationale: Chronic kidney disease (CKD) is characterized by progressive destruction of the renal parenchyma and the loss of functional nephrons which ultimately lead to end stage renal failure (ESRF). Understanding the pathophysiology of CKD progression is a key challenge for medical planning. ► POC: In human CKD (Autosomal dominant polycystic kidney disease (n = 84), oligomeganephronia (n = 11), IgA nephropathy (n = 12)). LCN2 was increased particularly in patients who rapidly progressed to end-stage renal failure. Selling points: Patent: EP10 306 077.8 on 2010/10/01 PCT/EP2011/067236 on 2011/10/03 Granted: EP(GB, ES, DE, IT, FR) • Scientific Publication(s):

• <u>J Clin Invest</u>, 2010 November 1, *Viau A. et al.*, doi: 10.1172/JCI42004

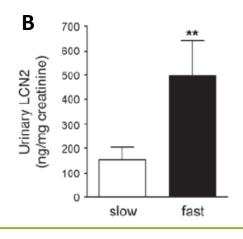
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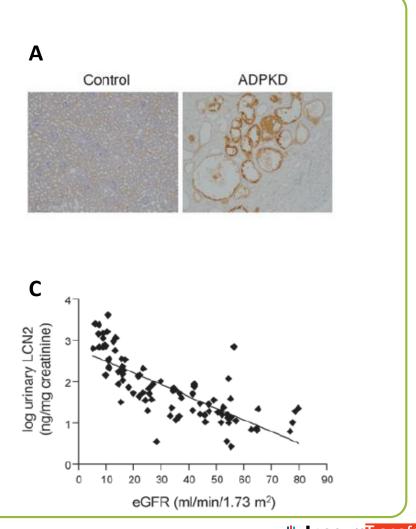
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NGAL AS A BIOMARKER FOR PREDICTING THE EVOLUTION AND TREATMENT OF CHRONIC KIDNEY DISEASE (BIO10870)

Proof of concept

- Human POC: Lcn2 is overexpressed in polycystic kidney disease in humans and correlates with CKD progression
 - (A) LCN2 staining in kidneys from controls (n = 9) and patients with ADPKD (n = 9). Original magnification, ×100.
 - (B) Urinary LCN2 excretion in patients with slow progression (eGFR decline <4.5 ml/min/1.73 m2 per year) as compared with fast progressors (eGFR decline >4.5 ml/min/1.73 m2 per year) toward ESRF.
 - (C) Urinary LCN2 excretion inversely correlates with eGFR in patients with ADPKD (r = -0.77, P < 0.0001). Data are mean ± SEM; n = 87 for ADPKD patients. Mann-Whitney U test: **P < 0.01 slow versus fast progressors.





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