

# Urine Neutrophil Gelatinase-Associated Lipocalin (uNGAL) as an Early Biomarker in Diagnosis of Acute Kidney Injury in Cirrhotic Patients

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### Background:

- The delayed diagnosis of acute kidney injury (AKI) by traditional markers in cirrhotic patients causes the undesired outcome.
- The potential of urine neutrophil gelatinase-associated lipocalin (uNGAL) which has been demonstrated as a novel early marker of AKI in various clinical setting has never been investigated in cirrhotic patients.
- This prospective cohort study was conducted to determine the accuracy of using uNGAL as a predictor for early identifying AKI in cirrhotic patients.

#### Methods:

- •The hospitalized cirrhotic patients who had risk factors for developing AKI at the time of admission were enrolled.
- The urine samples were obtained immediately after admission and subsequently next 24 hours for uNGAL testing which was operated by chemiluminescent microparticle immunoassay technique (ARCHITECT® analyzer, Abbott Diagnostics, Abbott Park, IL, USA).
- Other clinical and laboratory parameters were prospectively assessed and followed up.
- The primary endpoint was the development of AKI by AKIN criteria within 48 hours.
- The secondary end point was the 30-day mortality.

**Table 1** The sensitivity, specificity, PPV, NPV and accuracy of different cut-off values of uNGAL on admission in cirrhotic patients

Cutoff level of	sensitivit	specificity	PPV	NPV	Accuracy
uNGAL (ng/ml)	У				
46.2	95	70	54.5	97.7	79.2
65.7	84.2	81	60	94	82
150	68.4	96.6	86.7	90.3	89.6
190	47	97	81.8	84.8	84.4
259	42	100	100	84.1	85.7

## Conclusion:

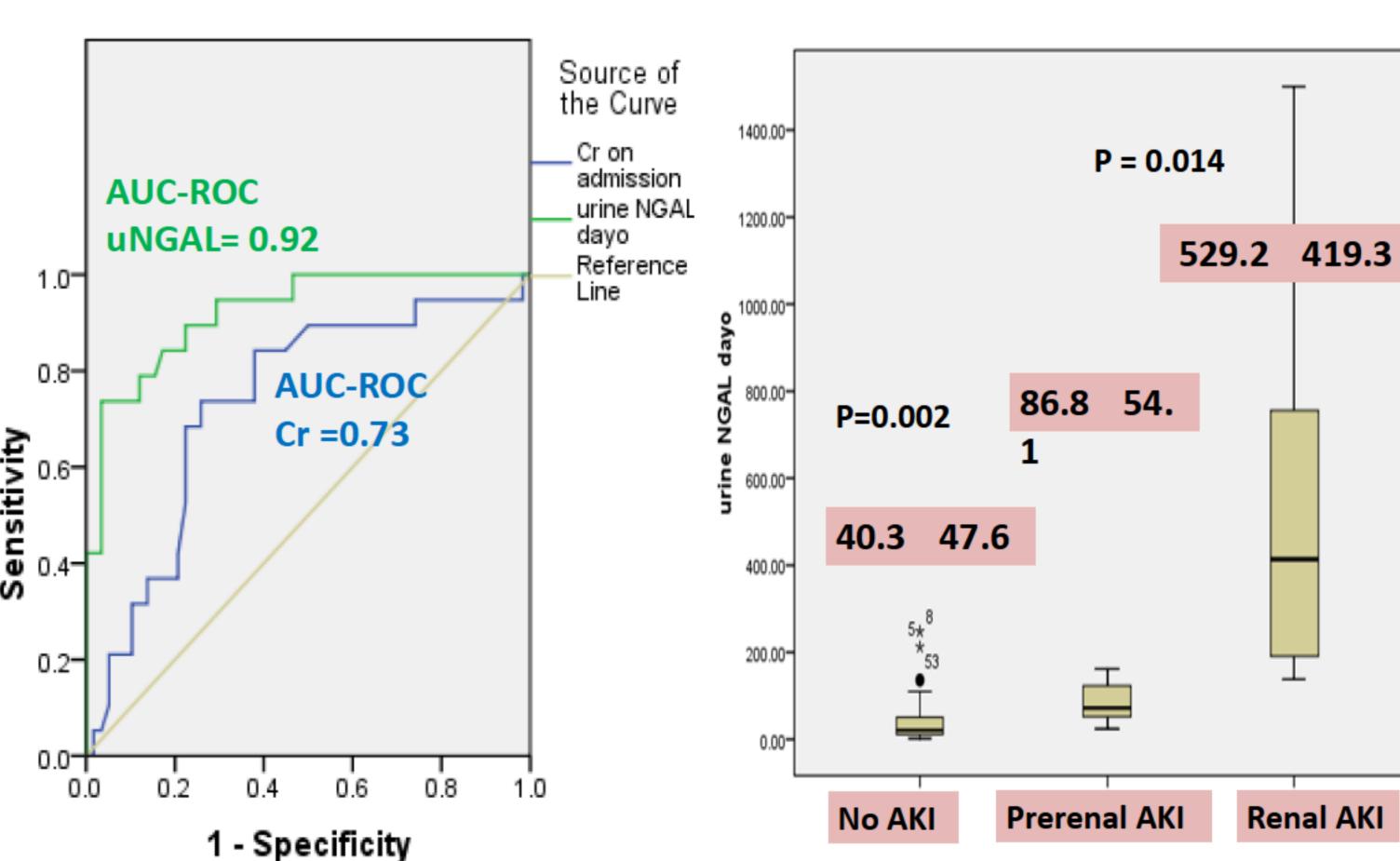
• The uNGAL is an excellent early biomarker for AKI in cirrhotic patients. The early AKI diagnosis by uNGAL might lead to the early effective treatments and improve the patient survival.

#### Results:

- 77 patients were enrolled, 24.7% of patients developed AKI within 48 h after enrollment. Child Turcotte Pugh A, B and C patients are 18.2%, 42.9% and 39% respectively.
- The mean uNGAL at enrollment was significantly higher in AKI group than non-AKI group (366.2 395.5 vs 40.3 47.6 ng/ml, p=0.002) while the serum Cr were comparable (1.06 0.32 vs 0.82 0.26 mg/dl, p =0.08).
- Using uNGAL level with cut-off 65.7 ng/mL, it could predict AKI with sensitivity and specificity of 84% and 81% as well as PPV and NPV of 60%, 94% and accuracy of 82%.
- The AUC-ROC of uNGAL for predicting AKI was 0.92 (95%CI; 0.86 to 0.98).
- The 30-day mortality was significantly higher in patients with high level of uNGAL than those with lower level of uNGAL (32.3% vs. 4.3%, p=0.001).
- Cirrhotic patients with prerenal AKI had lower uNGAL than those with renal AKI (86.8 54.1 vs 529.2 419.3ng/ml, p=0.014).

Fig. 1 The AUC-ROC of serum Cr and uNGAL in diagnosis of AKI

Fig.2 Difference values of uNGAL between cirrhotic patients with no AKI, prerenal and renal AKI



Poster

presented at: