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## Hormonal Mechanisms

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### 6.9 Plasma Aldosterone and Albumin Urinary Excretion in Hypertension

G.L. Colussi (1), C. Catena (1), A. Chiuch (1), E. Nadalini (1), A. Di Fabio (1),  
L. Sechi (1)

(1)Clinica Medica e Cattedra di Medicina Interna, Università di Udine,  
Udine, Italy

**Introduction.** Preclinical studies indicate that exposure to increased aldosterone levels might result in renal damage but the clinical evidence supporting this role of aldosterone is preliminary. Recent evidence indicates that aldosterone-related renal damage might in part reflect functional and potentially reversible abnormalities initiated by glomerular hyperfiltration.

**Methods.** In this study we examined the relationship between plasma aldosterone (ALDO) and urinary albumin excretion in patients with hypertension. We analyzed data of 222 grade 1-2 hypertensive patients (age: 49±12; 116 M, 106 F) who were consecutively referred to our unit. Secondary causes of hypertension were excluded after extensive diagnostic work-up that was performed after appropriate drug washout. Anthropometric indices and measurements of plasma lipids, uric acid, glucose, insulin, and HOMA-index were obtained after an overnight fast. Renal function was assessed by 24-h creatinine clearance and urinary albumin excretion, and the albumin-to-creatinine ratio (ACR) was calculated. Plasma active renin and ALDO were measured according to current guidelines and values were referred to the urinary Na excretion.

**Results.** When patients were subdivided according to the median value of ACR (8.6 mg/g) patients with higher values were found to have significantly greater BP, ALDO, insulin, and uric acid levels and HOMA. ACR was directly correlated with mean BP ( $P<0.001$ ), ALDO ( $P<0.001$ ), uric acid ( $P=0.005$ ), and C-peptide ( $P=0.033$ ). Multivariate regression analysis revealed that both mean BP and ALDO were independently correlated with ACR (both  $P<0.001$ ). ALDO was also correlated with plasma insulin and HOMA.

**Conclusions.** This study demonstrates that ALDO is related to urine albumin excretion independent of BP values and renal function in hypertensive patients and supports a possible role of this hormone in the development of renal damage.