

Prevalence of Microalbuminuria and its Relationship to Non-Traditional Risk Factors in Recently Diagnosed Type 2 Diabetes Mellitus: Observations from the ADOPT Study

GIANCARLO VIBERTI¹, MARTIN I FREED², RURY HOLMAN³, JOHN LACHIN⁴, MARK A HEISE² and THE ADOPT STUDY GROUP

¹London, UK; ²King of Prussia, PA; ³Oxford, UK; and ⁴Rockville, MD.

Microalbuminuria (MA) is a risk factor for cardiovascular (CV) disease and early mortality in type 2 diabetes mellitus (T2DM). The prevalence and associations of MA, defined as albumin:creatinine ratio (ACR) > 30 mg/g, were studied in 4,134 drug-naive T2DM patients (FPG ≤180 mg/dl) diagnosed within 3 years, entering a randomized double-blind comparative drug intervention trial (ADOPT). The overall prevalence of MA was 15.2% and was not affected by disease duration or age. Patients with MA (MA+) were more frequently male, significantly more obese ($P < 0.0001$), and had a significantly higher white blood cell count (WBC) ($P < 0.001$). Additionally, MA+ patients had higher blood pressure (BP) and prevalence of hypertension (HTN), as well as worse metabolic control than patients with normoalbuminuria (MA-).

| Risk Factor | MA+ | MA- | P-value |
|-----------------------|--------------|---------------|----------|
| ACR, mg/g | 87.2, 43–138 | 4.0, 3.5–10.0 | |
| HbA _{1c} , % | 7.5 ± 0.99 | 7.3 ± 0.92 | < 0.0001 |
| FPG, mg/dl | 155.6 ± 28.6 | 151.2 ± 26.1 | < 0.0001 |
| Systolic BP, mmHg | 137.0 ± 16.4 | 132.1 ± 15.2 | < 0.0001 |
| Diastolic BP, mmHg | 81.1 ± 9.3 | 79.4 ± 8.7 | < 0.0001 |
| Dx HTN*, % | 83.3 | 76.3 | < 0.0001 |

Mean ± SD, or Geometric Mean, IQR for ACR, *prior diagnosis of HTN or BP ≥ 130/85

Treatment with ACE Inhibitors and/or All Receptor Blockers was also more frequent in MA+ (21.5%) vs MA- (17.7%) patients ($P < 0.024$). LogACR significantly correlated with HbA_{1c} ($r = 0.056$, $P = 0.0004$), FPG ($r = 0.054$, $P = 0.0006$), SBP ($r = 0.110$, $P < 0.0001$), DBP ($r = 0.085$, $P < 0.0001$) and WBC ($r = 0.086$, $P < 0.0001$). MA was significantly related to traditional and non-traditional CV risk factors and its prevalence, in our cohort, was high and similar to the 12.3% reported in newly diagnosed T2DM by the UKPDS. This emphasizes the need for more aggressive, comprehensive treatment of MA, hyperglycemia, hypertension and other associated CV risks in T2DM.