

HIGHER ASPARTATE AMINOTRANSFERASE LEVELS ARE ASSOCIATED WITH HIGHER ALL-CAUSE MORTALITY IN MAINTENANCE HEMODIALYSIS PATIENTS

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BACKGROUND

In Maintenance Hemodialysis (MHD) patients, liver disease is a comorbidity and may be associated to poor survival. Some studies have observed positive association between elevated liver enzymes and mortality risk in MHD patients but to date, the relationship between Aspartate Aminotransferase (AST) and all-cause mortality risk in MHD patients has not been well studied. We hypothesized that higher levels of AST would be associated with increased all-cause mortality in MHD patients

OBJECTIVE

To evaluate association between AST and all-cause mortality and to determine variables associated with higher AST level in a contemporary cohort of US dialysis patients.

RESULTS

Patients were 61±15 years old and included 45% women, 32% blacks, and 58% diabetics. Using AST <20 U/L as reference, there was a positive association between increasing AST levels and all-cause mortality across all levels of multivariable adjustment. In fully adjusted models, highest risk of death was observed in patients with AST levels ≥80 U/L (HR: 1.70, 95%CI: 1.56-1.85)

We used baseline AST levels in fractional polynomial models in fully adjusted models, We found a significant positive association between increasing AST levels above 20U/L and all-cause mortality.

METHODS

In this study we analyzed a database of 114,267 DaVita MHD patients followed up to 8 years (2001-2009) to examine the association of AST with all-cause mortality. We used 5 categories of baseline AST (<20, 20-<40, 40-<60, 60-<80, ≥80 U/L) in Cox proportional hazard models with 3 levels of adjustment: unadjusted, case-mix, and case-mix covariates plus markers of the malnutrition and inflammation complex (MICS).

Figure 1. All-cause mortality hazard ratio

Unadjusted, Case-Mix adjusted, Case mix+MICS adjusted

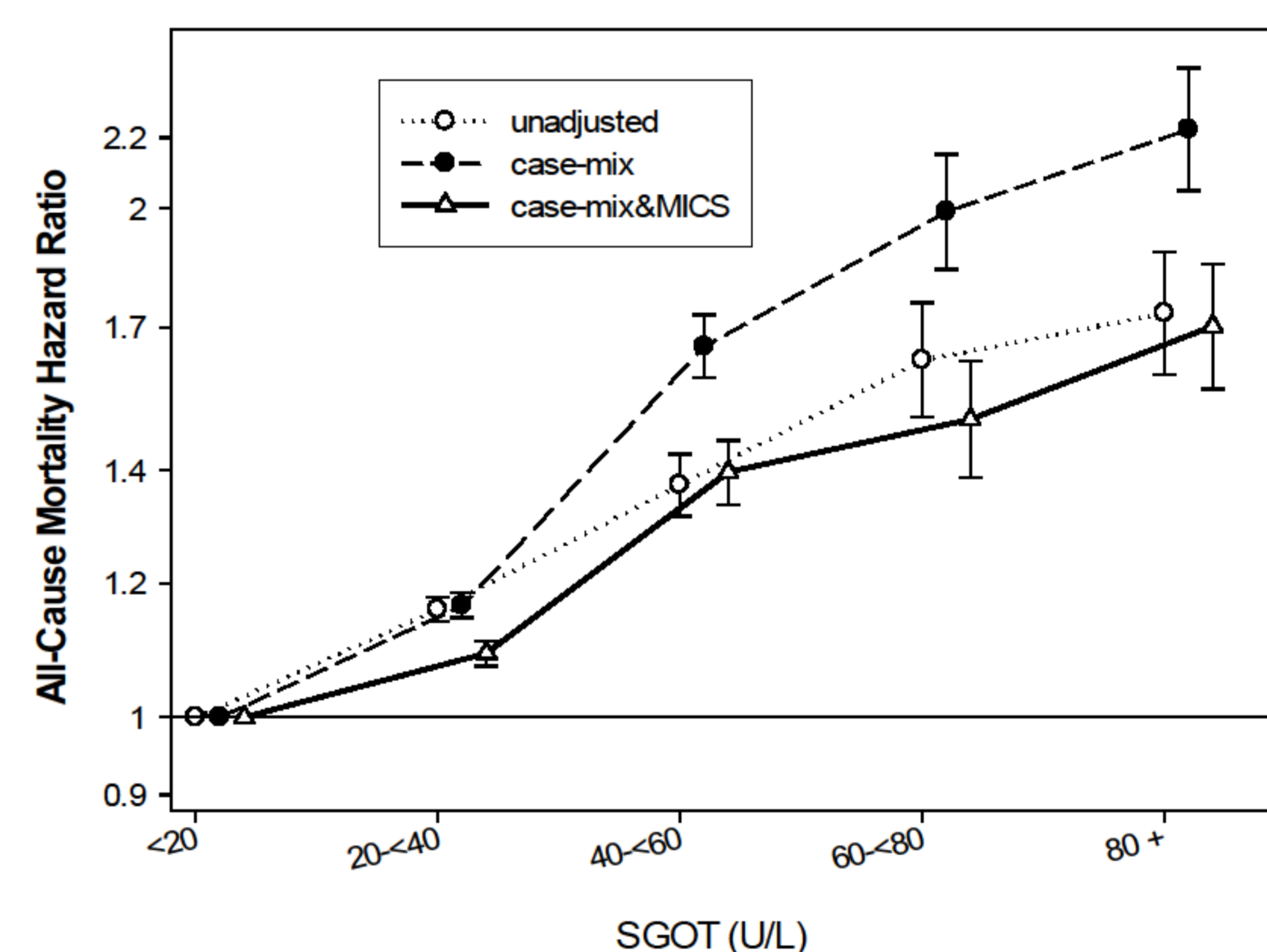


Figure 2. Mortality spline

Case mix+MICS adjusted

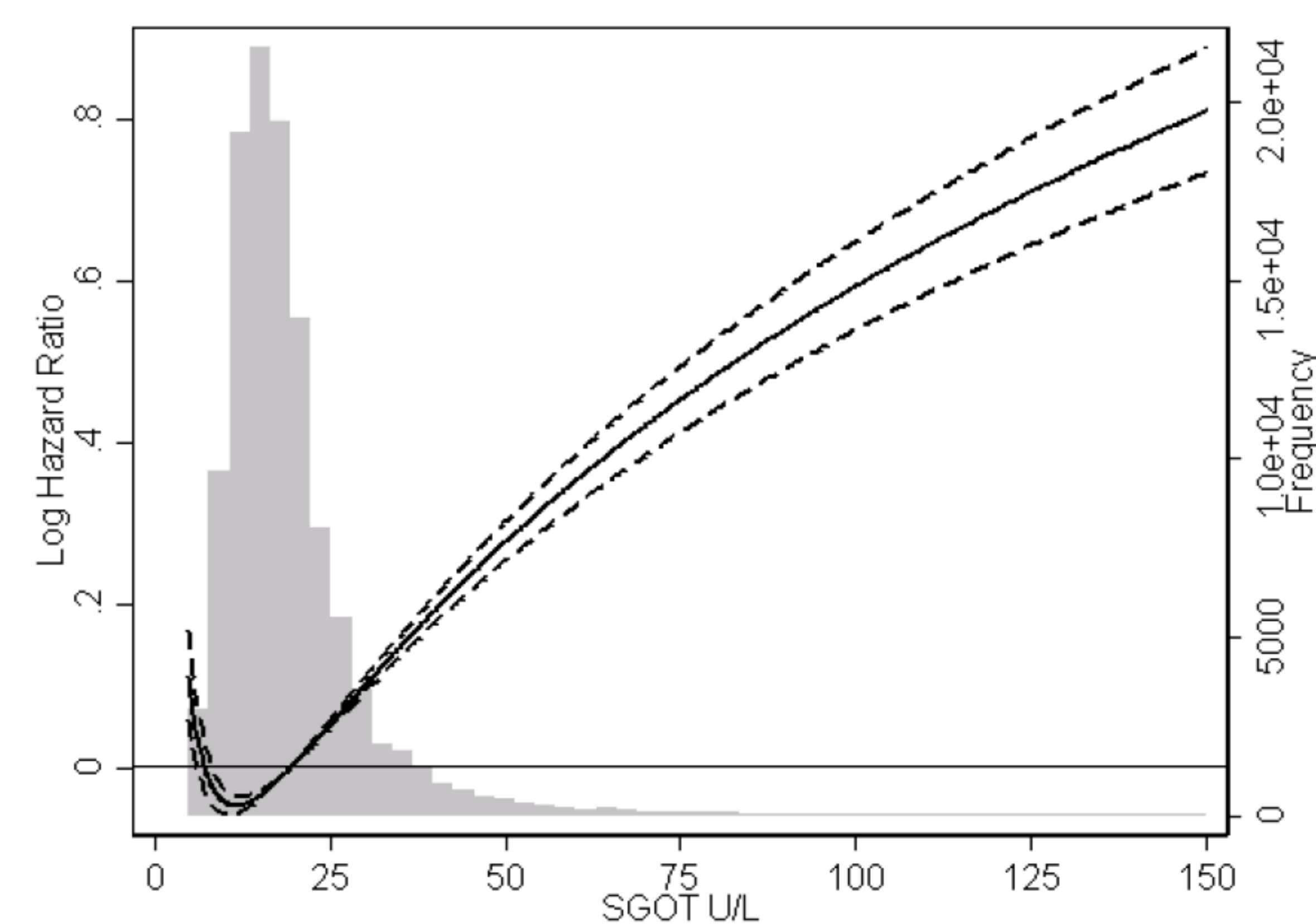


Table 1. Demographic Characteristics of 114,267 MHD patients and in 5 AST groups

Variables	All	AST (U/L)					P valueREG
		<20	20 to <40	40 to <60	60 to <80	80+	
No of patients	114,267	74,949	34,301	3,367	896	754	
Age (years)	61±15	61±15	62±15	57±14	57±14	56±15	<.0001
Gender (% women)	45	45	46	45	43	42	0.3653
Race (Black%)	32	32	32	42	42	43	<.0001
Tobacco Use(%)	4	5	5	6	7.7	5.4	0.0047
Drug Dependence(%)	1	1	2	3	3.1	3.8	<.0001
Alcohol Dependence (%)	1	1	2	3	3.8	4.2	<.0001
Diabetes mellitus (%)	57	58	58	57	54	56	0.6564
History of Hypertension(%)	79	80	79	75	74	74	<.0001
AIDS(%)	1	1	1	3	3.2	4	<.0001
Cancer(%)	4	4	5	4	4	4	0.6805
albumin (g/dL)	3.67±0.46	3.72±0.44	3.6±0.49	3.42 ±0.55	3.3±0.58	3.29±0.57	<.0001
creatinine (mg/dL)	8.02±3.32	8.38±3.32	7.34±3.22	7.48±3.37	7.37±3.20	7.45±3.28	<.0001
Body Mass Index (kg/m ²)	26.69±6.93	27.1±7.0	26.03±6.63	25.25±6.24	24.7±5.7	24.86±6.48	<.0001

CONCLUSION

In MHD patients, increasing levels of serum AST above 20 U/L are associated with a linear increased risk of all-cause mortality even after adjustment for MICS markers. Further studies are needed to confirm findings and determine mechanistic pathways of the AST—mortality association

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