



INCREASED LEVELS OF SOLUBLE FMS-LIKE TYROSINE KINASE 1 (SFLT-1) ARE ASSOCIATED WITH WORSE OUTCOMES IN OUTPATIENTS WITH HEART FAILURE

ACC Moderated Poster Contributions
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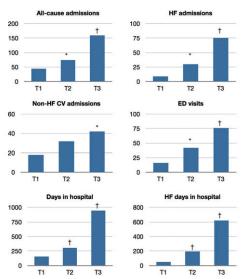
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Background: Soluble fms-like tyrosine kinase-1 (sFLT-1) antagonizes vascular endothelial growth factor and placental growth factor (PIGF) and regulates apoptosis in vascular smooth muscle cells. sFLT-1 levels predict clinical events in systolic heart failure (HF); however, the association of sFLT-1 with other outcomes in HF has not been reported.

Methods: We examined the association of baseline levels of sFLT-1, PIGF, and sFLT-1 to PIGF ratio with (1) clinical events (death, transplantation, ventricular assist device implantation) and (2) admissions and emergency department [ED] visits in 173 stable HF outpatients (age, 57±12 yrs; 63% men; 58% white; 38% black; ejection fraction [EF] 29±15%) enrolled in a prospective cohort study.

Results: Over 32±8 months (total: 465 person-years), there were 27 (15.6%) clinical events (22 deaths, 4 transplants, 1 ventricular assist device), 413 all-cause admissions (167 [40.4%] for HF), and 199 ED visits. Baseline sFLT-1, PIGF, and sFLT-1/PIGF were 339±83 pg/ml, 19.2±5.1 pg/ml, and 18.9±7.4, respectively. Compared to the lower sFLT-1 tertile, patients in the upper tertile had (1) increased risk for clinical events (HR 4.5; 95% CI 1.2-17.3; P=0.029) and (2) higher healthcare resource utilization rates (Figure) in models adjusted for age, gender, race, systolic blood pressure, creatinine, NYHA class and EF. PIGF and sFLT-1/PIGF were not predictive of outcomes.

Conclusion: Increased sFLT-1 but not PIGF levels are associated with worse outcomes in HF outpatients.



All rates per 100 patient-years; T1-T3: sFLT-1 level tertiles; *P<0.01; †P<0.001 vs T1