

Table 2. Major cardiac and pulmonary events and all-cause mortality according to beta-blocker use among MCBS cohort members with CVD and COPD (N=1,062) ^a

| Outcome (n=no. of events) | Unadjusted HR (95% CI) | PS-adjusted ^{c,d} HR (95% CI) | Covariate- adjusted ^b HR (95% CI) |
|--------------------------------------|-----------------------------------|---|---|
| <i>Cardiac event</i> (n=179) | 1.35 (1.00, 1.81) | 1.18 (0.85, 1.62) | 1.12 (0.83, 1.52) |
| <i>Pulmonary event</i> (n=389) | 1.05 (0.86, 1.29) | 0.91 (0.73, 1.12) ^e | 0.86 (0.71, 1.06) ^e |
| <i>Death</i> (n=255) | 0.97 (0.76, 1.23) | 0.87 (0.67, 1.13) | 0.82 (0.63, 1.07) ^f |

Abbreviations: CI, confidence interval; COPD, chronic obstructive pulmonary disease; CVD, cardiovascular disease; HR, hazard ratio; MCBS, Medicare Current Beneficiary Survey; PS, propensity score.

^a The HRs for β -Blocker use were created with Cox proportional hazards analysis. The referent groups were β -blocker nonusers.

^b All covariate models are controlled by entry year. Cardiac model additionally includes: heart failure, ACE/ARB use, and Elixhauser Comorbidities. Pulmonary model includes: total non β -Blocker medications, heart failure and Elixhauser Comorbidities. Mortality model includes: Elixhauser Comorbidities, total number of non β -Blocker medications, prior MI, cognitive impairment, limited social activities, ESRD, heart failure, and ACE/ARB use.

^c Variables used to generate the propensity score are marked in Table 1.

^d All PS models are controlled by PS and entry year. Pulmonary PS model additionally includes: physical function and Elixhauser Comorbidities. Mortality model includes: cognitive impairment.

^e One outlier removed; outlier did not have event.

^f Two outliers removed; outliers did not have event.