Fibrinogen, chronic obstructive pulmonary disease (COPD) and outcomes in two United States cohorts

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Background: Fibrinogen is a marker of systemic inflammation and may be important in the pathogenesis and progression of chronic obstructive pulmonary disease (COPD).

Methods: We used baseline data from Atherosclerosis Risk in Communities and Cardiovascular Health Studies to determine the relation between fibrinogen levels and COPD and to examine how fibrinogen levels at baseline affected outcomes of death, development of COPD, lung function decline, and COPD-hospitalizations.

Results: Our study sample included 20,192 subjects, of whom 2995 died during the follow-up period. The mean fibrinogen level was 307.6 mg/dL and 10% of the sample had levels >393.0 mg/dL. Subjects with Stage 3 or 4 COPD were more likely to have a fibrinogen level >393.0 mg/dL (odds ratio 2.28, 95% confidence interval [CI]: 1.79–2.95). In the longitudinal adjusted models, fibrinogen levels >393 mg/dL predicted mortality (hazards ratio 1.54, 95% CI: 1.39–1.70), COPD-related hospitalization (hazards ratio 1.45, 95% CI: 1.27–1.67), and incident Stage 2 COPD (odds ratio 1.36, 95% CI: 1.07–1.74). Similar findings were seen with continuous fibrinogen levels.

Conclusion: In the Atherosclerosis Risk in Communities/Cardiovascular Health Studies cohort data, higher fibrinogen levels are predictors of mortality, COPD-related hospitalizations, and incident Stage 2 COPD.

Keywords: COPD, fibrinogen, epidemiology, mortality, hospitalization

Background

Chronic obstructive pulmonary disease (COPD) is a heterogeneous disease with varied clinical presentations. 1 COPD became the third leading cause of death in the United States in 2008. 2 Approximately one in 20 deaths in the United States had COPD as the underlying cause. 2 Understanding the natural history of COPD has been important in the field of pulmonary medicine, dating back to the work of Burrows et al. 3 and Fletcher et al. 4 Over subsequent years researchers have championed different hypotheses about COPD development, including the British hypothesis stating that the presence of cough and sputum was the key factor 5 and the Dutch hypothesis stating that the presence of increased Airways responsiveness was the major factor. 6

Fibrinogen is an inflammatory marker that is increased in COPD, as well as many other inflammation-associated diseases. 7 The Coronary Artery Risk Development in Young Adults CARDIA study showed fibrinogen as a marker for chronic low-grade inflammation and is associated with modest deterioration of lung function in healthy young adults. 8 According to Dahl and colleagues, increased levels of fibrinogen were associated with reduced lung function and increased risk of COPD,
and these associations were independent of smoking status. Groenevogen and colleagues demonstrated that besides lung function impairment systemic inflammation manifested by elevated fibrinogen levels was an independent risk factor for exacerbations of COPD. Hyperfibrinogenemia is linked with asthma and smoking. There is little information on the relationship between plasma fibrinogen levels and lung function decline; although recent work has suggested that systemic processes may be important in these processes.

We hypothesized that fibrinogen can be used as a tool for stratifying COPD patients in clinical trials by identifying populations at higher risk for poor outcomes such as development of COPD, rapidly declining lung function, COPD hospitalizations, and death. This paper examines the descriptors of normal and elevated fibrinogen levels at baseline in a cohort of US adults and determines the relationship between elevated fibrinogen levels and incident respiratory outcomes including development of COPD, rapidly declining lung function, COPD hospitalizations, and death controlling other risk factors, including cardiovascular disease (CVD).

Methods
Study population
The study population originated from the combined cohorts of the Atherosclerosis Risk in Communities (ARIC) and Cardiovascular Health Study (CHS). Both the ARIC and CHS were population-based National Institute of Health cohorts initiated in the late 1980s.

The ARIC study was initiated in 1987 as a longitudinal, population-based study of the etiology and clinical sequelae of atherosclerosis in 15,792 adults. Study protocols were approved for protection of human subjects. Participants were selected from the entire population by probability sampling from four US communities: Forsyth County, NC; Minneapolis, MN; Washington County, MD; and Jackson, MS (where only African Americans were sampled). Specific details of the ARIC study are published elsewhere. Our analysis was limited to ARIC participants aged 45–64 years old at baseline, who provided information on respiratory symptoms and diagnoses, medical history, and smoking status, and who underwent a clinical exam and spirometric testing at baseline and 4 years. Details of the CHS are published elsewhere.

Definition of variables
Demographic data used in this analysis were age (in 5 year categories for ARIC and 4 year categories for CHS), sex, race (black, white, other), education (less than 12 years, 12 years, and more than 12 years), The study participants were categorized into normal and elevated fibrinogen values with a cut off value of greater than 393 mg/dL (corresponding to the top decile) and six lung function categories based on modified Global Initiative for Chronic Obstructive Lung Disease (GOLD) criteria; classification is based on “pre-bronchodilator” response. GOLD Stage 3 or 4 (forced expiratory volume in 1 second (FEV1)/forced vital capacity (FVC) <0.70 and FEV1 <50% predicted), GOLD Stage 2 (FEV1/FVC <0.70 and FEV1 >50 to <80% predicted), GOLD Stage 1 (FEV1/FVC <0.70 and FEV1 >80%), restricted (FEV1/FVC >0.70 and FVC <80% predicted), GOLD Stage 0 (presence of respiratory symptoms in the absence of any lung function abnormality), and no lung disease.

We included in the analyses potential confounders of CVD at baseline (subjects reporting a diagnosis of a previous myocardial infarction, stroke, heart failure, angina or transient ischemic attack, and excluding hypertension). Smoking status included current, former, and never smokers. Respondents with a positive response to “Have you ever smoked cigarettes?” and “Do you now smoke cigarettes?” were classified as “ever smokers” and “current smokers,” respectively. Pipe or cigar smokers were also considered as “smokers.” Subjects were classified as having diabetes if they reported either a diagnosis of diabetes at baseline or had impaired fasting or post-glucose load levels (140 mg/dL) upon examination. Body mass index was categorized as “Underweight” (<20), “Normal” (20–24), “Overweight” (25–29), and “Obese” (≥30).

Laboratory methods – fibrinogen
In ARIC, blood was drawn after an 8-hour fasting period with minimal trauma from an antecubital vein. Samples were processed by a standardized protocol and stored at −70°C until assayed at the ARIC Hemostasis Laboratory at the University of Texas Medical School, Houston, TX. Detailed methods for blood processing and measurement of hemostatic variables have been published elsewhere. Fibrinogen was measured by the thrombin-time titration method. Reliability coefficients
obtained from repeated testing of individuals over several weeks were 0.72 for fibrinogen.\textsuperscript{16}

In CHS, baseline fibrinogen levels in 1989–1990 were measured with a BBL fibrometer (Becton Dickinson, Cockeysville, MD) by the Clauss methods with Dade fibrinogen calibration reference (Baxter-Dade, Bedford, MA) and bovine thrombin (Parke-Davis, Lititz, PA).\textsuperscript{17}

**Pulmonary function testing**

In both studies, spirometry was conducted using a volume displacement, water-sealed spirometer. At least three acceptable spirograms were obtained from a minimum of five forced expirations. The best single spirogram was identified by computer and confirmed by a technician. Quality assurance was provided by the CHS Pulmonary Function Center for CHS and the ARIC investigators for ARIC, and the procedures followed contemporary American Thoracic Society guidelines.\textsuperscript{18} Several measures of lung function were used in this analysis: the FEV\textsubscript{1}, the FVC, and the FEV\textsubscript{1}/FVC ratio. We used the prediction equations derived from the Third National Health and Nutrition Examination Survey to define the predicted values of FEV\textsubscript{1} and FVC.\textsuperscript{19} We defined a subject as having a respiratory symptom if they reported cough, phlegm, dyspnea, or wheeze.

**Definition of outcomes**

Death was defined as all-cause mortality. COPD-related hospitalizations were people with hospitalization code of ICD-9 490–496 at any time after baseline. Rapidly declining lung function comprised of people who, between the first and second spirometries, were in the highest quartile of FEV\textsubscript{1} decline (determined as an absolute difference in the baseline minus the follow-up FEV\textsubscript{1} and divided by the interval between the two tests to obtain FEV\textsubscript{1} loss per year). Lung function decline was also determined as a proportion of the baseline FEV\textsubscript{1} and relative to the predicted FEV\textsubscript{1} at baseline, but only the absolute difference was used in the analyses. We also determined the proportion of people who were unable to obtain a follow-up spirometry. The development of COPD was defined as GOLD Stage 2 or higher disease in people who at baseline did not have GOLD Stage 2 or higher disease.

**Statistical analyses**

Data analysis was completed with SAS (v 9.2; SAS Institute, Cary, NC) and SUDAAN (v 9.0; RTI, Research Triangle Park, NC). Descriptive statistics and frequency distributions were calculated for the studied population and its relationship to fibrinogen levels and predictors (GOLD Status, CVD, smoking status, diabetes mellitus). Linear regression model using SUDAAN procedure REGRESS was used to determine correlates of fibrinogen levels controlling for age, sex, race/ethnicity, body mass index, smoking status, modified GOLD stage, diabetes, CVD, and educational level. These were replicated examining fibrinogen levels in the top decile (>393.0 mg/dL) using the SUDAAN procedure RLOGIST, controlling for these same potential confounders. Our primary outcome of interest in the survival models was time to death and COPD-related hospitalization. Cox proportional hazard regression models were developed using the SUDAAN procedure SURVIVAL to account for differential follow-up in cohort participants. (Censoring occurred at the date of death on certificate or the date the participant was last known to be alive for mortality, and date of hospitalization or date last known to be alive for COPD-related hospitalization.) The SUDAAN procedure RLOGIST was used to determine the relation between fibrinogen levels and elevated fibrinogen levels and not having a follow-up spirometry, being in the highest quartile of FEV\textsubscript{1} decline, and having incident COPD.

**Results**

The combined cohorts of ARIC and CHS included 20,993 participants at baseline. We excluded 311 who had not obtained fibrinogen levels, 209 who did not have pulmonary function testing done, and 281 subjects missing other key baseline data leaving 20,192 subjects in the combined cohorts. The mean follow-up time was 9.7 years, with a maximum of 12.1 years. By the end of the follow-up time, 2995 subjects (14.8\%) had died.

Table 1 shows the distribution of age, sex, race/ethnicity, body mass index, smoking status, modified GOLD stage, diabetes mellitus, CVD, and educational level, including the actual numbers of studied subjects and the percentage. This Table also reports the mean fibrinogen levels, the proportion of subjects with fibrinogen levels >393.0 mg/dL, and the deaths per 1000 person-years of follow-up.

Tables 2 and 3 report the correlates of fibrinogen levels (Table 2) and the correlates of elevated fibrinogen levels (Table 3). There was considerable overlap between these two analyses; for example, in both analyses older age, current smoking, and the presence of diabetes or CVD were associated with higher fibrinogen levels. The presence of severe or very severe COPD was one of the strongest predictors of fibrinogen levels, with a mean increase of 24.71 mg/dL (standard error 3.13 mg/dL) in the linear regression models and an odds ratio for elevated fibrinogen of 2.28 (95\% confidence interval: 1.79–2.95).
Table 4 shows the proportion of participants with the outcomes of interest. During follow-up, 14.8% of the cohort died, 7.5% experienced a COPD-related hospitalization, and 16.3% did not obtain a follow-up spirometry. The table also shows 16,935 subjects had a follow-up spirometry and of these, 14,848 did not have Stage 2 or higher COPD at baseline. Of the latter, 4.9% were found to have Stage 2 or higher COPD in follow-up.

The quartiles of lung function change are displayed in Table 5. The most rapidly declining quartile lost 127 mL of FEV$_1$ annually, corresponding to 4.7% (annually) of their baseline value and 4.3% of their predicted baseline value.

Table 6 displays the unadjusted and fully adjusted models for either continuous fibrinogen or elevated fibrinogen (>393 mg/dL) predicting death, COPD-related hospitalization, missing spirometry, rapidly declining lung function, and incident COPD. Fibrinogen was a significant predictor of all of these outcomes with the exception of being in the most rapidly declining FEV$_1$ quartile.
Figures 1–3 depict the interaction between fibrinogen levels >393 mg/dL and modified GOLD stage in predicting mortality, COPD-related hospitalization, and incident COPD.

**Discussion**

This analysis determined that fibrinogen levels were related to spirometrically determined obstructive lung disease, with evidence of a dose-response effect for COPD. Subjects with more advanced COPD (Stages 3 or 4) had a greater elevation in fibrinogen (24.71 mg/dL, \( P = 0.0000 \)) than GOLD Stage 2 or GOLD Stage 1 disease (12.04 mg/dL, \( P = 0.0000 \) and 7.68 mg/dL, \( P = 0.0000 \) respectively), relative to people with normal lung function. In addition, we determined that higher fibrinogen levels were related to an increased risk of death, COPD-related hospitalization, and the development of COPD.
Table 4 Proportion of subjects with the outcome of interest: death, any COPD-related hospitalization during follow-up, those missing follow-up spirometry, highest quartile of FEV₁ decline, and incident Stage 2 or higher COPD (among those free of COPD at baseline)

| Covariate                  | Death n = 20,192 | COPD-related hospitalization n = 20,192 | Missing follow-up spirometry n = 20,192 | Highest quartile of FEV₁ decline n = 16,935 | Incident Stage 2 COPD n = 14,648 |
|----------------------------|-----------------|----------------------------------------|----------------------------------------|-------------------------------------------|-----------------------------------|
| Age                        |                 |                                        |                                        |                                           |                                   |
| 45–49                      | 3.9             | 1.1                                    | 10.5                                   | 24.4                                      | 2.7                               |
| 50–54                      | 6.4             | 2.4                                    | 10.3                                   | 26.1                                      | 3.1                               |
| 55–59                      | 10.6            | 4.4                                    | 10.9                                   | 26.6                                      | 4.4                               |
| 60–64                      | 16.9            | 7.9                                    | 14.1                                   | 27.9                                      | 5.4                               |
| 65–68                      | 17.8            | 15.6                                   | 21.8                                   | 22.4                                      | 8.9                               |
| 69–72                      | 24.4            | 18.4                                   | 25.7                                   | 21.9                                      | 9.2                               |
| 73–76                      | 33.0            | 20.4                                   | 31.9                                   | 19.9                                      | 10.5                              |
| 77–80                      | 48.1            | 22.9                                   | 40.7                                   | 18.8                                      | 13.7                              |
| >80                        | 72.1            | 19.4                                   | 60.1                                   | 18.8                                      | 10.7                              |
| Sex                        |                 |                                        |                                        |                                           |                                   |
| Male                       | 11.7            | 6.0                                    | 15.8                                   | 18.6                                      | 4.5                               |
| Female                     | 18.8            | 9.3                                    | 16.8                                   | 33.3                                      | 5.3                               |
| Race-ethnicity             |                 |                                        |                                        |                                           |                                   |
| White                      | 14.9            | 8.6                                    | 14.9                                   | 26.1                                      | 5.4                               |
| Black                      | 14.5            | 3.2                                    | 21.5                                   | 21.1                                      | 3.1                               |
| Body mass index            |                 |                                        |                                        |                                           |                                   |
| <20                        | 25.8            | 15.3                                   | 23.0                                   | 22.5                                      | 7.9                               |
| 20–25                      | 14.8            | 7.9                                    | 16.0                                   | 24.6                                      | 5.7                               |
| 25–30                      | 14.7            | 6.9                                    | 15.2                                   | 26.2                                      | 4.7                               |
| >30                        | 13.4            | 6.6                                    | 17.2                                   | 24.4                                      | 4.0                               |
| Smoking status             |                 |                                        |                                        |                                           |                                   |
| Current smoker             | 17.8            | 12.3                                   | 19.2                                   | 31.0                                      | 8.6                               |
| Former smoker              | 16.3            | 8.7                                    | 16.1                                   | 26.3                                      | 5.1                               |
| Never smoker               | 11.8            | 3.6                                    | 14.7                                   | 20.8                                      | 2.9                               |
| GOLD stage                 |                 |                                        |                                        |                                           |                                   |
| Stage 3 or 4               | 44.8            | 46.5                                   | 38.3                                   | 19.6                                      | N/A                               |
| Stage 2                    | 23.5            | 19.6                                   | 23.5                                   | 25.5                                      | N/A                               |
| Stage 1                    | 21.6            | 10.6                                   | 19.6                                   | 31.7                                      | 15.0                              |
| Symptoms only              | 13.0            | 4.3                                    | 14.4                                   | 27.0                                      | 2.6                               |
| Restrictive                | 15.3            | 6.5                                    | 19.2                                   | 13.4                                      | 8.3                               |
| None                       | 8.2             | 1.8                                    | 11.0                                   | 26.6                                      | 1.6                               |
| Diabetes mellitus          |                 |                                        |                                        |                                           |                                   |
| Yes                        | 29.0            | 10.3                                   | 27.0                                   | 23.5                                      | 5.9                               |
| No                         | 12.8            | 7.1                                    | 14.7                                   | 25.3                                      | 4.8                               |
| Cardiovascular disease     |                 |                                        |                                        |                                           |                                   |
| Yes                        | 29.6            | 15.8                                   | 25.7                                   | 27.9                                      | 7.5                               |
| No                         | 12.2            | 6.0                                    | 14.6                                   | 24.7                                      | 4.5                               |
| Education level            |                 |                                        |                                        |                                           |                                   |
| <12                        | 22.6            | 11.8                                   | 25.7                                   | 23.4                                      | 5.8                               |
| 12 years                   | 12.1            | 6.9                                    | 14.3                                   | 24.8                                      | 5.1                               |
| >13 years                  | 12.5            | 5.4                                    | 12.5                                   | 26.2                                      | 4.3                               |
| Fibrinogen >393            |                 |                                        |                                        |                                           |                                   |
| Yes                        | 28.5            | 14.8                                   | 29.1                                   | 25.3                                      | 8.4                               |
| No                         | 13.3            | 6.7                                    | 14.9                                   | 25.1                                      | 4.6                               |
| Total                      | 14.8            | 7.5                                    | 16.3                                   | 25.1                                      | 4.9                               |

Abbreviations: COPD, chronic obstructive pulmonary disease; FEV₁, forced expiratory volume in 1 second.

Our findings are consistent with those of Dahl et al⁹ who showed that increased levels of plasma fibrinogen were associated with reduced lung function and increased risk of COPD, independent of smoking status. Jiang et al²⁰ also found that higher levels of baseline fibrinogen were cross-sectionally associated with lower lung function and greater longitudinal declines in FEV₁/FVC ratio in the elderly. Two population based studies²¹,²² showed that stable COPD patients have a pro-inflammatory state with increased levels of acute-phase reactants. Kalhan et al⁸ found...
that participants in the highest year 7 fibrinogen had greater FEV$_1$ and FVC decline.

This analysis demonstrates that fibrinogen levels in this cohort were significantly related to a number of factors, including age, sex, race/ethnicity, current smoking status, overweight and obesity, presence of chronic diseases such as CVD, diabetes mellitus, and COPD. A previous study$^{22}$ has shown that obstructive and restrictive lung diseases were predictors of increased levels of plasma fibrinogen. As fibrinogen levels were only determined at baseline we cannot speculate whether elevated fibrinogen caused COPD or COPD caused elevated levels of fibrinogen. It is still unknown and uncertain how and why individuals with COPD develop systemic inflammation. Whatever is the mechanism of COPD development, previous studies have shown individuals with accelerated decline in lung function are at an increased risk of COPD hospitalizations in the future. Engstrom et al showed increased incidence of hospital admissions for COPD in those with raised fibrinogen.$^{23}$

We have demonstrated an effect of elevated fibrinogen on mortality, COPD-hospitalizations, and incident Stage 2 COPD in the overall cohort (Table 6). The finding of elevated fibrinogen and COPD is consistent with findings from previous studies.$^{24}$

### Table 5 Quartiles of lung function decline among subjects who had baseline and follow-up pulmonary function measurement (n = 16,935) and baseline FEV$_1$, the mean annualized change in their absolute FEV$_1$, annualized FEV$_1$ change as a percentage of the baseline value, and annualized FEV$_1$ change as a percentage of the baseline predicted FEV$_1$ value

| FEV$_1$ change quartiles | n  | Baseline FEV$_1$ (percent predicted) | Change in FEV$_1$ (SD) | Change in FEV$_1$ as percentage of baseline (SD) | Change in FEV$_1$ as percentage of predicted (SD) |
|-------------------------|----|-------------------------------------|------------------------|-----------------------------------------------|-----------------------------------------------|
| 1                       | 4322 | 98.1                                 | −127 (63)              | −4.7 (2.6)                                     | −4.3 (2.2)                                     |
| 2                       | 4306 | 93.9                                 | −61 (10)               | −2.4 (0.9)                                     | −2.2 (0.5)                                     |
| 3                       | 4307 | 91.2                                 | −28 (9)                | −1.2 (0.6)                                     | −1.0 (0.4)                                     |
| 4                       | 4310 | 88.0                                 | 33 (72)                | 1.9 (6.0)                                      | 1.2 (2.8)                                      |

**Abbreviations:** FEV$_1$, forced expiratory volume in 1 second; SD, standard deviation.

### Table 6 Results of unadjusted and fully adjusted logistic and Cox proportional hazards models predicting death, any COPD-related hospitalization during follow-up, those missing follow-up spirometry, highest quartile of FEV$_1$ decline, and incident Stage 2 or higher COPD (among those free of COPD at baseline), with fibrinogen (per 100 mg/dL increase) or fibrinogen >393 mg/dL as the main predictors

| Outcome                      | Risk per 100 mg/dL increase in fibrinogen (unadjusted) | Risk per 100 mg/dL increase in fibrinogen (adjusted)$^*$ | Risk among subjects with fibrinogen levels >393 mg/dL (unadjusted) | Risk among subjects with fibrinogen levels >393 mg/dL (adjusted)$^*$ |
|------------------------------|-------------------------------------------------------|----------------------------------------------------------|------------------------------------------------------------------|------------------------------------------------------------------|
| Death n = 20,192             | 1.69 (1.62, 1.77)                                      | 1.31 (1.24, 1.38)                                         | 2.39 (2.18, 2.62)                                                | 1.54 (1.39, 1.70)                                                |
| Hazard ratio (95% confidence interval) | 1.76 (1.66, 1.87)                                      | 1.30 (1.21, 1.39)                                         | 2.52 (2.22, 2.87)                                                | 1.45 (1.27, 1.67)                                                |
| COPD-related hospitalization n = 20,192 | 1.67 (1.58, 1.76)                                      | 1.25 (1.18, 1.33)                                         | 2.35 (2.12, 2.61)                                                | 1.52 (1.35, 1.71)                                                |
| Hazard ratio (95% confidence interval) | 1.00 (0.95, 1.06)                                      | 1.07 (1.01, 1.14)                                         | 1.01 (0.89, 1.14)                                                | 1.08 (0.95, 1.23)                                                |
| Missing follow-up spirometry n = 20,192 | 1.49 (1.34, 1.65)                                      | 1.17 (1.03, 1.33)                                         | 1.90 (1.52, 2.38)                                                | 1.36 (1.07, 1.74)                                                |
| Odds ratio (95% confidence interval) | 1.49 (1.34, 1.65)                                      | 1.17 (1.03, 1.33)                                         | 1.90 (1.52, 2.38)                                                | 1.36 (1.07, 1.74)                                                |

**Note:** $^*$Adjusted for age, sex, race, education level, body mass index, smoking status, diabetes mellitus, cardiovascular disease, and GOLD stage.

**Abbreviations:** COPD, chronic obstructive pulmonary disease; FEV$_1$, forced expiratory volume in 1 second.
fibrinogen and the higher risk of mortality, COPD-hospitalizations, and incident Stage 2 COPD raises an option that fibrinogen could be used as a biomarker of disease activity in COPD and potential target for therapeutic interventions.

To our knowledge, this is one of the few studies with follow-up data, and although fibrinogen level was only available at baseline, partial temporal association of fibrinogen and lung function can be taken into account, although other cohort studies with lung specimens, exacerbations, and imaging are needed to validate the findings. The study has adequate sample size and power. Bias due to investigator’s knowledge of disease or risk factor seems unlikely since plasma fibrinogen was measured without the knowledge of lung function test results or disease status of subjects.

This analysis has several limitations. Analysis was only done using baseline fibrinogen levels, as follow-up fibrinogen levels were not available on all subjects. Another limitation is the unavailability of post-bronchodilator lung function-measurement, information on COPD exacerbations other than hospitalized exacerbations, and lung imaging for COPD. Not all subjects completed pulmonary function testing, biasing our sample towards a healthier population.

Figure 1 Risk of mortality by modified GOLD stage and elevated fibrinogen level.
Abbreviations: ARIC, Atherosclerosis Risk in Communities Study; CHS, Cardiovascular Health Study; GOLD, Global Initiative on Chronic Obstructive Lung Disease.

Figure 2 Risk of chronic obstructive pulmonary disease-related hospitalization by modified GOLD stage and elevated fibrinogen level.
Abbreviations: ARIC, Atherosclerosis Risk in Communities Study; CHS, Cardiovascular Health Study; GOLD, Global Initiative on Chronic Obstructive Lung Disease.
In addition, although we have adjusted for potential confounders there remains residual confounding due to COPD disease pathology based on genetic constitution, nutritional intake, environmental exposures, alcohol intake, and other infections causing systemic inflammation. These findings need to be replicated in other cohorts, including those with longitudinal measurements of fibrinogen.

**Conclusion**

In conclusion, we have demonstrated that elevated plasma fibrinogen was associated with COPD; predicting higher risk of mortality, COPD-hospitalizations, and incident Stage 2 disease. Fibrinogen can serve as a biomarker of the systemic component of COPD and may provide an option for targeting of interventions to patients with evidence of systemic inflammation or provide a means of stratifying patients in clinical trials. Future research is needed to evaluate whether targeted reduction in plasma fibrinogen levels will result in better COPD outcomes.

**Authors’ contributions**

DMM and DV conceived of the study, and participated in its design and coordination, completed the data analyses, and helped to draft the manuscript. HM and RTS participated in the design and coordination of the study and helped to draft the manuscript. All authors read and approved the final manuscript.

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**Disclosures**

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