

Chronic Obstructive Pulmonary Diseases: Journal of the COPD Foundation



Brief Report

Chronic Obstructive Pulmonary Disease Outcomes at Veterans Affairs Versus Non-Veterans Affairs Hospitals

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Abbreviations: chronic obstructive pulmonary disease, **COPD**; U.S. Centers for Medicare and Medicaid Services, **CMS**; Veterans Health Administration, **VHA**; Veterans Affairs, **VA**; confidence interval, **CI**

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Introduction

Randomized clinical trials of various interventions designed to reduce hospital readmissions in patients with chronic obstructive pulmonary disease (COPD) have demonstrated inconsistent results, including reductions in readmissions,^{1,2} an increase in the risk of readmissions or emergency department visits,³ and an increase in the risk of death⁴ compared to usual care. Results of these and other clinical trials in COPD⁵ suggest that efforts to reduce readmissions may be variably successful depending on the intervention, patient population, and health care setting.

In the last decade, the U.S. Centers for Medicare and Medicaid Services (CMS) and the Veterans Health Administration (VHA) have prioritized reducing avoidable readmissions. Among Medicare beneficiaries, an observational study found that the implementation of CMS financial incentives to reduce 30-day readmissions in patients hospitalized for heart failure or pneumonia was associated with significant increases in 30-day post-hospital mortality.⁶ These increases in mortality were primarily driven by patients who were not readmitted. In contrast,

among U.S. veterans hospitalized for heart failure, readmission rates declined while mortality rates remained stable.⁷ Little is known about the relative performance of Veterans Affairs (VA) hospitals and non-VA hospitals on COPD readmissions and mortality following the implementation of policies^{8,9} intended to incentivize hospitals to reduce COPD readmissions.

The primary objective of this study was to characterize 30-day readmission and mortality rates of patients hospitalized for COPD at VA and non-VA hospitals. Given the potential for harm when hospitals implement strategies to reduce readmissions, we also examined the association between readmission and mortality rates across all U.S. hospitals and separately in VA and non-VA hospitals. Moreover, we examined the association between readmission and mortality rates according to various hospital characteristics to determine whether the association is consistent across different hospitals. Preliminary results were presented at the May 2020 American Thoracic Society International Conference.¹⁰

Methods

Hospital-level risk-standardized 30-day readmission and mortality rates in both VA and non-VA hospitals were obtained from CMS Hospital Compare and included Medicare beneficiaries hospitalized for COPD over a 3-year period from July 1, 2015 to June 30, 2018.^{11,12} Risk-standardized readmission and mortality rates accounted for patient age, comorbid conditions, and indicators of frailty.^{12,13}

We further classified non-VA hospitals according to teaching status, safety-net status, and urban versus rural location. The data required to classify hospitals using these characteristics were obtained from the 2015 Final Rule Impact File¹⁴ and were not available for VA hospitals and non-VA critical access hospitals¹⁵ since they do not receive reimbursement adjustments under the CMS Inpatient Prospective Payment System. Teaching hospitals were defined¹⁶ as having a resident-to-bed ratio >0. Safety-net hospitals were defined as hospitals in the top quartile of the CMS Disproportionate Share Hospital patient percentage, consistent with previous studies.¹⁷ Urban or rural designation by CMS was determined by the core-based statistical area in which the hospital is located.¹⁸

Comparisons between VA and non-VA hospitals were evaluated using *t*-tests and 95% confidence intervals. Associations between hospital-level readmission and mortality rates were evaluated using linear regression models in all hospitals and within VA and non-VA strata. We report the beta coefficient with its 95% confidence interval (CI) to quantify the relationship between the hospital-level readmission and mortality rates. Effect modification by VA/non-VA status and by hospital characteristics was tested using an interaction term in separate linear regression models. If effect modification by hospital characteristic was present, we conducted additional stratified analyses. A 2-tailed *p*-value<0.05 was considered significant. Analyses were performed using R (version 3.6.0, Vienna, Austria).

Results

The mean (range) risk-standardized 30-day readmission rates were 15.3% (11.8% to 20.2%) at 127 VA hospitals and 19.5% (15.8% to 26.1%) at 3575 non-VA hospitals (Table 1). The mean (range) risk-standardized 30-day mortality rates were 6.0% (3.2% to 10.5%) at 126 VA hospitals and 8.5% (4.9% to 14.3%) at 3538 non-VA hospitals. VA hospitals had a 4.2% lower (95% CI -4.5% to -3.9%; *p*-value<0.001) mean readmission rate and 2.6% lower (95% CI -2.8% to -2.4%; *p*-value<0.001) mean mortality rate compared to non-VA hospitals. Among non-VA hospitals, those classified as non-teaching hospitals, non-safety-net hospitals, and rural hospitals had significantly lower readmission rates and higher mortality rates compared to teaching hospitals, safety-net hospitals, and urban hospitals, respectively.

Data for both readmission and mortality rates were available in 126 VA hospitals and 3523 non-VA hospitals. When VA and non-VA hospitals were aggregated, higher readmission rates were associated with significantly higher mortality rates (Figure 1) and VA/non-VA status did not modify the association (interaction *p*-value=0.44). However, in analyses stratified by hospital type, higher readmission rates were associated with significantly lower mortality rates among non-VA hospitals (Figure 2), with 50 fewer deaths per 1000 more readmissions. A similar, but non-significant inverse association was observed among VA hospitals.

Table 1. Risk-standardized 30-day All-Cause Readmission and Mortality Rates Among Patients Hospitalized for COPD Exacerbations by Hospital Characteristic, 2015 to 2018

Hospital Characteristic	30-day Readmissions, % Mean (SE)	Difference, ^a % Mean (95% CI)	p-value	30-day Mortality, % Mean (SE)	Difference, ^a % Mean (95% CI)	p-value
VA Hospitals ^b	15.3 (0.17)	-4.2 (-4.5, -3.9)	<0.001	6.0 (0.11)	-2.6 (-2.8, -2.4)	<0.001
Non-VA Hospitals ^c	19.5 (0.02)	Reference		8.5 (0.02)	Reference	
Among Non-VA Hospitals						
Teaching Hospitals	19.6 (0.04)	Reference		8.4 (0.04)	Reference	
Non-teaching Hospitals	19.5 (0.03)	-0.1 (-0.2, -0.02)	0.021	8.6 (0.03)	0.2 (0.1, 0.3)	<0.001
Safety-net Hospitals	19.7 (0.04)	Reference		8.4 (0.04)	Reference	
Non-Safety-net Hospitals	19.5 (0.02)	-0.2 (-0.3, -0.1)		8.6 (0.02)	0.2 (0.05, 0.2)	0.003
Urban Hospitals	19.6 (0.03)	Reference		8.5 (0.03)	Reference	
Rural Hospitals	19.4 (0.04)	-0.2 (-0.3, -0.1)	0.001	8.7 (0.04)	0.2 (0.08, 0.3)	<0.001
Missing ^d	19.4 (0.03)			8.6 (0.03)		

^aDifference is calculated as the risk-standardized 30-day readmission or mortality rate for hospitals with the indicated hospital characteristic minus the risk-standardized 30-day readmission or mortality rate for hospitals in the reference category.

^bVeterans Affairs hospitals. Readmissions data were available in 127 hospitals and mortality data were available in 126 hospitals.

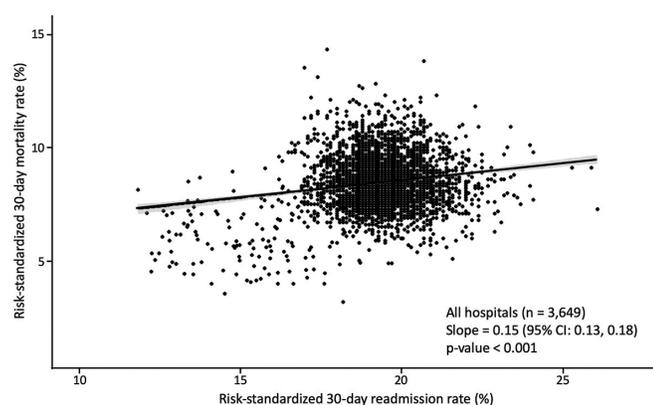
^cNon-Veterans Affairs hospitals include non-federal acute care hospitals in the United States. Readmissions data were available in 3575 hospitals and mortality data were available in 3538 hospitals.

^dOf the non-VA hospitals with data available for readmissions (n=3575) and mortality (n=3538), data was not available to classify hospitals according to teaching status, safety-net status, and urban/rural location in 759 hospitals and 734 hospitals, respectively.

Nearly all (95%) hospitals with missing data to classify according to these hospital characteristics were critical access hospitals, which by CMS definition are small (<25 inpatient beds) and located at least 35 miles driving distance from the nearest hospital.¹⁵

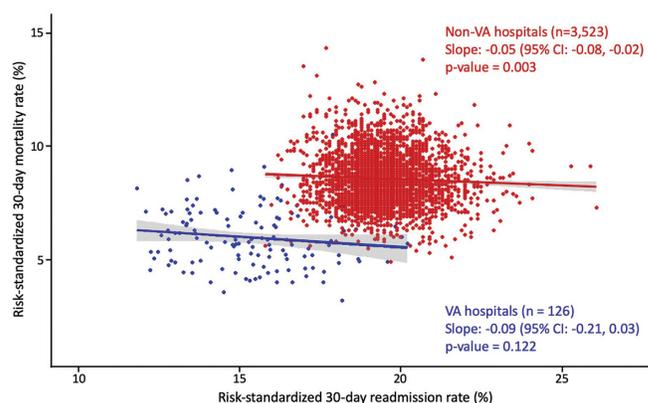
COPD=chronic obstructive pulmonary disease; SE=standard error; CI=confidence interval; VA=Veterans Affairs

Figure 1. Association Between 30-Day Readmission Rates and 30-Day Mortality Rates, Veterans Affairs and non-Veterans Affairs Hospitals Combined



The circles represent individual hospitals. The line and shading represent the results of a linear regression model and 95% confidence intervals. A 1% higher readmission rate was associated with a 0.15% (95% CI 0.13% to 0.18%; p-value<0.001) higher mortality rate across all hospitals.

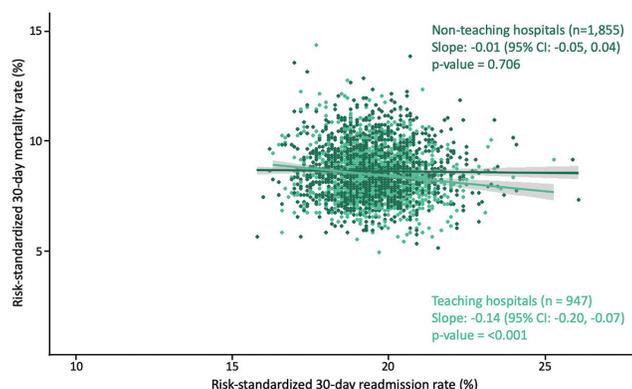
Figure 2. Association Between 30-Day Readmission Rates and 30-Day Mortality Rates, Stratified by Hospital Type



The circles represent individual hospitals (VA hospitals in blue, non-VA hospitals in red). The lines and shading represent the results of linear regression models and 95% confidence intervals. Among non-VA hospitals, a 1% higher readmission rate was associated with a 0.05% (95% CI -0.08% to -0.02%; p-value=0.003) lower mortality rate. Readmission rates and mortality rates were not significantly associated at VA hospitals (p-value=0.122).

Among the 3523 non-VA hospitals with available data for both readmission and mortality rates, data for teaching status, safety-net status, and urban/rural location were available in 2802 (80%) hospitals. Of the 721 non-VA hospitals without data available to classify according to those hospital characteristics, 687 (95%) were critical access hospitals. Teaching hospital status modified the association between readmission and mortality rates (interaction p -value=0.001) among non-VA hospitals, but this was not observed for other hospital characteristics (p -value for all interactions >0.5). Among teaching hospitals, we observed 140 fewer deaths per 1000 more readmissions (Figure 3).

Figure 3. Association Between 30-Day Readmission Rates and 30-Day Mortality Rates, non-Veterans Affairs Hospitals Stratified by Teaching Status



The circles represent individual hospitals (teaching hospitals in light green, non-teaching hospitals in dark green). The lines and shading represent the results of linear regression models and 95% confidence intervals. Among teaching hospitals, a 1% higher readmission rate was associated with a 0.14% (95% CI -0.20% to -0.07%; p -value<0.001) lower mortality rate. Readmission rates and mortality rates were not significantly associated at non-teaching hospitals (p -value=0.706).

Discussion

We found that VA hospitals had lower risk-standardized 30-day readmission and mortality rates in patients hospitalized for COPD compared to non-VA hospitals from July 2015 to June 2018, 3 years following the implementation of policies intended to reduce COPD readmissions.^{8,9} Among non-VA hospitals, higher readmission rates were also associated with lower mortality rates. The inverse association between readmissions and mortality observed among non-VA hospitals varied according

to teaching hospital status.

Our findings extend observations noted prior to and after implementation of policies to reduce readmissions. VA hospitals outperformed non-VA hospitals on mean risk-standardized 30-day readmissions (3.7% lower) and mortality (1.9% lower) in patients with COPD from 2012 to 2015,¹⁹ suggesting the gaps in readmissions and mortality may be widening (corresponding values in current report 4.2% and 2.6%, respectively). In an analysis limited to 121 of 306 U.S. hospital referral regions from 2014 to 2017, the risk-standardized 30-day mortality rate in patients with COPD also favored VA hospitals (3.5% lower).²⁰ Several possible explanations for results favoring VA hospitals in patients hospitalized for COPD warrant further investigation, including the availability of post-hospital transitional care services (e.g., arranging follow-up appointment), effects of financial penalties and reimbursement models on resources available to provide high-quality care, and differences in data quality or patient populations unaccounted for in the risk-adjustment models.

Interestingly, we observed a reversal of trends when comparing the association between readmission and mortality in all hospitals and when stratified by VA and non-VA hospital type. This reversal of trends is an example of Simpson's paradox, a statistical phenomenon that can occur when 1 or more unaccounted factors linked with the outcome are unequally distributed between groups.²¹ This finding highlights the challenges associated with interpreting administrative data, which lacks important clinical information; it is also relevant from a policy perspective. The Veterans Access, Choice and Accountability Act of 2014⁸ mandated the Secretary of Veterans Affairs to make patient quality and outcome data at VA hospitals publicly available through CMS Hospital Compare.¹¹ This effort to increase transparency and accountability for the VHA was followed by the VA MISSION Act of 2018,²² which further expanded veterans' access to health care outside the VA health care system. It is of interest to veterans to make informed choices about where to go for health care, and the data provided on Hospital Compare is intended to allow for comparisons between VA and non-VA hospitals. However, our finding of Simpson's paradox suggests the CMS-derived risk-adjustment models may be

inadequate for direct comparisons between VA and non-VA hospitals due to residual confounding. The lack of patient-level data in our study limits our ability to elucidate potential sources of residual confounding, but previous studies have noted that differences in coding practices and in social determinants of health among patients served by different hospitals also contribute to variations in risk-adjusted readmission and mortality rates.²³⁻²⁶ Taken together with our finding that teaching hospital status modified the association between readmissions and mortality, our study builds on existing evidence that variations in hospital readmission rates and mortality rates are likely related to the patients and communities which hospitals serve in addition to the quality of care they provide.

The inverse association between readmission and mortality rates in non-VA teaching hospitals in the current study is consistent with a randomized clinical trial that showed an increased risk of death with an intervention intended to reduce COPD readmissions.⁴ An observational study also found increases in post-hospital mortality in patients with COPD following implementation of CMS policies to reduce readmissions.²⁷ Although this finding may be partially explained by changes in coding practices,^{28,29} the potential for increases in post-discharge mortality accompanying reductions in readmissions has been reported for other conditions.⁶ It remains unclear if nationwide efforts to reduce readmissions improved or worsened patient outcomes, but our findings add to a body of literature that calls for re-examining these policies.

Strengths of our study include the use of well-established publicly reported metrics in thousands of U.S. hospitals. Our study also has several limitations. The lack of patient-level data about severity of illness or quality of care limits our ability to explain the observed patterns. Although CMS-derived risk-adjustment models do account for patient age, comorbid conditions, and indicators of frailty, they do not account for other determinants of clinical outcomes such as severity of illness at hospital discharge (e.g., hypercapnia) or adherence to prescribed care (e.g., use of home oxygen therapy for severe hypoxemia). To the extent these factors contribute to case-mix variation across hospitals, there may be residual patient-level confounding when comparing hospital-level outcomes. Also, the CMS

risk-adjustment formula for hospital readmissions does not account for deaths that occur during hospitalization or shortly after discharge. Thus, if the risk of readmission differs between those who die and those who survive, hospitals with higher numbers of deaths may have lower “risk-standardized” 30-day readmission rates as a result of selection bias.³⁰ Our analyses, which use hospital-level performance, may not necessarily hold for individual patients (ecologic fallacy). The cross-sectional study design also precludes establishing a causal association between readmissions and mortality.

Further research is needed to understand the patient, hospital, and health system determinants of readmission and mortality following hospitalization for COPD.³¹ Ongoing efforts to reduce readmissions should address modifiable risk factors that contribute to avoidable readmissions without discouraging appropriate readmissions.

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Declaration of Interest

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the submitted work. RGB is employed by the Veterans Health Administration. The submitted work does not necessarily reflect the views and positions of the Department of Veterans Affairs. VPC reports grants from the NIH during the conduct of the study and other from Vertex Pharmaceuticals outside the submitted work. SLL, YCC, EMS, and SB have nothing to disclose.

References

- Casas A, Troosters T, Garcia-Aymerich J, et al. Integrated care prevents hospitalisations for exacerbations in COPD patients. *Eur Respir J*. 2006;28(1):123-130.
doi: <https://doi.org/10.1183/09031936.06.00063205>
- Bourbeau J, Julien M, Maltais F, et al. Reduction of hospital utilization in patients with chronic obstructive pulmonary disease: a disease-specific self-management intervention. *Arch Intern Med*. 2003;163(5):585-591.
doi: <https://doi.org/10.1001/archinte.163.5.585>
- Aboumatar H, Naqibuddin M, Chung S, et al. Effect of a hospital-initiated program combining transitional care and long-term self-management support on outcomes of patients hospitalized with chronic obstructive pulmonary disease: a randomized clinical trial. *JAMA*. 2019;322(14):1371-1380.
doi: <https://doi.org/10.1001/jama.2019.11982>
- Fan VS, Gaziano JM, Lew R, et al. A comprehensive care management program to prevent chronic obstructive pulmonary disease hospitalizations: a randomized, controlled trial. *Ann Intern Med*. 2012;156(10):673-683.
doi: <https://doi.org/10.7326/0003-4819-156-10-201205150-00003>
- Prieto-Centurion V, Markos MA, Ramey NI, et al. Interventions to reduce rehospitalizations after chronic obstructive pulmonary disease exacerbations: a systematic review. *Ann Am Thorac Soc*. 2014;11(3):417-424.
- Wadhwa RK, Maddox KEJ, Wasfy JH, Haneuse S, Shen C, Yeh RW. Association of the Hospital Readmissions Reduction Program with mortality among Medicare beneficiaries hospitalized for heart failure, acute myocardial infarction, and pneumonia. *JAMA*. 2018;320(24):2542-2552.
doi: <https://doi.org/10.1001/jama.2018.19232>
- Parizo JT, Kohsaka S, Sandhu AT, Patel J, Heidenreich PA. Trends in readmission and mortality rates following heart failure hospitalization in the Veterans Affairs health care system from 2007 to 2017. *JAMA Cardiol*. 2020;5(9):1042-1047.
doi: <https://doi.org/10.1001/jamacardio.2020.2028>
- Veterans Access, Choice, and Accountability Act. Pub L No. 113-146, 128 Stat 1754 (2014).
- Centers for Medicare and Medicaid Services (CMS.gov). Hospital Readmissions Reduction Program. CMS.gov website. Updated August 24, 2020. Accessed August 2020.
<https://www.cms.gov/Medicare/Medicare-Fee-for-Service-Payment/AcuteInpatientPPS/Readmissions-Reduction-Program.html>
- LaBedz SL, Krishnan JA, Au DH, et al. Association of hospital mortality and readmission rates in COPD following implementation of the Hospital Readmissions Reduction Program in the United States. *Am J Respir Crit Care Med*. 2020;201:A5951.
doi: https://doi.org/10.1164/ajrccm-conference.2020.201.1_MeetingAbstracts.A5951
- Centers for Medicare and Medicaid Services (CMS). Hospital compare. CMS.gov website. Updated October 1, 2020. Accessed September 2020. <https://www.cms.gov/Medicare/Quality-Initiatives-Patient-Assessment-Instruments/HospitalQualityInits/HospitalCompare>
- Centers for Medicare and Medicaid Services (CMS). 2019 condition-specific readmission measures updates and specifications report. CMS.gov website. Published May 2020. Accessed June 2020. <https://www.qualitynet.org/inpatient/measures/readmission/resources#tab3>
- U.S. Department of Veterans Affairs. Strategic analytics for improvement and learning (SAIL) value model measure definitions. Va.gov website. Updated May 15, 2019. Accessed November 2020. https://www.va.gov/QUALITYOFCARE/measure-up/SAIL_definitions.asp
- Centers for Medicare and Medicaid Services (CMS). Files for FY 2015 final rule and correction order. CMS.gov website. Accessed March 2021. <https://www.cms.gov/Medicare/Medicare-Fee-for-Service-Payment/AcuteInpatientPPS/Acute-Inpatient-Files-for-Download-Items/FY2015-Final-Rule-CorrectionNotice-Files>
- Centers for Medicare and Medicaid Services (CMS). Critical access hospital. CMS.gov website. Published July 2019. Accessed March 2021. <https://www.cms.gov/Outreach-and-Education/Medicare-Learning-Network-MLN/MLNProducts/Downloads/CritAccessHospfctsh.pdf>
- Taylor DH Jr, Whellan DJ, Sloan FA. Effects of admission to a teaching hospital on the cost and quality of care for Medicare beneficiaries. *N Engl J Med*. 1999;340(4):293-299.
doi: <https://doi.org/10.1056/NEJM199901283400408>
- Joynt KE, Jha AK. Characteristics of hospitals receiving penalties under the Hospital Readmissions Reduction Program. *JAMA*. 2013;309(4):342-343.
doi: <https://doi.org/10.1001/jama.2012.94856>
- Snyder JE, Jensen M, Nguyen NX, Filice CE, Joynt KE. Defining rurality in Medicare administrative data. *Med Care*. 2017;55(12):e164-e169.
doi: <https://doi.org/10.1097/MLR.0000000000000607>
- Blay E, DeLancey JO, Hewitt DB, Chung JW, Bilimoria KY. Initial public reporting of quality at Veterans Affairs vs non-Veterans Affairs hospitals. *JAMA Intern Med*. 2017;177(6):882-885.
doi: <https://doi.org/10.1001/jamainternmed.2017.0605>
- Weeks WB, West AN. Veterans' health administration hospitals outperform non-Veterans health administration hospitals in most health care markets. *Ann Intern Med*. 2019;170(6):426-428.
doi: <https://doi.org/10.7326/M18-1540>
- Simpson EH. The interpretation of interaction in contingency tables. *J R Stat Soc Series B Stat Methodol*. 1951;13(2):238-241.
doi: <https://doi.org/10.1111/j.2517-6161.1951.tb00088.x>
- VA Mission Act. Pub L No. 115-182, 132 Stat 1393 (2018).

-
23. Lindenauer PK, Strait KM, Grady JN, et al. Variation in the diagnosis of aspiration pneumonia and association with hospital pneumonia outcomes. *Ann Am Thorac Soc*. 2018 May;15(5):562-569. doi: <https://doi.org/10.1513/AnnalsATS.201709-728OC>
-
24. Ibrahim AM, Dimick JB, Sinha SS, Hollingsworth JM, Nuliyalu U, Ryan AM. Association of coded severity with readmission reduction after the Hospital Readmissions Reduction Program. *JAMA Intern Med*. 2018;178(2):290-292. doi: <https://doi.org/10.1001/jamainternmed.2017.6148>
-
25. Barnett ML, Hsu J, McWilliams JM. Patient characteristics and differences in hospital readmission rates. *JAMA Intern Med*. 2015;175(11):1803-1812. doi: <https://doi.org/10.1001/jamainternmed.2015.4660>
-
26. Roberts ET, Zaslavsky AM, Barnett ML, Landon BE, Ding L, McWilliams JM. Assessment of the effect of adjustment for patient characteristics on hospital readmission rates: implications for pay for performance. *JAMA Intern Med*. 2018;178(11):1498-1507. doi: <https://doi.org/10.1001/jamainternmed.2018.4481>
-
27. Puebla Neira DA, Hsu ES, Kuo YF, Ottenbacher KJ, Sharma G. Readmissions reduction program: mortality and readmissions for chronic obstructive pulmonary disease. *Am J Respir Crit Care Med*. 2021;203(4):437-446. doi: <https://doi.org/10.1164/rccm.202002-0310OC>
-
28. Lindenauer PK, Yu H, Grady J, Dorsey K, Triche EW. Apparent increase in chronic obstructive pulmonary disease mortality likely an artifact of changes in documentation and coding. *Am J Respir Crit Care Med*. 2021;203(5):647-649. doi: <https://doi.org/10.1164/rccm.202009-3697LE>
-
29. Puebla Neira DA, Hsu ES, Kuo YF, Ottenbacher KJ, Sharma G. Reply to Lindenauer et al: apparent increase in chronic obstructive pulmonary disease mortality is likely an artifact of changes in documentation and coding. *Am J Respir Crit Care Med*. 2021;203(5):649-651. doi: <https://doi.org/10.1164/rccm.202010-3855LE>
-
30. Papanicolas I, Orav EJ, Jha AK. Is mortality readmissions bias a concern for readmission rates under the Hospital Readmissions Reduction Program? *Health Serv Res*. 2020;55(2):249-258. doi: <https://doi.org/10.1111/1475-6773.13268>
-
31. Press VG, Au DH, Bourbeau J, et al. Reducing chronic obstructive pulmonary disease hospital readmissions. an official American thoracic society workshop report. *Ann Am Thorac Soc*. 2019;16(2):161-170. doi: <https://doi.org/10.1513/AnnalsATS.201811-755WS>