

Improved Risk Adjustment for Comorbid Diagnoses in Administrative Claims Analyses of Orthopaedic Surgery

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INTRODUCTION:

Unlike the controlled environment and homogeneous patient populations employed in most RCTs, claims data represent “real world” data from heterogeneous populations. Thus, claims analyses require thorough risk adjustment for patients’ comorbid medical conditions. Many orthopaedic studies use only the diagnosis codes present on the index admission to identify comorbidities. However, information on coexisting medical conditions for inpatient admissions is often incomplete. The goal of this study is to utilize the best practices in comorbidity adjustment developed in other clinical disciplines and apply this to an orthopaedic population, assessing improvements in capture rates, model fit, and explanatory power. We focus on the implementation of the Elixhauser comorbidity measure in elderly hip fracture populations from three types of claims resources: Medicare, all-payer, and private-payer.

METHODS:

Inpatient admissions for hip fracture were identified through ICD-9&10-CM codes. Patients with concurrent fractures or polytrauma were excluded. Comorbidities from the Medicare cohort (2009-2018 20% RIF sample) were captured from MedPar (inpatient) and the Outpatient/Carrier base files. Comorbidities from the all-payer (2016-2020 NY HCUP) cohort were captured from inpatient, ambulatory, and emergency department (ED) databases. Comorbidities from the private payer cohort (2016-2020 insurance database) were captured from the inpatient and outpatient databases.

Elixhauser comorbidities were identified first using only the codes present on the index admission. Additional captures were then performed using increasing amounts of data: first, including inpatient hospitalizations (year prior), and second, including outpatient encounters (year prior). For HCUP, an additional capture was performed using ED data. For a comorbidity to be present, one claim from inpatient data or ≥ 2 claims occurring ≥ 30 days apart from outpatient/ED data were required to avoid “rule out” or erroneous diagnoses. Explanatory and discriminatory power of the capture strategies were assessed for in-hospital (death, LOS, and total payments/charges) and post-discharge (90-day readmission, and 90-day and 1-year death [Medicare only]) outcomes via multivariable binary logistic regression and Generalized Estimating Equations. The model performance for each capture strategy was assessed with the Akaike information criterion (AIC), and the explanatory power was quantified by the area under the receiver operating characteristic curve (AUC).

RESULTS:

The final cohorts included 389,357 patients from Medicare, 47,383 patients from NY HCUP, and 86,776 patients from insurance database. The percentage of each class of Elixhauser comorbidity missed when including only data from the index admission ranged from 9.3% to 65.6% for the Medicare cohort, 2.9% to 39% for the NY HCUP all-payer cohort, and 14.7% to 57.9% for the private-payer cohort (Figure 1). When including data from the index admission + inpatient data from the year prior between 2.0% and 45% of comorbidities were still missed for Medicare versus using all available data (0.2% to 8.5% for NY HCUP, and 2.9% to 40.0% for insurance database). The gains in capture rates with the inclusion of outpatient data from Medicare and insurance database were much greater than those achieved with the NY HCUP outpatient data, as the latter includes only claims from ambulatory surgery centers. Inclusion of ED data provided only 0.0% to 1.1% improvements in capture rates. For post-discharge metrics, the inclusion of inpatient and outpatient data from the year prior led to significant improvements in model performance ($p < 0.001$) in all cohorts (Table 1). For in-hospital death and LOS metrics, the comorbidities contained within the index admission, alone, provided the best model fit and explanatory power for all cohorts. For total inpatient payments/charges for the index hip fracture admission, the results were mixed, with the best explanatory power for the Medicare cohort resulting from all available data, whereas for the HCUP and insurance cohorts, the comorbidities captured on the index admission were sufficient.

DISCUSSION AND CONCLUSION:

Many comorbidities are missed when including only data from the index inpatient admission. The addition of inpatient and outpatient data from the year prior substantially improved the capture rate of Elixhauser comorbidities for Medicare and private payer data. For the all-payer NY HCUP data, inpatient data from the year prior substantially improved capture rates. However, data from ambulatory surgery centers and EDs provided only modest improvements. The inclusion of inpatient and outpatient data from the year prior provided substantive improvements in model performance and explanatory power for post-discharge outcomes in all cohorts. However, for in-hospital metrics, comorbidities captured on the index admission were most consequential and adding data from the year prior provided little to no improvement in model performance or explanatory power. In conclusion, our results support using all available data from the year prior wherever possible when capturing comorbidities for risk-adjusted modeling of post-discharge outcomes. These results also have significant implications for risk adjustment of quality metrics defined from claims data.

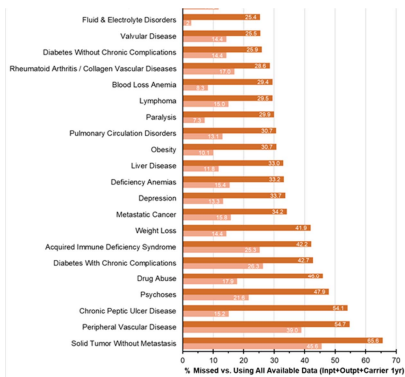


Figure 1. Percentage of comorbidities missed when using index data only and index+inpatient data from the year prior to admission versus using all available data (index + inpatient + outpatient/carrier) from the year prior to admission from the Medicare cohort.

Table 1. Model fit (lower AIC, significant improvement ≥ 2 point reduction) and explanatory power (higher AUC) for the different comorbidity capture strategies.

Cohort & Capture Strategy	In-Hospital Metrics				First Discharge Metrics				
	In-Hospital Death	LOS	30-Day Readmits for Cause of Admission	AUC	30-Day All-Cause Readmission	90-Day Death	1-Yr Death	AUC	
Medicare - No Comorbidities	246	8.92	106217	0.78171	35480	6.108	10024	0.821	0.828
Index Only	246	8.94	106229	0.78186	35492	6.108	10024	0.821	0.828
Ref 1 ^a	246	8.94	106229	0.78186	35492	6.113	10025	0.821	0.828
Ref 1 + Carrier for 2-yr	246	8.95	106221	0.78174	35487	6.117	10026	0.821	0.828
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^a Ref 1 represents 2-category comorbidity capture strategy for each outcome