

# Validation of the Combined Comorbidity Score in the ICD-10 Era: Application to High-Risk Populations

Justin Bohn, Sc.D. Emily Welch, M.P.H Sengwee Darren Toh, Sc.D.

Sentinel Operations Center Harvard Pilgrim Health Care Institute

## **Disclosures**



- The authors have no conflicts of interest to disclose
- This work was supported by the U.S. Food and Drug Administration (FDA) through the Department of Health and Human Services (HHS) Contract number HHSF223201400030I
- This presentation reflects the views of the authors and not necessarily those of the U.S. FDA

# **Background**



- In October 2015, U.S. transitioned to the 10<sup>th</sup> Revision of the International Classification of Diseases, Clinical Modification (ICD-10-CM)
  - General Equivalence Mappings (GEMs) developed by Center for Medicare & Medicaid Services
- Researchers must adapt diagnosis and procedure-based variable definitions
- Which mapping strategy will yield best possible confounding control?

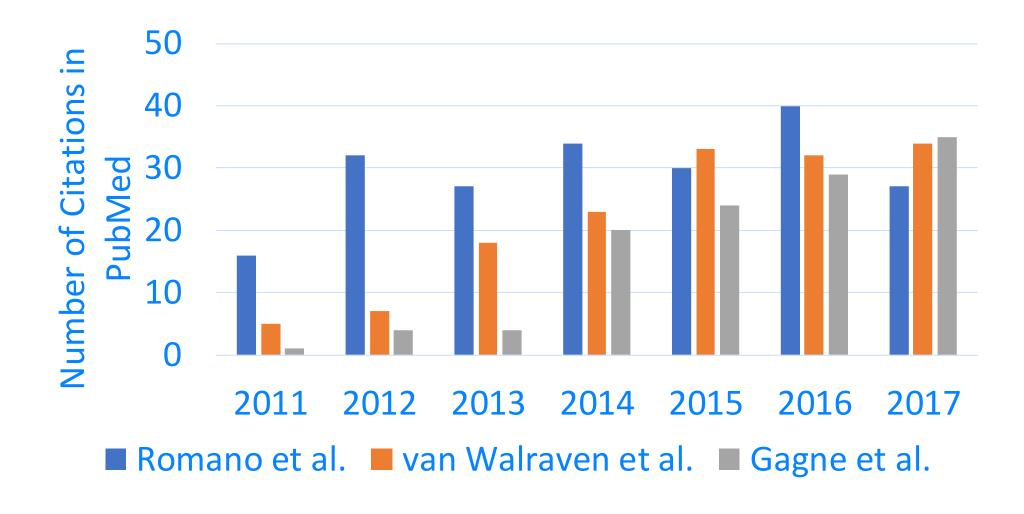
## Prior validation of ICD-10-CM-based CCIs



- In 2017, Sun et al.¹ validated four ICD-10-CM adaptations of the Charlson-Elixhauser Combined Comorbidity Index (CCI)
  - GEMs simple backward mapping (SBM)
  - GEMs forward-backward mapping (FBM)
  - Canadian mapping proposed by Quan et al.<sup>2</sup> (CA)
  - All three above mappings combined (ALL)
- Combined approach best discriminated between those rehospitalized within 30-days and those not re-hospitalized
  - Only ICD-10-CM data from Jan. Mar. 2016 available at time of study

# The rise of the combined comorbidity index





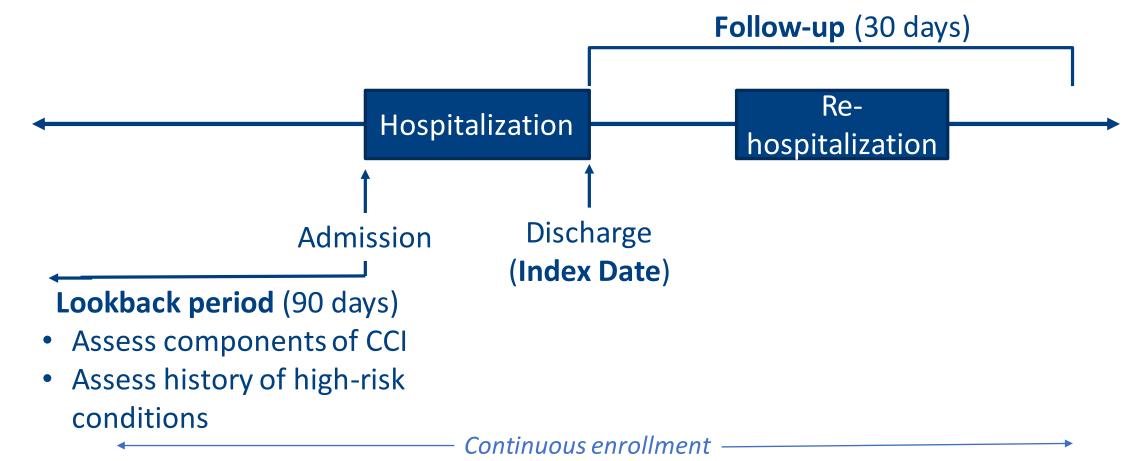
# **Objectives**



- Validate CCI as predictor of re-hospitalization in ICD-10-CM era
  - Using additional data through 2017
- Assess in commonly-studied, high-risk populations
  - Atrial fibrillation
  - Irritable bowel disease
  - Type 2 diabetes mellitus
- Sensitivity analysis
  - Vary lookback & follow-up periods: 30, 90, 183 days

# Study design & data source





Data source: Truven Health MarketScan® Research Databases

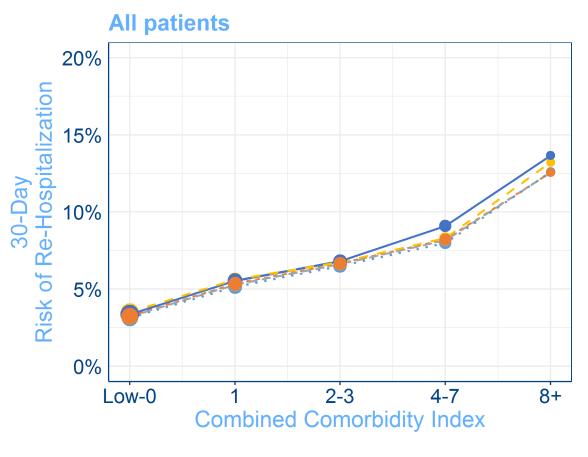
**Cohort identification:** 

Apr 2014 – Sep 2015 (ICD-9-CM) Oct 2015 – Mar 2017 (ICD-10-CM)



#### **Baseline characteristics**

Dascille characteristics			
	ICD-9-CM	ICD-10-CM	
Mean age, years	50.1	50.2	
Mean CCI	1.2	1.4 (SBM)	
		1.5 (FBM)	
		1.3 (CA)	
		1.6 (ALL)	



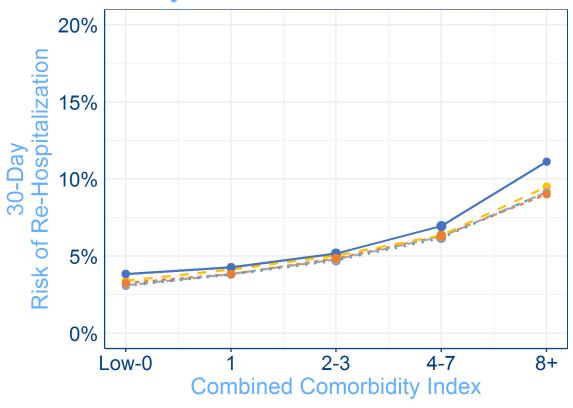
- **→** *ICD-9-CM*
- ICD-10-CM: Simple backward mapping
- Coding -- ICD-10-CM: Forward-backward mapping
  - → ICD-10-CM: Quan et al.
  - •• ICD-10-CM: All mappings



#### **Baseline characteristics**

	ICD-9-CM	ICD-10-CM
Mean age, years	74.2	74.6
Mean CCI	3.4	3.8 (SBM)
		3.9 (FBM)
		3.5 (CA)
		4.0 (ALL)

#### **History of atrial fibrillation**



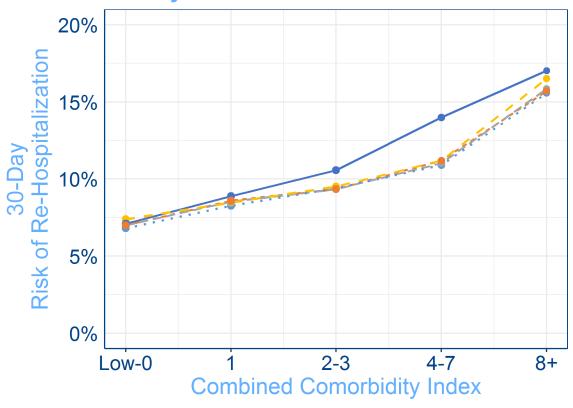
- **→** *ICD-9-CM*
- ICD-10-CM: Simple backward mapping
- **Coding** → ICD-10-CM: Forward-backward mapping
  - → ICD-10-CM: Quan et al.
  - •• ICD-10-CM: All mappings



#### **Baseline characteristics**

	ICD-9-CM	ICD-10-CM
Mean age, years	50.1	50.7
Mean CCI	1.4	1.7 (SBM)
		1.8 (FBM)
		1.5 (CA)
		2.0 (ALL)

#### **History of irritable bowel disease**



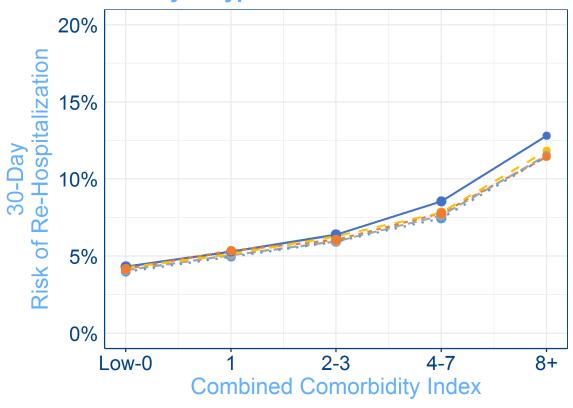
- **→** *ICD-9-CM*
- ICD-10-CM: Simple backward mapping
- Coding -- ICD-10-CM: Forward-backward mapping
  - → ICD-10-CM: Quan et al.
  - •• ICD-10-CM: All mappings



#### **Baseline characteristics**

	ICD-9-CM	ICD-10-CM
Mean age, years	63.9	64.7
Mean CCI	2.1	2.6 (SBM)
		2.8 (FBM)
		2.5 (CA)
		3.0 (ALL)

#### **History of type 2 diabetes**



- **→** *ICD-9-CM*
- ICD-10-CM: Simple backward mapping
- **Coding** → ICD-10-CM: Forward-backward mapping
  - → ICD-10-CM: Quan et al.
  - •• ICD-10-CM: All mappings

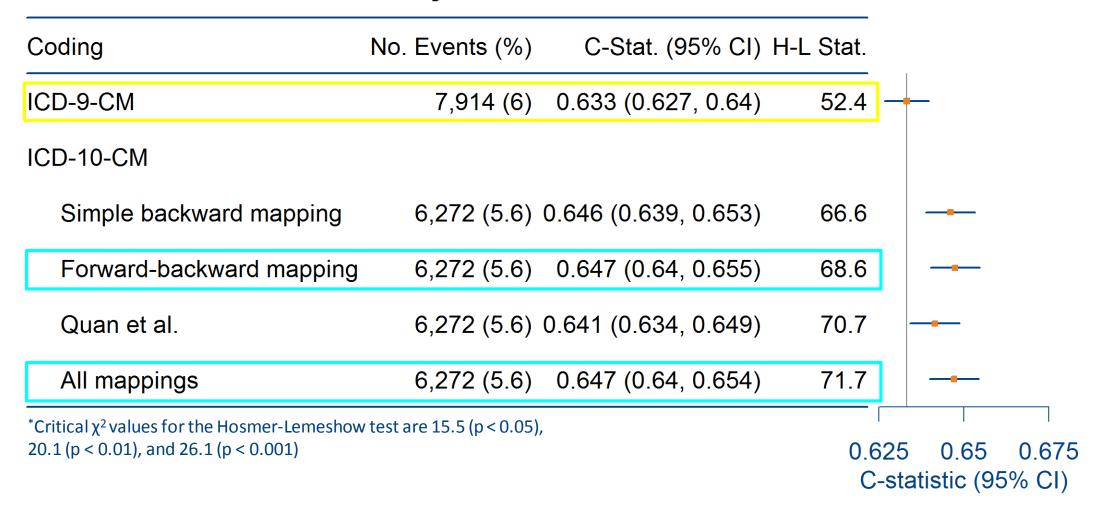


#### **All patients**

Coding	No. Events (%)	C-Stat. (95% CI)	H-L Stat.		
ICD-9-CM	74,161 (5.2)	0.645 (0.642, 0.647)	1,613.6	+	
ICD-10-CM					
Simple backward mapping	59,075 (5.2)	0.65 (0.648, 0.653)	1,405.2	-	
Forward-backward mapping	59,075 (5.2)	0.651 (0.649, 0.653)	1,344.6	-	
Quan et al.	59,075 (5.2)	0.64 (0.638, 0.643)	1,991.3		
All mappings	59,075 (5.2)	0.653 (0.65, 0.655)	1,326.2	] -	
*Critical $\chi^2$ values for the Hosmer-Lemeshov 20.1 (p < 0.01), and 26.1 (p < 0.001)	v test are 15.5 (p < 0.05)	,		625 0.65 -statistic (	



#### History of atrial fibrillation





## **History of irritable bowel disease**

Coding	No. Events (%)	C-Stat. (95% CI)	H-L Stat.	
ICD-9-CM	2,125 (9.3)	0.629 (0.617, 0.642)	10.7	
ICD-10-CM				
Simple backward mapping	1,635 (8.8)	0.633 (0.619, 0.647)	8.3	
Forward-backward mapping	1,635 (8.8)	0.633 (0.619, 0.647)	8.4	-
Quan et al.	1,635 (8.8)	0.626 (0.613, 0.64)	17.2	-
All mappings	1,635 (8.8)	0.634 (0.62, 0.648)	5.2	] ——
*Critical $\chi^2$ values for the Hosmer-Lemeshov	v test are 15.5 (p < 0.05)	),		
20.1 (p < 0.01), and 26.1 (p < 0.001)			0.6	125 0.6375
			C	-statistic (95% CI)



#### **History of type 2 diabetes**

Coding	No. Events (%)	C-Stat. (95% CI)	H-L Stat.	
ICD-9-CM	16,098 (6.3)	0.644 (0.64, 0.649)	144.6	-
ICD-10-CM				
Simple backward mapping	13,088 (6.2)	0.655 (0.65, 0.66)	129.6	
Forward-backward mapping	13,088 (6.2)	0.655 (0.65, 0.66)	101.8	
Quan et al.	13,088 (6.2)	0.651 (0.646, 0.656)	112.2	
All mappings	13,088 (6.2)	0.655 (0.65, 0.66)	81.8	
*Critical $\chi^2$ values for the Hosmer-Lemeshov 20.1 (p < 0.01), and 26.1 (p < 0.001)	v test are 15.5 (p < 0.05),	,	0.6 C-	25 0.65 0.6 statistic (95% 0

## **Limitations**



- Lack of mortality data
  - CCI initially validated as mortality predictor
- Discrimination & calibration ≠ confounding control
  - Empirical comparative safety and effectiveness examples needed
- Still early in use of ICD-10-CM
  - Future may bring validated ICD-10-CM algorithms for all CCI components

## **Conclusions**



- Replication and extension of work by Sun et al.
  - More ICD-10-CM experience, different commercial data population
- Combined mapping approach yields best discrimination in majority of settings
  - Robust to changes in lookback and follow-up duration
  - Calibration results equivocal

# **Acknowledgments**



- Sentinel Operations Center
  - Qoua Her, Pharm.D., S.M.
  - Catherine Panozzo, Ph.D.
- Brigham and Women's Division of Pharmacoepidemiology
  - Jenny Sun, S.M.
  - Jimmy Rodgers, M.S.
  - Joshua Gagne, Pharm.D., Sc.D.