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Validity and utility of ICD-10 administrative health data for identifying ST- and non–ST-elevation myocardial infarction based on physician chart review

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Abstract

Background: Health care administrative databases are useful for assessing the population-level burden of disease and examining issues related to access, costs and quality of care. In these databases, the diagnoses and procedures are coded with the use of the World Health Organization International Classification of Diseases (ICD). We examined the validity of 2 ICD-10 coding definitions for categorizing patients with acute myocardial infarction (MI) as having ST-elevation MI (STEMI) or non–ST-elevation MI (non-STEMI).

Methods: Charts of patients with acute MI discharged between April and June 2007 from 3 hospitals in Edmonton, were reviewed to define the acute MI subtype (i.e., STEMI v. non-STEMI). The agreement between clinician chart review and STEMI/non-STEMI classification based on the standard (ICD-10 I21.x) and the supplementary electrocardiogram (ECG) codes (R94.3x) was determined. We assessed the effect of these alternative definitions on in-hospital mortality estimates by applying them to the data for all patients with acute MI admitted to hospital in the province from April 2007 to March 2010.

Results: Of the 297 patients, 49.2% were identified as having STEMI based on chart review, 44.4% using the standard definition, and 44.1% using the ECG definition. Both the standard and ECG definitions provided high agreement (92% for STEMI and 100% for non-STEMI) with the chart review classification. In the larger population-level cohort (n = 15 148), use of the standard definition or the ECG definition did not affect in-hospital mortality estimates for patients with STEMI and those with non-STEMI.

Interpretation: The standard definition appears equivalent to the definition using supplementary ECG codes to subcategorize patients with acute MI as having STEMI or non-STEMI. These findings may be relevant for the development of later versions of ICD codes.

ealth care administrative databases, such as those maintained by the Centers for Medicare and Medicaid Services in the United States and by provincial and federal agencies in Canada and other countries, are extremely useful for assessing the population-level burden of disease and examining issues related to access, costs and quality of care.^{1–5} In these databases, the diagnoses and procedures are coded with the use of the World Health Organization International Classification of Diseases (ICD).⁶ As of October 2015, all developed countries, including the US, are using version 10 of the ICD. Because the World Health Organization permits its member countries to add countryspecific diagnosis codes or specify conditions using subcodes in ICD-10, several countries have developed their own ICD-10 versions.

The incidence and clinical characteristics of patients presenting with ST-elevation myocardial infarction (STEMI) and those of patients with non–ST-elevation myocardial infarction (non-STEMI) vary greatly.^{7,8} These differences underline the importance of accurately classifying acute myocardial infarction (MI) subtypes when examining patient outcomes. Historically, owing primarily to limitations in coding algorithms, studies using administrative data to examine treatment and outcomes of acute MI have grouped STEMI and non-STEMI together.^{9,10} However, the improved ICD-10 codes distinguish between patients with STEMI and those with non-STEMI.¹¹ In April 2007, the Canadian version of ICD-10 introduced secondary diagnosis codes for cardiovascular function based on the electrocardiogram (ECG) findings.¹² It is not known whether these supplementary codes are

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more accurate than the standard ICD-10 codes in differentiating between the acute MI subtypes.

The primary objective of our study was to evaluate the agreement between chart review by clinicians and the standard definition (based on ICD-10 Q-wave codes alone) and ECG definition (based on supplementary ECG codes) for classifying STEMI and non-STEMI. We also examined the extent to which in-hospital mortality estimates were affected by the use of the alternative definitions.

Methods

Defining STEMI and non-STEMI in chart data

The charts of all patients discharged from hospital with a diagnosis of acute MI in Edmonton (population of about 1 million), undergo review by a trained chart abstractor. This review is conducted as part of a dedicated quality-assurance program linked to the Vital Heart Response clinical program. This program is a regional reperfusion program focused on delivering expeditious evidence-based care to patients with STEMI.

To validate the standardized chart abstraction process and confirm appropriate patient selection, we obtained the charts of all patients discharged between Apr. 1 and June 30, 2007, from the 3 major hospitals in the Vital Heart Response program. The charts of patients with a diagnosis of acute MI were independently abstracted to obtain an approximately equal sample of patients identified as having STEMI and non-STEMI. The ECGs were then blinded. Subsequently, 2 experienced clinicians (S.S. and W.T.) interpreted the 12-lead ECG and reviewed peak myocardial enzymes to determine the clinical diagnosis of non-STEMI versus STEMI.¹³ If the 2 physicians disagreed on the diagnosis, a third physician (R.C.W.) read the ECG to determine STEMI or non-STEMI. The classification of acute MI subtype based on clinician chart review was used as the reference standard.

Defining STEMI and non-STEMI in administrative data

We used 2 definitions to categorize patients with acute MI as having STEMI or non-STEMI. The standard definition was the presence of the following codes in the primary diagnosis field: ICD-10 codes I21.0–3 for STEMI and I21.4x for non-STEMI.

A patient presenting with an ECG indicative of STEMI can progress to Q-wave MI, a non–Q-wave MI or an aborted MI, whereas those with no ST-elevation on the ECG are considered to have non-STEMI and can progress only to non–Q-wave MI or unstable angina.¹² This means that all patients with a code for Q-wave MI (I21.0–3) should have STEMI, but some with a code for non–Q-wave MI (I21.4x) could also have STEMI. Thus, the supplementary ECG codes that can distinguish between ECG results suggestive of STEMI (R94.30) or non-STEMI (R94.31) could be useful in accurately categorizing patients into the 2 subtypes.¹² The ECG codes were introduced in April 2007 in Canada, and therefore we used the charts of patients who were discharged from the 3 hospitals between April and June 2007. In our study, the ECG

definition to categorize patients with acute MI as having STEMI or non-STEMI was the presence of ICD-10 code R94.30 in any diagnosis field for STEMI and ICD-10 code R94.31 in any diagnosis field for non-STEMI.

Statistical analysis

We linked the chart data with hospital discharge abstract data from April to June 2007 using the unique identifier of personal health number. The discharge abstract data include age, sex, length of stay, up to 25 diagnoses, up to 16 procedures and discharge status. We extracted these linked Discharge Abstract Database records using the date of admission and the most responsible diagnosis of I21 (acute MI). We removed the data for patients with invalid personal health numbers to create our final sample.

We compared the agreement between the clinician chart review classification and the STEMI versus non-STEMI classification based on the standard definition and the ECG definition. The data for patients with an unspecified MI (I21.9) or an unspecified ECG result (R94.38) were excluded from the analysis. The agreement between chart review and the 2 subtype definitions was calculated overall and by sex and age.

To determine the effects of subtype definition on in-hospital mortality estimates, we applied the standard and ECG definitions to a cohort of all patients admitted to hospital with acute MI in the province of Alberta between Apr. 1, 2007, and Mar. 31, 2010. The data set for this cohort contained information on age, sex, length of stay, diagnoses, procedural interventions and in-hospital mortality. Patients who had a primary diagnosis code of I21.x were included in this assessment. A record of R94.3x (supplementary ECG code) in any of the secondary diagnosis fields was captured. The unit of analysis was the patient. To test the utility of the coding definitions, we calculated the in-hospital mortality among patients with STEMI and those with non-STEMI identified with the use of the standard and ECG definitions.

We followed the Standards for Reporting of Diagnostic Accuracy (STARD) guidelines for reporting studies of diagnostic accuracy.¹⁴ Data analysis was conducted with the use of SAS version 9.2.

Ethics approval

The ethics board of the University of Alberta felt the study was for quality-assurance purposes and did not require ethics approval.

Results

The 2 physicians reviewed 319 charts of patients with a primary diagnosis of acute MI (Figure 1). They disagreed on the classification of the acute MI as STEMI or non-STEMI in 3 cases (0.9%).

Agreement between administrative and chart data

Administrative data were linked with chart review data for 302 of the 319 patients (Figure 1). Once duplicate records and

records with invalid personal health numbers were removed, we had linked data for 297 patients. Seven patients were excluded from the validation of the standard definition because they had a code of I21.9 ("Acute myocardial infarction, unspecified"), and 8 patients were excluded from the validation of the ECG definition because they had a code of R94.38, which does not specifically define the STEMI or non-STEMI subtype, or had missing or conflicting R94.3x coding.

The number of patients classified as having STEMI or non-STEMI by chart review, standard definition and ECG definition are presented in Table 1. For STEMI, the standard definition had a higher level of agreement with chart review overall (92.3%) than did the ECG definition (91.6%). This higher level of agreement remained for men and for both younger (age < 65 yr) and older (age \geq 65 yr) patients. The ECG definition had a higher level of agreement with chart review for women (89.1%) than did the standard definition (87.0%). With both definitions, the level of agreement was lower for older patients than for younger patients.

For non-STEMI, the level of agreement between the chart review and each of the standard and ECG definitions was perfect; that is, all patients identified by chart review as having non-STEMI were classified as having non-STEMI with both definitions. There were, however, patients with STEMI who were categorized as having non-STEMI. The level of misclassification was as follows: 7.7% of patients with STEMI were classified as having non-STEMI with the standard definition, and 8.4% were classified as having non-STEMI with the ECG definition.

Effect of definitions on in-hospital mortality estimates

We applied the standard and ECG coding definitions for STEMI and non-STEMI to 15 148 patients admitted to hospital with a primary diagnosis of acute MI (I21.x) in Alberta between April 2007 and March 2010. Based on the standard definition, 5713 patients (37.7%) were classified as having STEMI and 9435 (62.3%), non-STEMI. There were 528 patients (3.5%) with an unspecified abnormal ECG result (R94.38) or missing supplementary R94.3x code, and 20 patients (0.1%) had both the code R94.30 and the code R94.31. When only the ECG definition was used to define the acute MI subtypes, 5530 patients (36.5%) were classified as having STEMI and 9070 (59.9%), non-STEMI. Table 2 summarizes these findings and shows that the calculation of



Figure 1: Steps in defining a study sample for validating coding of acute myocardial infarction (MI) subtypes. ECG = electrocardiogram, PHN = personal health number, STEMI = ST-elevation myocardial infarction. *Excluded from analysis.

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in-hospital mortality was similar irrespective of which definition was used.

Interpretation

Our study examining the validity of 2 definitions, one using standard ICD-10 codes and the second using supplementary ECG codes, for categorizing patients with acute MI as having STEMI or non-STEMI has several interesting findings. First, we found that both definitions were good at differentiating between the 2 subtypes in a sample of patients with confirmed acute MI. The agreement between both definitions and clinician chart review was high across all patient subgroups with the exception of the coding of STEMI for patients older than 65 years, where the agreement was slightly lower. Second, for STEMI, the standard definition had a slightly higher level of agreement with chart review than did the ECG definition. Third, for non-STEMI, the agreement with the chart review for both the standard definition and the ECG definition was perfect, that is, all patients identified by chart review as having non-STEMI were classified as having non-STEMI with both definitions. And finally, when we applied the 2 definitions to a database of patients with acute MI admitted to hospital, we found that the calculation of in-hospital mortality was similar irrespective of which definition was used.

The validation of STEMI and non-STEMI coding relies on the validation of acute MI first. Studies have shown that acute MI is coded well with the use of ICD. In a recent systematic review, Metcalfe and associates¹⁵ reported that there have been several studies validating the case definitions for acute MI, with most studies focused on ICD-9 and no studies validating exclusively ICD-10 codes. The reported validity of these diagnostic codes for identifying acute MI has varied from a sensitivity of 66.0% to 95.1% and a specificity of 80.2% to 100.0%, depending on the subgroup, database and area studied.¹⁵ Although that systematic review did not identify any studies validating the acute MI subtypes as defined using recently revised ICD coding definitions, ICD-9-CM¹⁶ and ICD-10¹¹ codes for acute MI have been used to differentiate between patients with STEMI and those with non-STEMI.

Studies have been conducted evaluating the validity of ICD-9 coding for acute MI. Varas-Lorenzo and colleagues¹⁷ studied Saskatchewan data from November 1999 to December 2001 and found that ICD-9 codes for acute MI had a positive predictive value of 0.95 (95% confidence interval 0.91 to 0.98) for identifying patients with acute MI based on detailed chart review as the gold standard. Although many studies validating the case definitions of acute MI using ICD coding have shown similar high positive predictive values or sensitivity,¹⁵ they have not distinguished the validity of acute MI subtypes using ICD-coded data.

Studies have been conducted in the US validating the acute MI subtype coding definitions for ICD-9-CM. With the current ICD-9-CM coding definitions, patients with non-STEMI are those with a diagnosis code of 410.7, and patients with codes 410–410.8 are classified as having STEMI.¹⁸ One US study with a very large sample (over 1 million) showed that 10.9% of patients who were found to have STEMI based on an ECG received the code 410.71, and 38.8% of patients with ECG indicative of non-STEMI had an ICD-9 code other than 410.71.¹⁹ Our study shows that the Canadian ICD-10 codes are less likely to misclassify both STEMI and non-STEMI than has previously been reported for ICD-9-CM.

Using data from 2000–2001, Woo and associates²⁰ found that MI documentation in medical charts was incomplete and inconsistent in its ability to distinguish between STEMI and non-STEMI and that up to 20% of medical records could not be used for retrospective MI-subtype–specific research. Inconsistencies in the medical record are one possible reason for the misclassification of coded information. If the information in the chart is not recorded completely and correctly, coders cannot transfer this information to the administrative data codes that could then be used to distinguish between acute MI subtypes. When we applied the subtype coding definitions in our study, we found that the charts of about 0.1% of patients in our sample had inconsistent coding, including

Table 1: Validity of the International Classification of Diseases version 10, Canadian edition, coding for acute MI subtypes												
	Reference standard		Standard definition* n = 290				ECG definition† n = 289					
	(chart	review)	No. of p	oatients	Agreen	nent, %	No. of patients		Agreement, %			
Group	STEMI	Non- STEMI	STEMI	Non- STEMI	STEMI	Non- STEMI	STEMI	Non- STEMI	STEMI	Non- STEMI		
Overall	143	147	132	158	92.3	100.0	131	158	91.6	100.0		
Women	46	55	40	61	87.0	100.0	41	60	89.1	100.0		
Men	97	92	92	97	94.8	100.0	90	98	92.8	100.0		
Age < 65 yr	82	67	81	70	98.8	100.0	79	69	96.3	100.0		
Age ≥ 65 yr	61	80	51	88	83.6	100.0	52	89	85.2	100.0		

Note: ECG = electrocardiogram, MI = myocardial infarction, STEMI = ST-elevation MI.

*I21.0-3 for STEMI v. I21.4x for non-STEMI.

†R94.30 ("ECG suggestive of STEMI") v. R94.31 ("ECG suggestive of NSTEMI"). n = 289 because the chart review for the ECG definition is missing 1 male patient under 65 years of age.

having an R94.3x code representing one subtype and an R94.3x code representing a different subtype. The lower level of agreement for the coding of STEMI for patients aged

Table 2: Application of the standard and ECG codingdefinitions to data for all patients admitted to hospitalwith acute MI in Alberta between Apr. 1, 2007, andMar. 31, 2010 ($n = 15\ 148^*$)

Group; acute MI subtype	Coding definition	Mortality, %	No. of patients	
Overall				
Standard definition				
STEMI	l21.0–3	5.51	5713	
Non-STEMI	l21.4x	3.95	9435	
ECG definition				
STEMI	R94.30	5.42	5530	
Non-STEMI	R94.31	3.89	9070	
Female				
Standard definition				
STEMI	121.0–3	8.79	1525	
Non-STEMI	l21.4x	4.82	3174	
ECG definition				
STEMI	R94.30	8.63	1460	
Non-STEMI	R94.31	4.77	3038	
Male				
Standard definition				
STEMI	l21.0–3	4.32	4188	
Non-STEMI	l21.4x	3.51	6261	
ECG definition				
STEMI	R94.30	4.28	4070	
Non-STEMI	R94.31	3.45	6032	
Age < 65 yr				
Standard definition				
STEMI	l21.0–3	2.15	3342	
Non-STEMI	l21.4x	0.62	4009	
ECG definition				
STEMI	R94.30	2.20	3266	
Non-STEMI	R94.31	0.57	3868	
Age ≥ 65 yr				
Standard definition				
STEMI	121.0–3	10.25	2371	
Non-STEMI	l21.4x	6.41	5426	
ECG definition				
STEMI	R94.30	10.07	2264	
Non-STEMI	R94.31	6.36	5202	

Note: ECG = electrocardiogram; MI = myocardial infarction; STEMI = ST-elevation MI.

*For ECG definition $n = 14\ 600\ (528\ patients had an unspecified abnormal ECG result [R94.38] or missing supplementary R94.3x code, and 20 patients had both the code R94.30 and the code R94.31).$

65 years or more in our study may have been due to less complete documentation by physicians for older patients with several conditions than for younger patients. It is also possible that coders spent less time on older patients' records when there were large amounts of information on conditions.

Limitations

Our study has limitations. It does not provide information on the validity of acute MI coding. Instead it examines only the validity of subcategorizing patients with acute MI as having STEMI or non-STEMI. Therefore, given that our study included only patients with acute MI, we were unable to provide the positive predictive value, negative predictive value, sensitivity or specificity associated with the acute MI diagnosis. Future studies based on a random sample of all patients admitted to several acute care hospitals in Alberta would be useful in obtaining these measures. The reference standard for this study was a sample of charts of patients discharged over a 3-month period from 3 hospitals in 1 Canadian city. To ensure the generalizability of our findings, it would be valuable to conduct this type of validation study for ICD-10 coding of acute MI subtypes in different regions of the country as well as internationally. Although our study was limited to a smaller study area, it covered a range of hospital settings in a large urban centre.

Conclusion

Among patients with acute MI, the standard ICD-10 definition to classify patients as having STEMI or non-STEMI was as accurate as the supplementary definition based on ECG codes. Our findings show that the acute MI subtype codes are valid and can be used in population-based outcome studies. These findings may be relevant for the development of later versions of ICD codes.

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Contributors: Robert Welsh, Hude Quan and Padma Kaul conceived the study; Alka Patel, Hude Quan and Padma Kaul wrote the manuscript; Alka Patel conducted the analysis and wrote the first draft of the paper; Jessica Deckert-Sookram and Ian Surdhar collected the data and reviewed the manuscript for content; and Wayne Tymchak, Sunil Sookram and Robert Welsh conducted the chart reviews and reviewed the manuscript for content. All of the authors approved the final version to be published and agreed to act as guarantors of the work.

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