

Current evidence and emerging methods regarding validation of the AHRQ QIs

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Outline and learning objectives

- No disclosures (except support from AHRQ)
- To describe key domains of validity for health care quality measures
- To illustrate these key domains of validity with examples based on the AHRQ Quality Indicators
 - To summarize methods and opportunities for AHRQ QI users who may wish to be involved in validation activities







- The degree to which a measure accurately represents the true state of the phenomenon of interest (i.e., "free of systematic error")
 Does this measure what it purports to
 - measure (quality)?





Validation – A conceptual framework

Face validity is the degree to which a measure "appears" to measure the phenomenon of interest

- Content validity is a related concept, focused on whether the content of a measure adequately samples all relevant domains of the concept of interest (coverage)
- Criterion (concurrent) validity is the degree to which a measure generates data that agree with data from a better ("gold standard") approach.
- Predictive validity is the degree to which a measure successfully predicts an outcome of interest.
- Construct (convergent) validity is the degree to which a measure correlates with other measures, based on a construct that is grounded in prior literature or a sound conceptual framework



Face validity: perspectives

Developers
Expert panels
Users and stakeholders





Face validity: NQF

- Health outcomes: "A rationale supports the relationship of the health outcome to at least one healthcare structure, process, intervention, or service."
 - Measure specifications must be "consistent with the evidence presented to support the focus of measurement..."
- Intermediate outcomes: "Quantity, quality, and consistency of... evidence that the measured intermediate clinical outcome leads to a desired health outcome."
- Processes or structures: "...evidence that the measured healthcare process leads to desired health outcomes in the target population with benefits that outweigh harms."
 - Patient experience: "Evidence that the measured aspects of care are those valued by patients and for which the patient is the best and/or only source (OR evidence that patient experience... is correlated with desired outce the



AHRQ expert panel process

- Intended to establish consensual (face) validity
- Modified RAND/UCLA Appropriateness Method
- Physicians of various specialties/subspecialties, nurses, other professionals (e.g., midwife, pharmacist)
- Potential PSIs were rated by 8 multispecialty panels; surgical PSIs were also rated by 3 surgical panels
- All panelists rated all assigned indicators (1-9) on:
 - Overall usefulness
 - Likelihood of being preventable
 - Likelihood of being due to medical error
 - Likelihood of being clearly charted in the medical record
 - Susceptibility to bias due to case mix





AHRQ expert panel process

- Pre-conference ratings and comments
- Individual ratings returned to panelists with distribution of ratings and other panelists' comments
- Telephone conference call(s) focusing on highvariability items and panelists' suggestions
- Suggestions adopted only by consensus
- Post-conference ratings and comments
- Exclude indicators rated "Unclear," "Unclear-," or "Unacceptable":
 - Median score <7, OR
 - At least 2 panelists rated the indicator in each of the extreme 3point ranges





Potential PSIs not adopted Only 18 "accepted" from original list of 48

"Experimental" PSIs

Aspiration pneumonia CABG after PTCA Decubitus ulcer in high risk patients In-hospital fractures possibly related to falls Intraoperative nerve compression injuries Malignant hyperthermia Postoperative acute myocardial infarction Postoperative iatrogenic complications cardiac system Postoperative introgenic complications – nervous system Reopening of surgical site Suture of laceration Obstetric wound complications- cesarean Obstetric wound complications- vaginal Other obstetric complications Postpartum urinary tract infection **Uterine rupture**

"Rejected" PSIs

Dosage complications latrogenic hypotension Intestinal infection due to C. difficile Postop iatrogenic complications digestive complications Postop iatrogenic complications respiratory complications Postop iatrogenic complications urinary complications Postop iatrogenic complications – vascular complications Postoperative pneumonia Unexpected LOS/Conditional LOS Obstetric thrombosis or embolism **Puerperal infection**



Face validity varies by proposed purpose

Application	Qua Improv	•			Compa Repor						y for rmance	
Level of Reporting	Large Pl Group	•	Area	Level	Payer	Level	Large Pl Group	•	Payer	Level	Large P Group	•
Panel	Delphi	NG	Delphi	NG	Delphi	NG	Delphi	NG	Delphi	NG	Delphi	NG
COPD and asthma (40 y+)	6*	7++	6*	6*	6*	6*	6*	7++	5*	5*	5.5*	7++
Asthma (<39 y)	7++	7++	6*	7++	6*	7++	5*	7++	6*	6*	6*	7++
Hypertension	5*	7++	6*	7++	5*	7++	4*	6*	5*	7+	4*	5.5*
Angina	6*	4*	5*	4.5*	5*	4*	5*	3-	4*	4*	4*	3-
CHF	7++	7++	6*	7++	6*	7++	7++	7++	6*	5*	6*	6*
Perforated appendix	4*	3-	5*	3-	5*	3 -	4*	3.5-	3.5-	2.5 -	4*	2-
Diabetes short term complications	7++	7++	6*	6*	6*	7++	6*	7++	5*	5*	6*	5*
Diabetes long-term complications	6*	7++	6*	7++	5*	6*	6*	6*	5*	4*	5*	4*
Lower extremity amputation in diabetics	6*	7++	7++	7++	6*	7++	5.5*	4*	5*	5*	5*	4*
Bacterial pneumonia	6*	6*	6*	5*	5*	5.5*	5*	6*	5*	5*	5*	6*
UTI	5*	6*	5*	6*	4*	5*	4*	4*	4*	3 -	4*	3-
Dehydration	5*	3 -	5*	5*	4*	3 -	3-	3 -	4*	3 -	3-	3 -

TABLE 3. Overall Usefulness Ratings (Median Panel Scores)

Numbers represent median usefulness ratings, as measured on a 9-point scale (1 = Highly discourage use; 9 = Highly recommend use).

Overall usefulness ratings: major concern-; some concern*; majority support+; full support++.

CHF indicates congestive heart failure; COPD, chronic obstructive pulmonary disease; NG, Nominal group; UTI, urinary tract infection.

Face validity varies by method

Table 2: Concordance between Delphi and Nominal Group (NG) on Combinations Rated

	Delphi Full Support	Delphi General Support	Delphi Some Concern	Delphi Major Concern
NG full support	8	2	21 (6)*	0
NG general support	0	0	$1 (1)^{*,\dagger}$	0
NG some concern	0	0	34	0
NG major concern	0	0	12 (5) *	3

*Numbers in parentheses are the number of instances in that cell where |Median (Delphi) - Median (NG)| > 1. The median difference between groups was < 1 in all other combinations. [†]The support level can only deemed "General Support with Some Concern" if statistical disagreement exists within the panel.



Is the purported "gold standard" really a gold standard? How do we know? Administrative data perspective versus registry data perspective Coding perspective versus clinical perspective (whose truth) Validity can change... dramatically Finding false positives is easy, but what about false negatives?





Criterion validity: NQF

"Empirical evidence of validity of BOTH data elements AND measure score within acceptable norms; AND Identified threats to validity (lack of risk) adjustment/stratification, multiple data types/methods, systematic missing or "incorrect" data) are empirically assessed and adequately addressed so that results are not biased."

Could be satisifed by either criterion or construct testing

Criterion validity: Present on admission coding vs. chart review

SERVICES

	Percentage not POA (%)				
PDI	NACHRI	Mich	СА	NY	Мауо
PSI 1: Complications of Anesthesia		100	100	100	94
PSI 3: Decubitus Ulcer	60	42	11	14	18
PSI 5: Foreign Body Left During Proc	80	80	64	76	54
PSI 6: latrogenic Pneumothorax	89	100	73	65	78
PSI 7: Infection Due To Medical Care	57	36	65	65	60
PSI 8: Postop Hip Fracture		0	21	26	22
PSI 9: Postop Hemorrhage or Hematoma	97	100	79	71	87
PSI 10: Postop Physiologic or Metabolic		91	77	64	46
PSI 11: Postop Respiratory Failure	83	100	94	93	74
PSI 12: Postop DVT or PE		67	46	43	40
PSI 13: Postoperative Sepsis	60	60	73	70	76
PSI 14: Postop Wound Dehiscence	90				100
PSI 15: Accidental Puncture/Laceration	93	84	87	87 🚄	85
PSI 16: Transfusion Reaction	71	N/A	58	78	IKR



Criterion validity of PSIs linked to NSQuIP

Romano PS, et al. *HSR* 2009; 44(1):182 Cima RR, et al. *Surg* 2011; 150:943 Koch CG, et al. *J Am Coll Surg* 2012

	S	ensitivity	PPV		LR+	
Indicator	V2	V3/V4	V2	V3/V4		
Postoperative sepsis		37% (VA) 5-10% (C/M)		45% (VA) 19-44% (C/M)		131
Postoperative thromboembolism	56%	58-72% (C/M)	22%	42-53% (C/M)	65	
Postoperative respiratory failure		63% 21-22% (C/M)		68% 42-61% (C/M)		147
Postop physiologic/ metabolic derangement		48% 12% (M)		63% 89% (M)		744
Postop abdominopelvic wound dehiscence	29%	22% (M)	72%	47% (M)	160	

VA=Veterans Affairs; C/M=Cleveland Clinic/Mayo Clinic Rochester Sensitivity = TP/(TP+FN) – are all the real cases captured? PPV = TP/(TP+FP) – are all the flagged cases real? LR = Sensitivity/(100-Specificity) – how many times more likely is the event?





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Indicator	AHRQ	NSQIP Registry
Postoperative sepsis	Sepsis or septicemia diagnosed by MD (pts with POA infection excluded)	SIRS w "definitive" evidence of any infection
Postoperative thromboembolism	DVT or PE diagnosed by MD (implied treatment)	DVT or PE treated by MD (PE requires imaging)
Postoperative respiratory failure	Diagnosed by MD or unplanned reintub >0 days or postop vent >1 day	Postop vent >48 hrs or unplanned reintub (any)
Postop physiologic/ metabolic derangement	Diagnosed by MD and new onset of dialysis (or DKA or hyperosmolar)	New onset of dialysis or ultrafiltration
Postop abdominopelvic wound dehiscence	Surgery to repair postop wound disruption	Postop wound separation w fascial disruption





PPV of the PSIs based on chart review by nurse abstractors

Indicator	VA %PPV (95%Cl)	AHRQ %PPV (95%CI)	UHC %PPV (95%CI)
Pressure Ulcer	30 (22-40)	_	32 (30-35)
Foreign Body Left In	46 (36-55)		
latrogenic Pneumothorax	73 (64-81)	78 (73-82)	
CVC-related Bloodstream Infection	38 (29-47)	61 (51-71)	
Postop Hip Fracture	28 (15-43)		
Postop Hemorrhage/Hematoma	75 (66-83)	78 (62-95)	
Postop Phys/Met Derangement	63 (54-72)	85 (78-92)	_
Postop Respiratory Failure	67 (57-76)	_	83 (77-89)
Postop PE or DVT	43 (34-53)	47 (42-52)	44 (37-51)
Postop Sepsis	53 (42-64)	41 (28-54)	
Postop Wound Dehiscence	87 (79-92)	_	_
Accidental Puncture or Laceration	85 (77-91)	91 (86-94)	

Rosen, Med Care, 2012; Sadeghi, Am J Med Qual, 2010; Zrelak, J Healthc Qual, 2011; HRR White, Med Care, 2009; Utter, Ann Surg, 2009; Utter, J Am Coll Surg, 2010



Limitations of chart abstraction for criterion validation

Information needed to verify complication may not be available via chart review Complication was not properly evaluated or described by physicians; vicarious process Absence of evidence vs. evidence of absence Time constraints limit abstractor's ability to assess some aspects of care (e.g., urinary catheter), may lead to premature termination Much cheaper to look for FPs than FNs Inter-hospital variation in physician documentation and nurse abstraction Volunteer samples (except VA)



Comparing coding vs. clinical perspective for Postop DVT/PE

UHC Cohort (n=450)	Coding	Clinical
Sensitivity	80% (46-100%)	100%
Specificity	99.5% (99.3-99.6%)	98.6% (98.6-99.2%)
Positive Predictive Value	72% (67-79%)	44% (36-52%)
Negative Predictive Value	99.6% (98.9-100%)	100%
VA Cohort (n=112)		
Positive Predictive Value		43% (34-53%)
AHRQ Cohort (n=121)		
Positive Predictive Value	84% (72-95%)	48% (42-67%)

University HealthSystem Consortium cohort includes 505 flagged, randomly sampled surgical cases from 33 volunteer hospitals in 21 states; 450 cases were fully abstracted and submitted to UHC. 453.4 Acute venous embolism and thrombosis of deep vessels of lower extremity

453.40 Acute venous embolism and thrombosis of unspecified deep vessels of lower extremity Deep vein thrombosis NOS

453.41 Acute venous embolism and thrombosis of deep vessels of proximal lower extremity Femoral, Iliac, Popliteal, Thigh, Upper leg NOS

453.42 Acute venous embolism and thrombosis of deep vessels of distal lower extremity

Calf, Lower leg NOS, Peroneal, Tibial

453.5 Chronic venous embolism and thrombosis of deep vessels of lower extremity Excludes: personal history of venous thrombosis and embolism (V12.51)

453.50 Chronic venous embolism and thrombosis of unspecified deep vessels of lower extremity

453.51 Chronic venous embolism and thrombosis of deep vessels of proximal lower extremity

453.52 Chronic venous embolism and thrombosis of deep vessels of distal lower extremity

453.6 Venous embolism and thrombosis of superficial vessels of lower extremity

453.7 Chronic venous embolism and thrombosis of other specified vessels

Excludes: personal history of venous thrombosis and embolism (V12.51)

453.71 Chronic venous embolism and thrombosis of superficial veins of upper extremity

453.72 Chronic venous embolism and thrombosis of deep veins of upper extremity

453.73 Chronic venous embolism and thrombosis of upper extremity, unspecified

453.74 Chronic venous embolism and thrombosis of axillary veins

453.75 Chronic venous embolism and thrombosis of subclavian veins

453.76 Chronic venous embolism and thrombosis of internal jugular veins

453.77 Chronic venous embolism and thrombosis of other thoracic veins

453.79 Chronic venous embolism and thrombosis of other specified veins

453.8 Acute venous embolism and thrombosis of other specified veins

Excludes: cerebral, coronary, intracranial sinus, nonpyogenic, mesenteric, portal, precerebral, pulmonary

453.81 Acute venous embolism and thrombosis of superficial veins of upper extremity

453.82 Acute venous embolism and thrombosis of deep veins of upper extremity

453.83 Acute venous embolism and thrombosis of upper extremity, unspecified

453.84 Acute venous embolism and thrombosis of axillary veins

453.85 Acute venous embolism and thrombosis of subclavian veins

453.86 Acute venous embolism and thrombosis of internal jugular veins

453.87 Acute venous embolism and thrombosis of other thoracic veins

453.89 Acute venous embolism and thrombosis of other specified veins

453.9 Of unspecified site (embolism of vein, thrombosis (vein))



Methods to reassess criterion validity of PSI 12

- Two parallel studies were conducted to update previous PPV estimates for PSI 12 and to identify actionable opportunities to improve care:
 - 7 volunteer hospitals recruited through AHRQ QI listserve, including flagged cases only
 - 15 academic health systems recruited through UHC, including both flagged and unflagged cases with TKA surgery
- AHRQ PSI 12 Version 4.1 software was applied to eligible cases from participating hospitals, using "present on admission" (POA) flags.
 - Hospital's own data (AHRQ) or Clinical Database (UHC)





Methods to reassess criterion validity of PSI 12

- Flagged cases were reviewed by trained QI nurses at each hospital, using detailed chart abstraction tool and guidelines.
- Detailed review of discrepant cases to identify possible reasons for the discrepancy:
 - Present on admission (note one hospital did not apply denominator exclusions)
 - Location of thrombosis (upper extremity and superficial thromboses are clinical FPs)
 - Chronic vs. acute embolism (based on radiographic criteria)
- Records from volunteer hospitals in AHRQ study were sampled in sequential reverse order from 6/30/2010 back to 10/1/2009, up to N=30





Findings: 7 volunteer hospitals

From a total of 171 audited charts, 15 cases were excluded post hoc (because hospital did not properly apply POA)
 30 cases were False Positive:

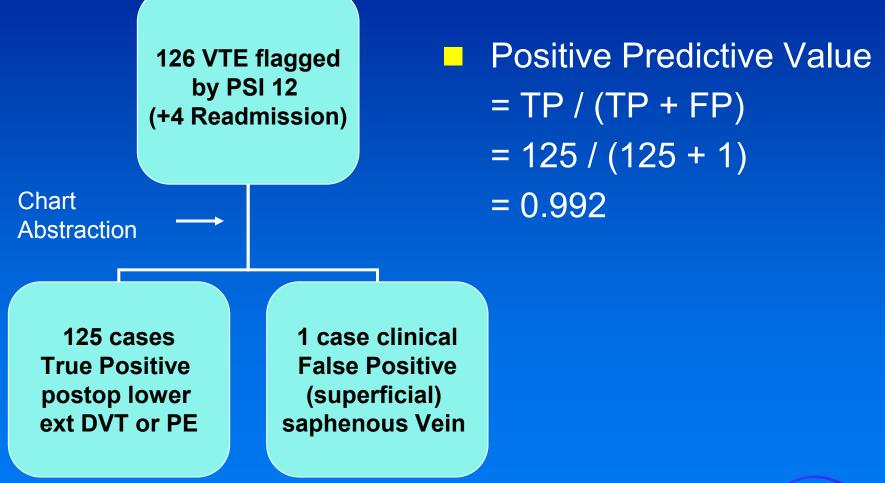
 15 cases were POA
 8 cases with upper extremity VT

- 1 case with SVC (central VT)
- 3 cases with superficial VT
- 3 cases were chronic
- Overall PPV = 81%





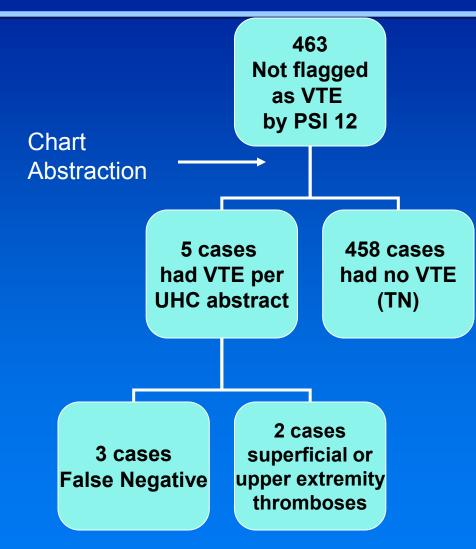
Findings: AMCs with TKA patients PSI-12 flagged cases







Findings: AMCs with TKA patients PSI-12 unflagged controls



Negative Predictive Value
 = TN / (FN + TN)
 = 458 / (458+3) = 0.993



Estimating sensitivity: Looking for a needle in a haystack

- Retrospective cross-sectional study
 - 27 hospitals from 11 states
 - 2006-2009 PSI-negative hospitalizations
- Stratified sample
 - By hospital, risk of being falsely negative
 - Oversampled cases at risk
- Medical records abstracted locally
 - Trained staff, standard instrument

Analysis with survey statistical methods

- "Verification-biased sampling" approach, used model-based weights in analysis
- Incorporated previous estimates of PPV

Suspicious records: Could they be false negatives?

Indicator

OF HEALTH & AL



X

X

X

Foreign Body Left In	X		X
latrogenic Pneumothorax	X		X
Central Venous Catheter Infection	X	X	
Postop Hemorrhage/Hematoma	X		
Postop Phys/Met Derangement	X	X	X
Postop Wound Dehiscence	X		X
Accidental Puncture or Laceration	X		X



Sampling scheme for estimating sensitivity of selected PSIs

Indicator	Risk level	Sampling frame	Abstracted records
			n
	%		n %
Foreign Body Left In		Low	
	664,956	99.91	295
	0.04		
		High	589
	0.09	5	21
	3.56		
latrogenic Pneumothorax		Low	
	535,648	99.92	269
	0.01		
		High	425
	0.08		11
	2.59		
Central Venous Catheter Infection	Low		453,138
	99.29		197
	0.04		
	<u> </u>	High	3,250

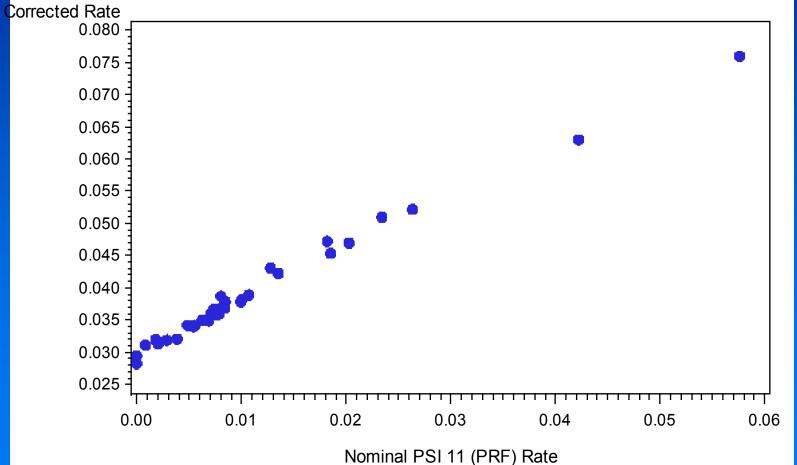
Estimated sensitivity of selected AHRQ PSIs

Indicator	Abstracted records	False negative records	Sensitivity
	N % (95% CI)	All	High risk
Foreign Body Left In	316 100 (0-100)	0	_
latrogenic Pneumothorax	279 25 (8-58)	9	7
Central Venous Catheter Infection	223 11 (1-60)	3	2
Postop Hemorrhage/Hematoma	281 49 (26-72)	32	30
Postop Phys/Met Derangement	231	6	6



Simulating how false negatives and false positives would affect hospital-level rates

Corrected vs. Nominal PSI Rate

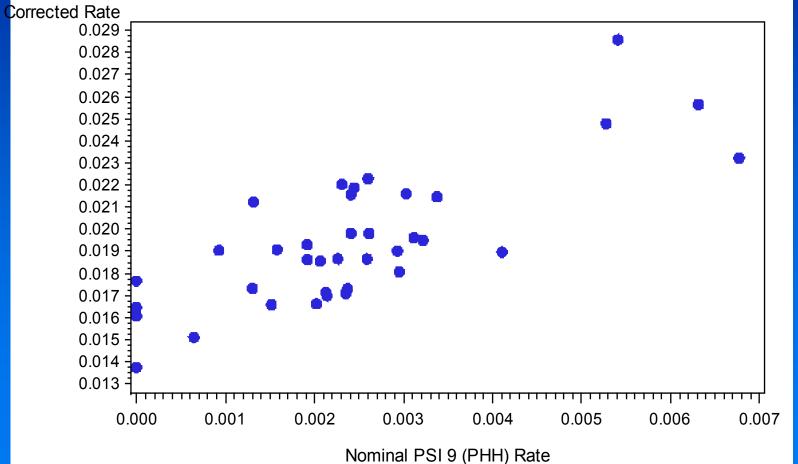






Simulating how false negatives and false positives would affect hospital-level rates

Corrected vs. Nominal PSI Rate





Improving the specification for Postoperative Hemorrhage/Hematoma

Indicator	Sensitivity % (95% CI)	PPV % (95% CI)
Current PSI 9 definition	49 (26-72)	78 (59-90)
PSI 9 definition + codes for treatment of PHH	71 (48-87)	76 (58-88)
PSI 9 definition + codes for treatment of PHH + codes for eval of bleeding	87 (69-95)	77 (61-87)
PSI 9 definition - requirement for procedure code	94 (80-98)	48 (30-66) AHR R



Predictive validity: perspectives

Predicting what?

- Mortality
- Length of stay, charges
- Readmission
- Time window for prediction?Implied gold standard





Impact of preventing a PSI on mortality, LOS, charges NIS 2000 analysis by Zhan & Miller, JAMA 2003;290:1868-74 VA PTF 2001 analysis by Rivard et al., Med Care Res Rev; 65(1):67-87

Indicator	Δ Mort (%)	ΔLOS (d)	Δ Charge (\$)
Postoperative septicemia	21.9-30.2	10.9-18.8	31264-57700
Selected infections due to medical care	2.7-4.3	9.5-9.6	13816-38700
Postop abd/pelvic wound dehiscence	9.6-11.7	9.4-11.7	18905-40300
Postoperative respiratory failure	21.8-24.2	8.6-9.1	39745-53500
Postoperative physiologic or metabolic derangement	19.8	8.9	54,800
Postoperative thromboembolism	6.1-6.6	5.4-5.5	7205-21700
Postoperative hip fracture	4.5	5.2	13,400
latrogenic pneumothorax	2.7-7.0	3.9-4.4	5633-17300
Decubitus ulcer	6.8-7.2	4.0-5.2	6713-10800
Postoperative hemorrhage/hematoma	3.0-5.1	3.9-3.9	7863-21400
Accidental puncture or laceration	2.2-3.2	1.3-1.4	3 <u>3</u> 59-8300

Excess mortality, LOS, and charges computed from mean values for PSI cases and matched controls.



PSIs also appear to predict later readmissions

7 state SID 2004 analysis by Friedman et al., Med Care 2009;47(5):583-90

Indicator	Inpatient death	Readmit 1 month	Readmit 3 months
Postoperative septicemia	4.70	0.99	1.26
Selected infections due to medical care	1.23	1.00	1.29
Postop abd/pelvic wound dehiscence	1.57	1.24	1.56
Postoperative respiratory failure	13.23	1.03	1.14
Postoperative physiologic or metabolic derangement	3.73	1.09	1.30
Postoperative thromboembolism	1.35	1.25	1.28
latrogenic pneumothorax	2.47	1.02	1.20
Postoperative hemorrhage/hematoma	1.03	1.10	1.18
Accidental puncture or laceration	1.52	1.25	1.16

Risk ratios adjusted as in AHRQ model, but also for payer group and APR DRG SOI and ROM levels Statistically significant risk ratios are highlighted in yellow.

Predictive validity must be interpreted in context with reported prevalence (England)

	Rates/1000 admissions		Matched cases: excess mortality	
Indicator	England (2005/6)	US (2000)	England (2005/6)	US (2000)
Decubitus ulcer	7.17	21.51	13.4	7.2
latrogenic pneumothorax	0.12	0.67	10.6	7.0
Infections due to medical care	1.06	1.99	5.7	4.3
Postoperative hip fracture	0.08	0.77	18.2	4.5
Postoperative sepsis	2.66	11.25	27.1	21.9
Obstetric trauma:				
Vaginal with instrument	60.34	224.21	*	0.0
Vaginal without instrument	29.39	86.61	0.01	0.0

Data reported by:

UK - Raleigh VS, Cooper J, Bremner SA, Scobie S. BMJ 2008, 337:a1702

US - Zhan C, Miller MR. JAMA 2003;290:1868-74.

Reported data do NOT suggest predictive validity for Birth Trauma (PSI 17), Obstetric trauma (PSI 18/19), Transfusion Reaction (PSI 16), and Complications of Anesthesia (EXP 01).



Is the construct sound?
Correlation with process measures

Do we have the right process measures?

Correlation with structural measures

Do we have the right structural measures?

Correlation with other outcome domains



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Case control study of PSI 12

Cases (up to 20):

- Uni-TKA or Bilat-TKA
- Oct 2008 to Mar 2010
- >40 yrs
- PSI-12 code for VTE within 90 days

Controls (up to 40):

- Uni-TKA or Bilat-TKA
- Oct 2008 to Mar 2010
- >40 yrs
- NO PSI-12 code for VTE within 90 days

No TKA or THA within 90 days prior
No VTE as principal ICD-9-CM diagnosis
No VTE as POA
No pregnancy, childbirth, or puerperium



Analysis of case control data

- Classified FDA-approved pharmacologic prophylaxis as receipt of the recommended dose at the recommended starting time (per package insert) before or after surgery, and continued until at least the day of discharge
- Patients who were diagnosed with VTE on the day of surgery or the day after surgery were not included in the case control analysis
- Other risk factors assessed included age, obesity (BMI), type of TKA, race/ethnicity, date of ambulation, personal or family history of VTE, and comorbid conditions
- Analysis adjusted for conditional stratified sampling of controls without VTE





Multivariable analyses of process factors

Multivariate adjusted odds ratios and 95% confidence intervals
 Outcome: Any VTE event diagnosed Day 2 of surgery or later

Excluded one hospital that screened TKA patients routinely for VTE

Predictive Factor	Odds Ratio (95% CI)	P value
Age	1.02 (0.99 – 1.05)	0.20
Gender (ref: male)	1.7 (0.9 – 2.9)	0.90
Ambulation (ref: no ambulation)		
 Taking steps day 1 or 2 	0.3 (0.1 – 0.9)	0.005
 Taking steps after day 2 	0.7 (0.2 – 2.1)	0.56
Type of TKA (ref: unilateral TKA)		
 Bilateral TKR 	4.2 (1.9 – 9.1)	0.004
Recommended pharmacologic prophylaxis (ref: only mechanical)	0.5 (0.3 – 0.8)	0.01
BMI ≥ 35 (ref: BMI < 35)	0.9 (0.5 – 1.6)	0.66



Implications of recent validation studies

- Generalizability to nonparticipating hospitals remains unclear
- Limited supervision/oversight of local abstractors who may have COI
- ICD-9-CM changes and associated coder training can substantially improve criterion validity of PSIs, in the right circumstances

Actionable opportunities to improve care may persist despite 100% compliance with TJC process measures, but often we don't know what processes to use for construct validation
Estimating sensitivity or FN rate is still hard and the sensitivity or FN rate is still hard and the sensitivity of the sens



Future directions in validation

- Encourage ongoing criterion validation work using previously developed tools or new learning collaboratives
- Continue to pursue opportunities to link registry or EHR data, especially laboratory, imaging, vital signs, etc.

Encourage case control and intervention studies when we know what processes to measure or change

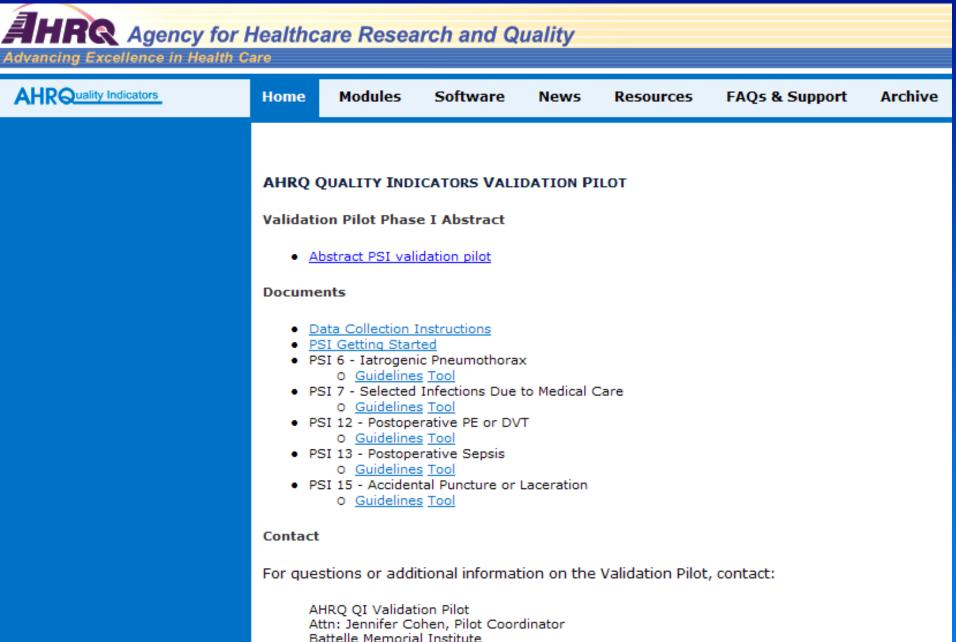




Improving indicator performance

Change indicator specifications to capture false negatives or exclude false positives Change ICD-9-CM codes, coding guidelines, or Coding Clinic advice Promote universal adoption of POA flag Work with hospitals to improve and standardize clinical documentation and coding Everything will change in ICD-10-CM/PCS (10/1/2014)

http://qualityindicators.ahrq.gov/ValidationPilot.aspx



Acknowledgments and references

- Participating hospitals and systems
- AHRQ Quality Indicators project team: Mamatha Pancholi, John Bott
- HCUP data partners
- Battelle validation support: Jeffrey Geppert, Jaime Liesmann
- UC Davis: Pat Zrelak, Banafsheh Sadeghi, Richard White, Garth Utter
- White RH, et al. How valid is the ICD-9-CM based AHRQ Patient Safety Indicator for postoperative venous thromboembolism? Med Care 2009; 47(12):1237-43.
- White RH, et al. Evaluation of the predictive value of ICD-9-CM coded adminstrative data for venous thromboembolism in the United States. Thromb Res 2010; 126(1):61-7.
- Sadeghi B, et al. Mechanical and suboptimal pharmacologic prophylaxis and delayed mobilization but not morbid obesity are associated with venous thromboembolism after total knee arthroplasty: A case-control study. J Hosp Med 2012; in press.

Go to AHRQ QI website for full list: http://qualityindicators.ahrq.gov/Resources/Publications.as





