

Leveraging Clinical Registries to Drive Quality and Value

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Disclosures

- Research
 - Boston Scientific
 - Medtronic
 - Janssen
- Speaker's Bureau
 - Bristol Myers Squibb
 - Pfizer

How do we define Quality and Value in Medicine?

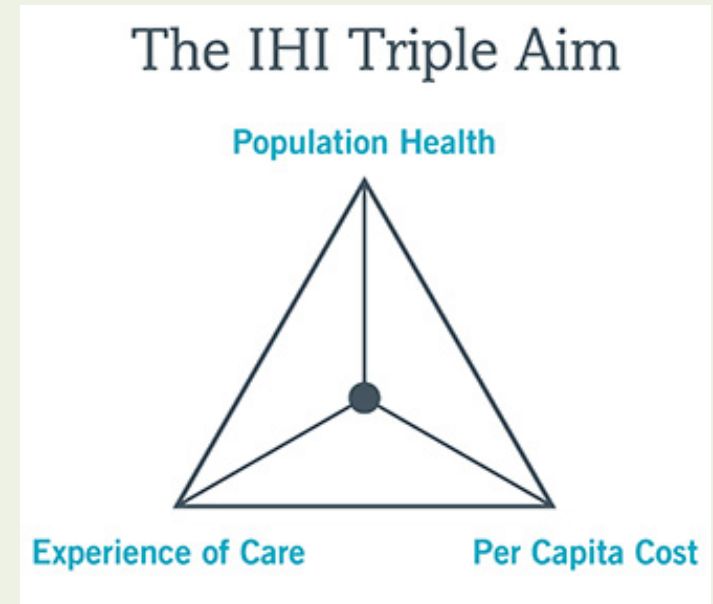
- Value can be defined as “...health outcomes achieved per dollar spent.”¹
- “Value — neither an abstract ideal nor a code word for cost reduction — should define the framework for performance improvement in health care. Rigorous, disciplined measurement and improvement of value is the best way to drive system progress. Yet value in health care remains largely unmeasured and misunderstood.”²
- In simple terms, $\text{value} = \text{quality} / \text{cost}$

¹Porter ME, Teisberg EO. Redefining health care: creating value-based competition on results. Boston: Harvard Business School Press, 2006.

²Porter ME. What is value in health care? N Engl J Med 2010; 363:2477-2481. DOI:10.1056/NEJMp1011024

The Triple Aim

- Improving the patient experience of care (including quality and satisfaction)
- Improving health of populations
- Reducing the per capita cost of health



www.ihl.org


An Example

ED Activation of Cath Lab


- Door-to-balloon time ↓ from 113 to 75 minutes
- Transfer time ↓ from 147 to 85 minutes
- Infarct size ↓ (creatinine kinase)
- Hospital stays ↓ by 2-3 days
- Cost ↓ by over 30 percent



Improved Satisfaction



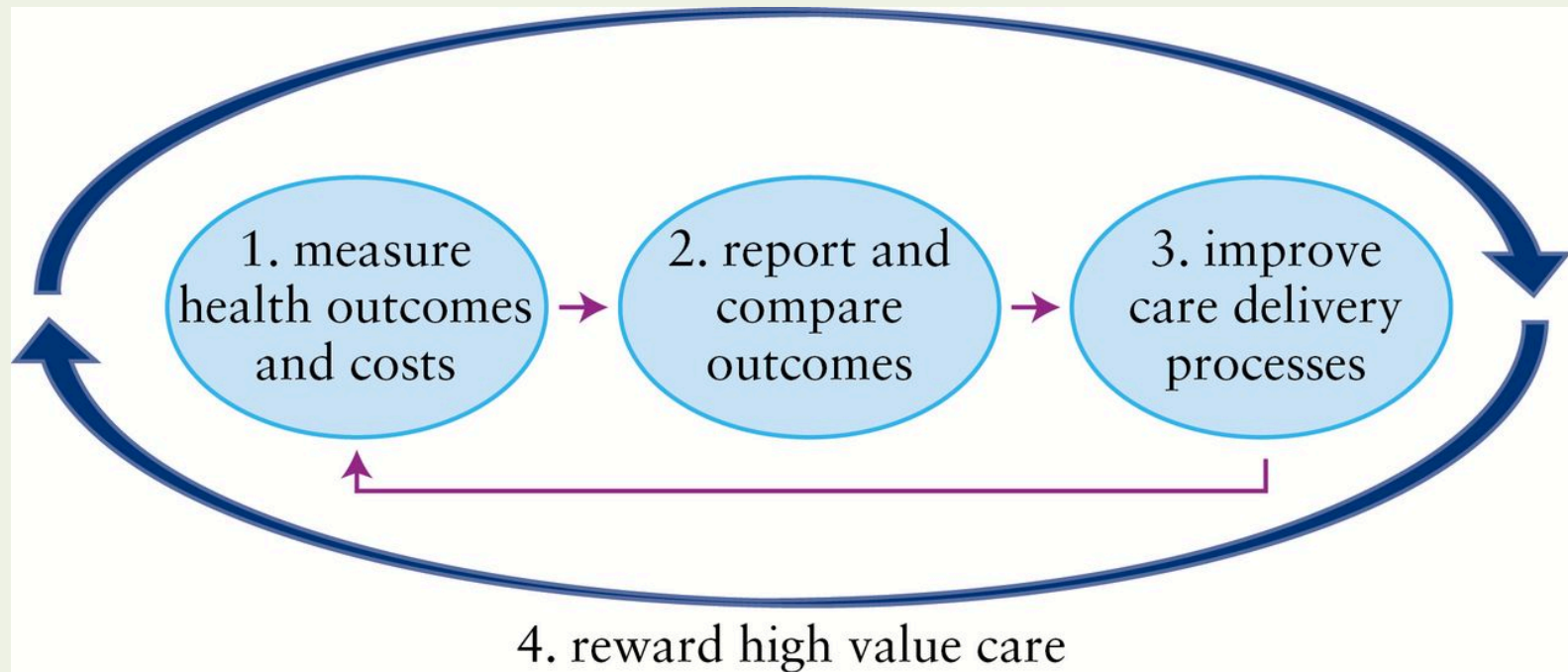
Improved Health



Reduced Cost

U. M. Khot et. Al. *Circulation*. 2007; 116

Steps to Improve Quality and Value





That was then...

Launched 1997

1 registry

Focused on quality patient care

This is now...

More than 2,500 hospitals and 1000 practices

Health plans and government regulator adoption

Industry uses for market research, clinical research, and to support best practice treatments

FDA uses NCDR data for post market assessment

This is our future...

One holistic registry

International expansion

Platform for clinical trials and CER

More post market assessment studies

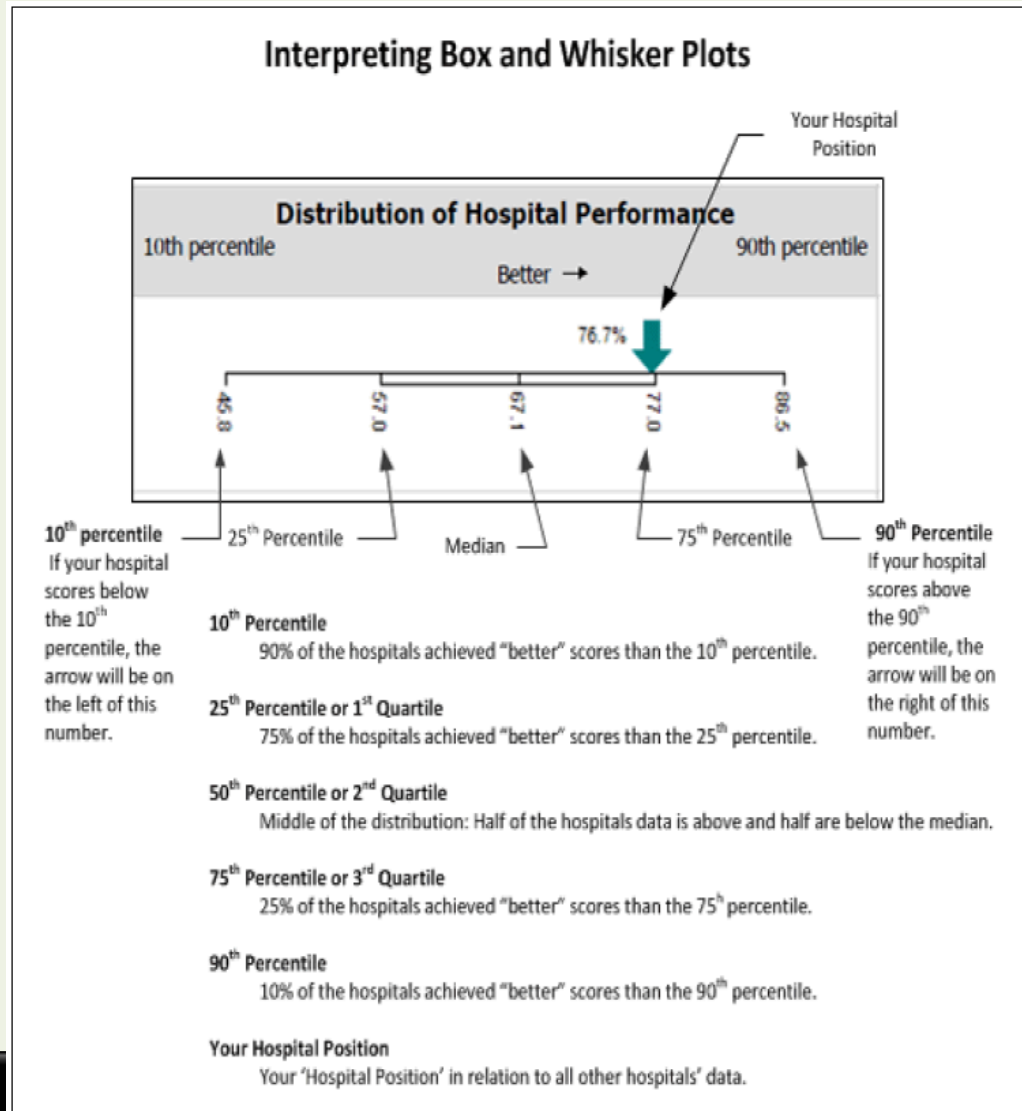
Implement physician reports to support MOC and MOL

EHR Integration

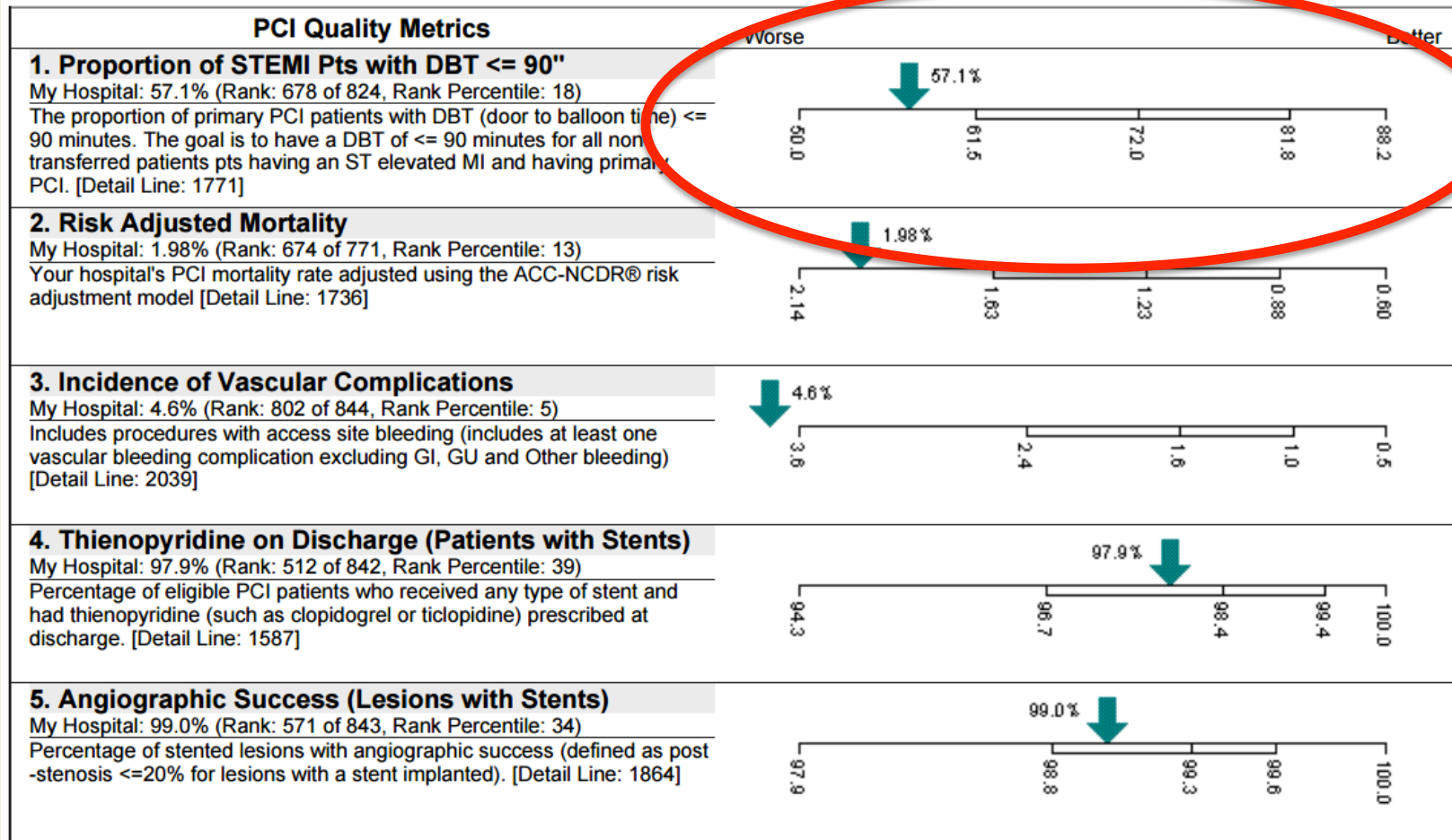
NCDR Registries

- Hospital Registries (for in-patients)
 - ACTION Registry-GWTG
 - High risk STEMI/NSTEMI patients
 - Afib Ablation (coming soon)
 - CathPCI Registry
 - Diagnostic catheterization and PCI
 - ICD Registry
 - IMPACT Registry
 - Pediatric and adult congenital heart disease
 - LAAO Registry (coming soon)
 - LAA occlusion procedures
 - PVI Registry
 - -Peripheral arterial catheterization procedures including carotid stenting
 - STS/ACC TVT Registry
 - -Transcatheter valve procedures
- Out-patient Registries
 - Diabetes Collaborative Registry
 - PINNACLE
 - CAD, hypertension, heart failure and atrial fibrillation

The Box and Whisker Plots



The History of D2B at IU Health Bloomington



How Did We Improve Our Quality?

- The team came together to review and compare the data, and improve the process
- Physicians can rarely do this on their own
 - EMS
 - Transmission of ECGs to the ED from the field
 - ED physicians and staff
 - Empowered to activate STEMI alert
 - Interventional cardiologists
 - Primary caregivers to all STEMI patients
 - Cath lab staff
 - Received feedback on all STEMI
 - Quality review specialists
 - Reviewed data at monthly meetings
 - Coders
 - -Educated on correct coding

Picture of AMI Mtg Attendees



Indiana University Health

STEMI Heroes

Date	9/15/2015
ED Arrival Time	07:14
EKG Time	STEMI 9/15/15
STEMI Alert	Jacobs Young, Elizabeth M [EJacobs3@IUHealth.org]
Cardiac Cath Lab Arrival	Sent: Tuesday, September 15, 2015 2:37 PM
MD Arrival	To: Brown, Jessica M [JBrown28@IUHealth.org]; Bryan, Jessica Lynne [jbryan@IUHealth.org]; Caudill, Gwendolyn M [gcaudill@IUHealth.org]; Elliott-Felton, Julia F [jelliottfelt@IUHealth.org]; Englert, Danielle [DEnglert@IUHealth.org]; Hatcher, Tara [THatcher@IUHealth.org]; Hickok, 'Benny' Richard E [RHickok@IUHealth.org]; Hobson, Penny L [PHobson@IUHealth.org]; Hoy, Jill C [JHoy2@IUHealth.org]; Jacobs Young, Elizabeth M [EJacobs3@IUHealth.org]; Lemons, Casey L [CLemons@bloomingtonhospital.org]; Monnier, Megan E [MMonnier@bloomingtonhospital.org]; Spicer, Kathryn R [kspicer1@IUHealth.org]; Sturgeon, Betty K [BSturgeon1@IUHealth.org]; Uland, Kari B [KUland@IUHealth.org]; Walls, Jordann Leigh [JLWalls@bloomingtonhospital.org]; Wieligman, Patti [PWieligman@IUHealth.org]; Adams DO, Robert T [RADams1@IUHealth.org]; Amber Chinn [axevas@stvincent.org]; Balaguras, Jean M [JBalaguras@IUHealth.org]; Bellamy, Linda [LBellamy@IUHealth.org]; Bill Turpen RN [wturpen@monroehospital.com]; Bob Page [lead2noclue@mac.com]; Breeden, Pamela [pbreeden@IUHealth.org]; Davis, Brenda R [bdavis14@IUHealth.org]; Brian Bomar [bbomar@careambulance.com]; Brian McCrate clinical Pharm. [bmccrate@iuhealth.org]; C. Jessee [cjessee@iuhealth.org]; Carter, Susan [SCarter@IUHealth.org]; Clark, Charlene [CClark@IUHealth.org]; Cook, Holly C [hcook@IUHealth.org]; Cris Lunsford [clunsford@ocems.net]; Crouch, Jason [JCrouch4@IUHealth.org]; Beachy DO, David [dave@clearstreams.net]; David Doane [david.doane@co.greene.in.us]; David Polley [dpolley@crh.org]; Dina Wood RN [dwood@monroehospital.com]; Don Smith Care Ambulance [donsmith@careambulance.com]; Dr Raza [steph@havcmd.com]; Dr. Hall [Dhall14@iuhealth.org]; Eisenhut Jr MD, Richard J [REisenhut@bloomingtonhospital.org]; Etter, Jason [JETter1@IUHealth.org]; Evans, J. Chris [JEvans15@IUHealth.org]; Fields, Derek B. [DFields5@IUHealth.org]; Fons, Mark [mark.fons@gmail.com]; Geyer, Susan L [SGeyer@bloomingtonhospital.org]; Grupenhoff, Marcy L [MGrupenhoff@IUHealth.org]; Hamm, Krystal M [KHamm2@IUHealth.org]; Hanania, Raja [RHanania1@IUHealth.org]; Harris, Michael D [MHarris11@IUHealth.org]; Hawkins, Sara L [SHawkins2@IUHealth.org]; Helms, Tricia [THelms@IUHealth.org]; Horn, LeAnne [MHorn@IUHealth.org]; Jeanette Hammerstein, MD [jhammerstein@monroehospital.com]; Jim Frasure [jim@frasurefam.com]; Hill, Reva J [RHill5@IUHealth.org]; John Gott [gottjohn@air-evac.com]; Jones, Maudie J. [MJones2@IUHealth.org]; Jones, Venus D [vjones5@IUHealth.org]; Joshua Davis [davis.josh@seals-ambulance.com]; Kay, June E. [JKay1@IUHealth.org]; Kelly Mazzier-Devitt [Kelly.Mazzier-Devitt@TheMedCo.com]; Fahr, Kenneth J [KFahr@bloomingtonhospital.org]; Kern, Kammi D [KKern3@IUHealth.org]; Kim Adams (Monroe Hosp) [kadams@monroehospital.com]; Kinder, Connie [CKinder@IUHealth.org]; King, Jason M [jking@IUHealth.org]; Olsen, Kristen P [kolsen@IUHealth.org]; Kurtis Cummings BHAS EMS [kcummings1@iuhealth.org]; Ladonna Stroud RN Paoli ED [lstroud3@iuhealth.org]; Lea Ann Camp [lea.camp@mygcgh.org]; Max Peters Lawrence Co, Perry Township VFD [petersfm47462@yahoo.com]; Mullis, Kelly D [KMullis@IUHealth.org]; Owen, Michael P [MOWen2@IUHealth.org]; Pam Underwood Seals Ambulance [underwood.pamela@seals-ambulance.com]; Paula Armas [parmas@careambulance.com]; Pavich, Emily M [epavich@IUHealth.org]; Reynolds, Chad [creynolds1@IUHealth.org]; Ridge, Michele A. [MRidge@IUHealth.org]; Ryan Bertram [bertram.ryan@seals-ambulance.com]; Sarah Tieman MD [satieman@iu.edu]; Sargent, J. Scott [JSargent1@IUHealth.org]; Sayers, Jennifer A [JSayers@IUHealth.org]; Schwartz, Patrick [pbschwar@iupui.edu]; Slone, Teresa J [TSlone1@IUHealth.org]; Sperring MD, Sally [SSperring@bloomingtonhospital.org]; Stackhouse, Rebecca J [RStackhouse@IUHealth.org]; Stacy Fiscus [stacy.fiscus@air-evac.com]; Steve Waldrige [swaldrige1@iuhealth.org]; Tim Hale [tim.hale@mygcgh.org]; Travis Keating [tkeating@iuhealth.org]; Troy Galtner [troy.galtner@mygcgh.org]; Vickie Nelson RN Paoli ED [vnelson2@iuhealth.org]; Wallace, Sandra J [SWallace1@IUHealth.org]; Watters MD, Andrew Kyle [awatters1@IUHealth.org]; Watts, Wylene S [WWatts1@IUHealth.org]; Webb, Nichole [nwebb2@iuhealth.org]; Wilcher, Amy [awilcher1@IUHealth.org]; Williams, Lindsey R [LWilliams2@IUHealth.org]; Zachary Shaw [zshaw1@iuhealth.org]; Blemker, M.D. David; Shenoy, M.D. Mohan; Ferguson, M.D. Andrew; Fix, M.D. James; Heumann, M.D. Gregory; Frey, P.A. Julie; Petersen, M.D. Jennifer; Rink, M.D. Lawrence; Strobel, M.D. John; Sutliff, M.D. Gregory; Williams, M.D. Eric; Zamani, Atiq; Zawacki, M.D. Kevin

Pre



RCA pre

Attachments: [0915 RL 407236173 Private ~1.pdf \(563 KB\)](#) [Open as Web Page]

This is a 86 yr old gentleman that began experiencing chest pain at 06:00. He arrived to IUH Bloomington ED at 07:14 by private vehicle. EKG was done at 07:19. STEMI page went out at 07:25.

Patient arrived to the cath lab at 07:32. Right femoral access was obtained at 07:39. Angiography revealed acutely occluded right coronary artery. Vessel was ballooned at 07:45. Vessel was later stented. Patient was transferred to CWR in stable condition at 08:40.

Elizabeth (Beth) Jacobs Young RN
Cardiovascular Program Coordinator
IU Health Bloomington Hospital
P 812-353-9080 f 812-353-3559
EJacobs3@iuhealth.org

EKG Dept



Section II: Quality Metrics – to support self assessment and quality improvement at the provider, hospital, and/or health care system level.

PCI Process Metrics

2 Proportion of elective PCIs with prior positive stress or imaging study

My Hospital	US Hospitals 50th Pctl	US Hospitals 90th Pctl
73.68%	66.86%	88.94%

Proportion of elective PCI procedures (excluding patients with ACS) with an antecedent stress or imaging study with a positive result (suggestive of ischemia) or with a fractional flow reserve value of ≤ 0.8 during the PCI procedure [Detail Line:1513]

3 Median time to immediate PCI for STEMI patients (in minutes)

My Hospital	US Hospitals 50th Pctl	US Hospitals 90th Pctl
55	60	49

Your hospital's median time from hospital arrival to immediate PCI for STEMI patients in minutes. Exclusions: Patients transferred in from another acute care facility; Reasons for delay does not equal none. [Detail Line:1502]

4 Proportion of STEMI patients receiving immediate PCI w/in 90'

My Hospital	US Hospitals 50th Pctl	US Hospitals 90th Pctl
99.10%	94.62%	100.00%

Proportion of STEMI patients with a time from your hospital arrival (or subsequent ECG if ST elevation first noted on subsequent ECG) to immediate PCI $\leq 90'$. Exclusions: Patients transferred in from another acute care facility; Reasons for delay does not equal none. [Detail Line:1503]

5 Median time from ED arrival at STEMI transferring facility to ED arrival at STEMI receiving facility among transferred patients.

My Hospital	US Hospitals 50th Pctl	US Hospitals 90th Pctl
69	73	48

Your hospital's median time from arrival at transferring facility to ED arrival at STEMI receiving facility among transferred patients. [Detail Line:1505]

6 Median time from ED arrival at STEMI transferring facility to immediate PCI at STEMI receiving facility among transferred patients (in minutes)

My Hospital	US Hospitals 50th Pctl	US Hospitals 90th Pctl
88	107	80

Your hospital's median time from arrival at referring facility to immediate PCI at STEMI receiving facility among transferred patients. Exclusions: Reasons for delay does not equal none. [Detail Line:1506]

7 Median fluoro time (in minutes)

My Hospital	US Hospitals 50th Pctl	US Hospitals 90th Pctl
10	10	7

Inclusion criteria: PCI of one vessel/lesion. Exclusion criteria: Prior CABG, or "other" procedure during the same lab visit; PCI of >1 vessel/lesion. [Detail Line:1633]

8 Proportion of patients with aspirin prescribed at discharge

My Hospital	US Hospitals 50th Pctl	US Hospitals 90th Pctl
99.7%	98.8%	100.0%

Proportion of patients (without a documented contraindication) with aspirin prescribed at discharge. [Detail Line:1997]

9 Proportion of patients with a P2Y12 inhibitor prescribed at discharge

My Hospital	US Hospitals 50th Pctl	US Hospitals 90th Pctl
99.7%	99.6%	100.0%

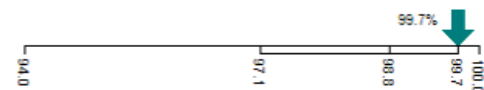
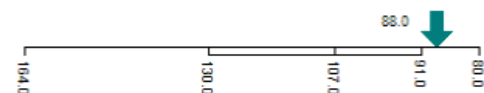
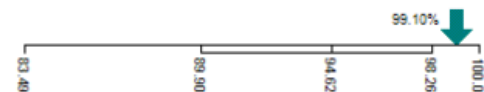
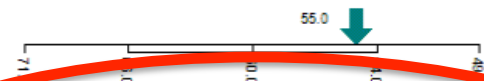
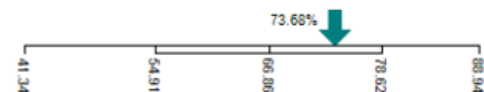
Proportion of patients (without a documented contraindication) with a stent implanted that had a thienopyridine/P2Y12 inhibitor prescribed at discharge. [Detail Line:2006]

Distribution of Hospital Performance

10th percentile

90th percentile

Better →



Documentation

- In many cases, a hospital's or physician's quality is fine, but documentation is poor.
- Develop and use tools to improve documentation.

Patient Baseline Characteristics

1. ☐ Cardiac arrest/sustained VT/VF (secondary prevention)
2. ☐ Ischemic cardiomyopathy ☐ Non-ischemic cardiomyopathy
3. ☐ EF _____ % measured by ☐ echo ☐ nuclear ☐ angiography on ____/____/____.
4. ☐ NYHA Heart Failure Class: ☐ I ☐ II ☐ III ☐ IV
5. ☐ ECG: ☐ LBBB ☐ RBBB ☐ IVCD ☐ Paced
☐ QRS duration _____ ms measured on ____/____/____.
6. ☐ Patient is on guideline-directed medical therapy (GDMT).
☐ ACEi or Contraindicated due to hypotension/CKD/intolerance or allergy
☐ ARB or Contraindicated due to hypotension/CKD/intolerance or allergy
☐ β -blocker or Contraindicated due to hypotension/intolerance or allergy
☐ Aldosterone inhibitor or Contraindicated due to hypotension/ hyperkalemia/intolerance or allergy
7. ☐ Documented prior MI greater than 40 days
☐ Pathologic Q waves on ECG
☐ Imaging evidence of a region of loss of viable myocardium that is thinned and fails to contract
☐ Pathological findings of a healed or healing MI

Medicare / CMS Guidelines for Empiric ICD

Secondary Prevention Indications

1. ☐ Documented episode of cardiac arrest due to ventricular fibrillation (VF), not due to a transient or reversible cause.
2. ☐ Documented sustained ventricular tachyarrhythmia (VT), either spontaneous or induced by an electrophysiology (EP) study, not associated with an acute myocardial infarction (MI) and not due to a transient or reversible cause.

Primary Prevention Indications

1. ☐ Documented familial or inherited conditions with a high risk of life-threatening VT, such as long QT syndrome or hypertrophic cardiomyopathy.
2. ☐ Coronary artery disease with a documented prior MI, a measured left ventricular ejection fraction (LVEF) \leq 35%, and inducible sustained VT or VF at EP study. (The MI must have occurred more than 4 weeks prior to defibrillator insertion. The EP test must be performed more than 4 weeks after the qualifying MI.)
3. ☐ Documented prior MI and a measured LVEF \leq 30% (includes NYHA class I, II, or III).
4. ☐ Patients with ischemic dilated cardiomyopathy (IDCM), documented prior MI, NYHA Class II and III heart failure, and measured LVEF \leq 35%.
5. ☐ Patients with non-ischemic dilated cardiomyopathy (NIDCM) $>$ 3 months, NYHA Class II and III heart failure, and measured LVEF \leq 35%.

Patient-Last Name, First Name, Middle Initial ☐ M ☐ F Age _____
Admission Number _____ Date _____ Birth Date _____
Physician Name _____
Patient Identification _____ Medical Record Number _____

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ICD IMPLANT DOCUMENTATION

011071

p. 1 of 2

Procedure Documentation 1062-011071

CRT Indications

1. ☐ LVEF \leq 35%, NSR, LBBB with QRSd \geq 150 ms, NYHA class II, III, or ambulatory class IV on GDMT. (class I)
2. ☐ LVEF \leq 35%, NSR, LBBB with QRSd 120-149 ms, NYHA class II, III, or ambulatory class IV on GDMT. (class Ila)
3. ☐ LVEF \leq 35%, NSR, non-LBBB with QRSd \geq 150 ms, NYHA class III, or ambulatory class IV on GDMT. (class Ila)
4. ☐ LVEF \leq 35%, permanent atrial fibrillation on GDMT if a) the patient requires ventricular pacing or otherwise meets CRT criteria and b) AV nodal ablation or pharmacologic rate control will allow near 100% ventricular pacing with CRT. (class Ila)
5. ☐ LVEF \leq 35% on GDMT, and are undergoing new or replacement device placement with anticipated requirement for significant ($>$ 40%) ventricular pacing. (class Ila)
6. ☐ LVEF \leq 30%, ischemic etiology of heart failure, NSR, LBBB with QRSd \geq 150 ms, NYHA class I on GDMT. (class IIb)
7. ☐ LVEF \leq 35%, NSR, non-LBBB with QRSd \geq 120-149 ms, NYHA class III, or ambulatory class IV on GDMT. (class IIb)
8. ☐ LVEF \leq 35%, NSR, non-LBBB with QRSd \geq 150 ms, NYHA class II on GDMT. (class IIb)

Exclusions/Contraindications for ICD Implant

1. ☐ New York Heart Association (NYHA) classification IV heart failure
2. ☐ Cardiogenic shock or symptomatic hypotension while in a stable baseline rhythm.
3. ☐ Coronary artery bypass graft (CABG) or percutaneous trans-luminal coronary angioplasty (PTCA) within past 3 months.
4. ☐ MI within past 40 days.
5. ☐ Clinical symptoms or findings that would make them a candidate for coronary revascularization.
6. ☐ Any disease, other than cardiac disease (e.g., cancer, uremia, liver failure), associated with a likelihood of survival less than 1 year.
7. ☐ Unable to give informed consent.
8. ☐ Irreversible brain damage from preexisting cerebral disease.

Patient-Last Name, First Name, Middle Initial ☐ M ☐ F Age _____
Admission Number _____ Date _____ Birth Date _____
Physician Name _____

Patient Identification _____

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ICD IMPLANT DOCUMENTATION

PINNACLE/EHR

Risk-adjusted...

- Many measures (mortality and morbidity) are risk-adjusted.
- For this reason, it is vital that risks are documented.
- Accurate characterization of the patient's acuity of illness and co-morbidities is important, but time-consuming.



Bloomington Hospital

Cardiovascular Services

Cardiac Documentation

Disclaimer: This form is not intended to be an all-inclusive diagnosis list. If the patient has a diagnosis not on this list, please document the diagnosis in "Other" category section on this form or in the other chart documentation.

X if yes	Diagnosis	Specify (Fill in the blank or place an X beside the appropriate information)	POA (Y = yes, N = no, U = Unable to determine)
CARDIAC			
<input type="checkbox"/>	STEMI	Site Anterior <input type="checkbox"/> Inferior <input type="checkbox"/> Lateral <input type="checkbox"/> Other <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	NSTEMI		Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
Occurrence:		STEMI Initial <input type="checkbox"/> or Subsequent <input type="checkbox"/> : Within 4 weeks <input type="checkbox"/> Within 8 weeks <input type="checkbox"/> NSTEMI Initial <input type="checkbox"/> or Subsequent <input type="checkbox"/> : Within 4 weeks <input type="checkbox"/> Within 8 weeks <input type="checkbox"/>	
<input type="checkbox"/>	Unstable Angina		Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Cardiogenic Shock		Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Stent Stenosis	Progression of CAD <input type="checkbox"/> or Complication of cardiac stent <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Valvular Heart Disease	Specify valve and disorder: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Atrial Fibrillation	Rate Controlled <input type="checkbox"/> or RVR <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Other Dysrhythmia	Specify: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Heart Failure	EF <input type="checkbox"/> % Acute <input type="checkbox"/> or Chronic <input type="checkbox"/> or Acute on Chronic <input type="checkbox"/> AND Systolic <input type="checkbox"/> or Diastolic <input type="checkbox"/> or Systolic/Diastolic <input type="checkbox"/> NYHA Heart Failure Class: I <input type="checkbox"/> II <input type="checkbox"/> III <input type="checkbox"/> IV <input type="checkbox"/> Contraindication to ACEI/ARB if EF < 40% <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Cardiomyopathy	Due to: HTN <input type="checkbox"/> or HF <input type="checkbox"/> or Ischemia <input type="checkbox"/> or Valvular <input type="checkbox"/> or Other: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	H/O PCI	Specify vessels: LAD <input type="checkbox"/> Diagonal <input type="checkbox"/> LCx <input type="checkbox"/> OM <input type="checkbox"/> RCA <input type="checkbox"/> PDA <input type="checkbox"/> Ramus <input type="checkbox"/> Other: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	H/O CABG	Specify vessels: LAD <input type="checkbox"/> Diagonal <input type="checkbox"/> LCx <input type="checkbox"/> OM <input type="checkbox"/> RCA <input type="checkbox"/> PDA <input type="checkbox"/> Other: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	H/O Cardiac Device	Specify Type: Pacemaker <input type="checkbox"/> ICD <input type="checkbox"/> and Single <input type="checkbox"/> Dual <input type="checkbox"/> CRT <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>

RENAL/HYDRATION

<input type="checkbox"/>	ERSD		Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	CKD	Stage I <input type="checkbox"/> or II <input type="checkbox"/> or III <input type="checkbox"/> or IV <input type="checkbox"/> or V <input type="checkbox"/> Cause: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Acute Renal Failure	Cause: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Dehydration		Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Fluid Overload	Due to HF <input type="checkbox"/> Other: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Urinary Tract Infection	Organism: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>

VASCULAR

<input type="checkbox"/>	Peripheral Artery Disease	Arteriosclerotic <input type="checkbox"/> or Other: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Hypertension	Benign <input type="checkbox"/> or Malignant/Accelerated <input type="checkbox"/> or Chronic <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>

PULMONARY

<input type="checkbox"/>	COPD	Exacerbation/Acute <input type="checkbox"/> or Without Exacerbation <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Pneumonia	Specify Type: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Chronic Respiratory Failure	End Stage Lung Disease <input type="checkbox"/> Home Oxygen Use <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Acute Respiratory Failure		Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Pleural Effusion	Associated with HF <input type="checkbox"/> or Malignant <input type="checkbox"/> or Acute on Chronic <input type="checkbox"/> AND Cardiogenic <input type="checkbox"/> or Post-op Complication <input type="checkbox"/> or Other: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Pulmonary Edema	Acute <input type="checkbox"/> or Chronic <input type="checkbox"/> or Acute on Chronic <input type="checkbox"/> AND Cardiogenic <input type="checkbox"/> or Post-op Complication <input type="checkbox"/> or Other: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>

ENDOCRINE/NUTRITIONAL/DIGESTIVE

<input type="checkbox"/>	Diabetes	Type I <input type="checkbox"/> or Type II <input type="checkbox"/> AND Controlled <input type="checkbox"/> or Uncontrolled <input type="checkbox"/> Specify Diabetic Complication: Retinopathy <input type="checkbox"/> Neuropathy <input type="checkbox"/> Nephropathy <input type="checkbox"/> PAD <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Hypokalemia		Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Hyperkalemia		Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Hyponatremia		Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Hypernatremia		Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
<input type="checkbox"/>	Hypothyroidism	Specify Cause: <input type="checkbox"/>	Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>