# the**quality**post

Р.

this

issue

Make Group Problem Solving More Effective

Distinguishing Lean From Taylorism

Clinical Documentation Improvement

Sustaining the Gains: Med Rec Kaizen

4+1 Metrics

Monthly Quality Improvement Newsletter for the Division of Hospital Medicine

July 2016 • Issue 67

#### Greetings from Cat, Nader and Sasha

QUALITY IMPROVEMENT DIVISION OF HOSPITAL MEDICINE Welcome to the 67th edition of The Quality Post. In this issue we feature a piece on the group problem solving and distinguishing Lean from Taylorism. We also highlight updates in CDI, a piece on where are now with med rec, and data on our 4+1 metrics.

## **Distinguishing Lean from Taylorism**

In a NEJM piece on Lean published earlier this year, Pamela Hartzband, MD and Jerome Groopman, MD claimed that the Toyota Production System (TPS)/Lean was inspired by the principles of "Taylorism" made popular by Frederick Taylor in the early 20th century. John Toussaint, MD, CEO of the ThedaCare Center for Healthcare Value wrote a response published by the Health Affairs blog distinguishing TPS/Lean from Taylorism.

Taylor believed that there was one way to accomplish a task, and it was up to management determine that way and make sure all workers executed the plan. In contrast, TPS/Lean is based on the teaching of W. Edwards Deming who argued that frontline works should be in charge of improvement processes

The application of TPS/Lean in healthcare is relatively nascent, and it is true that it has failed in some organizations, but advocates of Lean argue that this is likely due to misguided implementation. Healthcare institutions like ThedaCare, Stanford, and others that have adhered to the original principles of TPS, including respect for frontline workers, have seen extraordinary results.

"There remains too much unjustified waste and unwarranted complexity in delivering care to patients, exposing them to errors and complications. It is time to marry the science of management systems as embodied in TPS/Lean with the science of medicine to achieve care that is safe, efficient, effective, personalized, timely, and equitable." --Toussaint

As we move forward with our Lean journey at UCSF, it will be essential to learn from the pitfalls at other institutions and to participate in the movement towards building national standards for applying TPS in health care.

Source: http://healthaffairs.org/blog/2016/04/06/the-toyotaproduction-system-what-does-it-mean-and-what-does-it-mean-forhealth-care/

## Make Group Problem Solving More Effective

When groups get together to brainstorm, they actually come up with fewer ideas than the individuals in that group would have come up with on their own. That's why it's important to think about group problem solving in two phases: divergence and convergence.

Divergence happens when the group considers as many different potential solutions as possible. For example, "How many different uses can you find for a brick?"

Convergence happens when a large number of ideas are whittled down to a smaller set.

For the best results, have people work alone when generating ideas. Then collect those ideas and send them around to the aroup. Allow the divergence to continue as group members individually build on the ideas of their colleagues. Give the resulting ideas to everyone and let the group get together to pick the best ones. This way everyone can offer solutions without being unduly influenced by others' ideas.

Harvard Business Review: Adapted from "The Problem-Solving Process That Prevents Groupthink," by Art Markman



#### **GOALS OF THE PROGRAM:**

- Document, capture and code all diagnoses, procedures, co-morbidities and complications
- Accurately and completely reflect the clinical complexity of our patients and the quality of their care
- Improve publically reported measures and ratings

#### What is the relationship between our quality outcomes and how we document?

Hospitals are increasingly being judged on our quality outcomes-- outcomes like mortality, LOS, and hospital acquired complications; all of which are adjusted by patient's severity of illness.

Capturing severity of illness thus becomes increasingly important if we want our quality measurements to be accurate measures of our performance as physicians and as a medical center.

### How is Severity of Illness measured?

Severity of Illness is based on the **Case Mix Index (CMI)** of your patient population.



The CMI is used to **Risk Adjust** patient outcomes.

Every Medicare principle diagnosis or **MS-DRG** is associated with a case mix index.

#### **Complications and Comorbidities**

AND

Major Complications and Comorbidities

Add to the principle diagnosis or the DRG to increase the Case Mix Index and expected LOS of patients.

		р	atients	•
<u>.</u>	MS-DRG		CMI	LOS
diagnosis	446	Disorders of Biliary Tract w/o CC or MCC	0.7	2.4
groups of 44X differ		Secondary Diagnosis- leukocytosis		
in the presence or	445	Disorders of Biliary Tract w CC	1.0	3.5
lack of CC's or MCC's.		Secondary Diagnosis- bacteremia or cholar	gitis	
	444	Disorders of Biliary Tract w MCC	1.6	4.7
		Secondary Diagnosis- Severe sepsis		

Documenting *"Leukocytosis"* will result in severely under representing this patient's severity of illness.

If we want our quality outcomes and LOS to be judged fairly we need to pay attention to how we document co-morbidities and complications

# **Clinical Documentation Improvement**

Attention to just some simple specific terms can eliminate the need for most queries!

The old way	The new way
CKD	CKD and the stage
Volume overload Echo shows low EF HFpEF or HFrEF	Acute pulmonary edema Acute/Chronic systolic heart failure Acute/Chronic diastolic heart failure
Vent dependent Unable to wean from vent	Acute hypoxemic respiratory failure
Requiring pressors	Septic/Cardiogenic Shock
Rising Creatinine	AKI/ ATN (remember contrast nephropathy is ATN
GI bleed	Acute anemia 2/2 Blood Loss
AMS	Encephalopathy or Coma

#### Other more unique issues to be aware of:

#### SIRS & SEPSIS

- Look for SIRS with every infection
  - $\circ$  SIRS + Infection = Sepsis
- Look for organ dysfunction and document Severe Sepsis
- Document Sepsis in the discharge summary (let's the coders know this was a confirmed Dx)





Sijiary Sepsis

 SIRS without an infection is still a marker of severity (Think of a severe COPD patient)

#### **HEALTH CARE ASSOCIATED PNA**

- Not all PNA are created equal
- Health care associated PNAs carry a higher risk of readmission
- Unfortunately documenting HCAP is not enough!
  - Document broad spectrum antibiotics and the organisms the are covering for

HCAP: Treating with Zosyn for possible pseusomonas and vancomycin for possible MRSA

#### NSTEMI TYPE II vs. DEMAND ISCHEMIA

**The problem:** We use these terms interchangeably, but they are different codes! In the coding world: NSTEMI = Acute MI To be accurate use:

- NSTEMI Type II for TROPONIN + EKG changes, Typical Symptoms or wall motion abnormalities
- **Demand Ischemia** for isolated Troponin Elevation in the setting of non-cardiac disease

#### **ENCEPHALOPATHY**

#### What is encephalopathy?

A global or diffuse alteration in brain function associated with a systemic cause.

#### **Examples:**

- Toxic intoxication/withdrawal or over medication
- Infectious AMS from sepsis
- Hepatic Excess ammonia
- Ischemic Associated with stroke or shock

# Sustaining the Gains: Med Rec Kaizen

The Med Rec Kaizen took place in September of 2015, and brought together a multidisciplinary group of ED and Medicine pharmacists, housestaff, an attending and an RN. With the goal of improving the efficiency, safety, and quality of the medication reconciliation process, the team created standard work and training materials for housestaff, attendings, pharmacists and nurses. Since then, the UCSF Med Rec Committee and the Housestaff Incentive Program (HIP) residents have been continuing to work hard on sustaining the gains.

#### So how are we doing almost a year after the Kaizen? Not only have we improved the percent of patients with medications reconciled prior to discharge, but we now have data to show that we have improved the quality of the med rec process.

The percent of patients with no duplicate meds and the percent of patients with prior to admission meds has improved. Below are statistical process control charts showing a new set of upper and lower control limits (3 sigma above and below the centerline) based on our new and improved system.



#### Complete admission medication reconciliation prior to discharge

# DHM/Residency "4+1" Priorities

# Four Core Metrics:

Achieve ≥ 75% score for HCAHPS Communication MD "Explained in Understandable Way"					7	7 of 12 months						
FY2015 Baseline	July	Aug	Sept	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	Jun
<b>74.9</b> %	71.4	83.3	81.8	67.7	84.8	89.7	81.0	70.5	69.4	73.3	73.3	83.3
Sustain number of total phlebotomy draws by achieving ≤ 1.7 sticks per hospitalized patient per day						0	6 of 12 months					
FY2015 Baseline	July	Aug	Sept	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	Jun
1.7	1.56	1.51	1.51	1.64	1.60	1.51	1.50	1.42	1.45	1.55	1.54	1.61
Achieve ≥ 90% of patients who have had all medications reconciled before discharge				2	4 of 12 months							
FY2015 Baseline	July	Aug	Sept	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	Jun
77.7%	76.5	86.7	84.3	88.9	88.1	88.7	89.3	89.1	90.2	92.6	90.3	87.2
Achieve ≥ 20% of hospital medicine discharges by noon					7	8 of 12 months						
FY2015 Baseline	July	Aug	Sept	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	Jun
20.1%	20.4	16.0	15.7	17.1	18.5	18.9	18.3	23.8	16.8	19.6	16.7	20.0
Plus One Metrics: Achieve ≤ 23% of patients on telemetry until discharge (with LOS > 48hrs)						6 of 12 months						
48hrs)	on telem	etry unt	il discha	rge (wit	h LOS >	0			6 of 12	months		
48hrs) FY2015 Baseline	July	etry unt	il discha Sept	rge (wit Oct	h LOS > Nov	Dec	Jan	Feb	6 of 12 Mar	months Apr	May	Jun
48hrs) FY2015 Baseline 22.6%	July 23.0	etry unt Aug 24.2	il discha Sept 23.6	rge (with Oct 29.0	h LOS > Nov 17.5	Dec 22.1	Jan 21.2	Feb 18.3	6 of 12 Mar 20.0	Months Apr 20.7	May 22.1	Jun 17.9
48hrs) FY2015 Baseline 22.6% Achieve ≥ 75% patients wit	July 23.0 h High-C	Aug 24.2 Luality A	Sept 23.6	rge (with Oct 29.0	Nov 17.5	Dec 22.1	Jan 21.2	Feb 18.3	6 of 12 Mar 20.0 6 of 12	Months Apr 20.7 months	May 22.1	Jun 17.9
48hrs) FY2015 Baseline 22.6% Achieve ≥ 75% patients wit FY2015 Baseline	July 23.0 h High-C July	Aug 24.2 Quality A	Sept 23.6 VS Sept	rge (with Oct 29.0 Oct	Nov 17.5 Nov	Dec 22.1 Dec Dec	Jan 21.2 Jan	Feb 18.3 Feb	6 of 12 Mar 20.0 6 of 12 Mar	Months Apr 20.7 months Apr	May 22.1 May	Jun 17.9 Jun
48hrs) FY2015 Baseline 22.6% Achieve ≥ 75% patients with FY2015 Baseline 74.4%	July 23.0 h High-C July 65.1	Aug 24.2 Quality A Aug 69.1	il discha Sept 23.6 VS Sept 62.0	rge (with Oct 29.0 Oct 65.0	Nov 17.5 Nov 65.2	Dec 22.1 Dec 65.7	Jan 21.2 Jan 64.6	Feb 18.3 Feb 78.5	6 of 12 Mar 20.0 6 of 12 Mar 78.4	Months Apr 20.7 months Apr 75.3	May 22.1 May 69.4	Jun 17.9 Jun 65.3
48hrs) FY2015 Baseline 22.6% Achieve ≥ 75% patients with FY2015 Baseline 74.4% Achieve C Diff rate of ≤ 11.	July 23.0 h High-C July 65.1	Aug 24.2 Quality A Aug 69.1	Sept 23.6 VS Sept 62.0 tient da	rge (with Oct 29.0 Oct 65.0 ys)	Nov 17.5 Nov 65.2	Dec 22.1 Dec 65.7	Jan 21.2 Jan 64.6	Feb 18.3 Feb 78.5	6 of 12 Mar 20.0 6 of 12 Mar 78.4 6 of 12	months Apr 20.7 months Apr 75.3 months	May 22.1 May 69.4	Jun 17.9 Jun 65.3
48hrs) FY2015 Baseline 22.6% Achieve ≥ 75% patients with FY2015 Baseline 74.4% Achieve C Diff rate of ≤ 11. FY2015 Baseline	July 23.0 h High-C July 65.1 1 (per 10 July	Aug 24.2 Quality A Aug 69.1 D,000 pa Aug	Sept 23.6 VS Sept 62.0 tient dav	rge (with Oct 29.0 Oct 65.0 ys) Oct	Nov 17.5 Nov 65.2 Nov	Dec 22.1 Dec 65.7	Jan 21.2 Jan 64.6 Jan	Feb 78.5 Feb	6 of 12 Mar 20.0 6 of 12 Mar 6 of 12 Mar	Months Apr 20.7 months Apr 75.3 months Apr	May 22.1 May May	Jun 17.9 Jun 65.3

Achieve 50% of patients (not full code) with POLST completion					tion	0		6 of 12 months					
FY2015 Baseline	July	Aug	Sept	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	Jun	
40.2%	37.1	52.1	51.5	50.8	62.2	52.1	59.3	45.6	66.7	63.4	57.4	57.7	