

Opportunities and Obstacles in Using Routinely Collected Electronic Data for Health Services Research

The Example of Identifying Infection Risk Factors and Outcomes

CIRAR Seminar Series
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The Electronic Medical Record: Projections and Concerns for Policy

- Potential for long-term cost savings
- Possible perverse consequences of decision support: flagging patient safety issues has not necessarily reduced incidence
- Limitations of ICD-9 coding for reimbursement

Little discussion on the challenges of using electronic data for research purposes...

Exciting Possibilities for Research Using Electronic Data

- Post-marketing pharmaceutical research
- Clinical decision support systems
- Policy and payment changes
- Study of hospital phenomena (e.g. antimicrobial resistance)

Obstacles: Through the Lens of Antimicrobial Resistant Infection

1. Cooperation from Institutions and Individual Departments
2. Timing Issues
3. Integrating Disparate Datasets
4. Limits of Automated Case Finding
5. Definition Variation
6. Construct Validity
7. Data Reliability
8. Free Text
9. Time Stamps

“Distribution of the Costs of Antimicrobial Resistance” (Grant R01 NR010822): Aims

1. Estimate hospital costs of resistant infections
2. Assess extent to which patients other than the initially infected patient experience higher costs due to cases of antibiotic resistance
3. Examine the distribution of the costs of resistant infection between hospitals, third-party payers, and patients
4. Model the impact of alternative payment policies on incentives to invest in reducing resistant infections
5. Evaluate whether observed patterns of investment in the control of resistant infections are related to the distribution of the costs of infection

Performance Sites

New York Presbyterian Hospital

Columbia campus

- Milstein/Presbyterian Hospital – 700 bed tertiary
- Morgan Stanley Children's Hospital – 257 bed children's hospital

Cornell campus

- New York Hospital – 797 bed tertiary

Allen pavilion

- 212 bed community

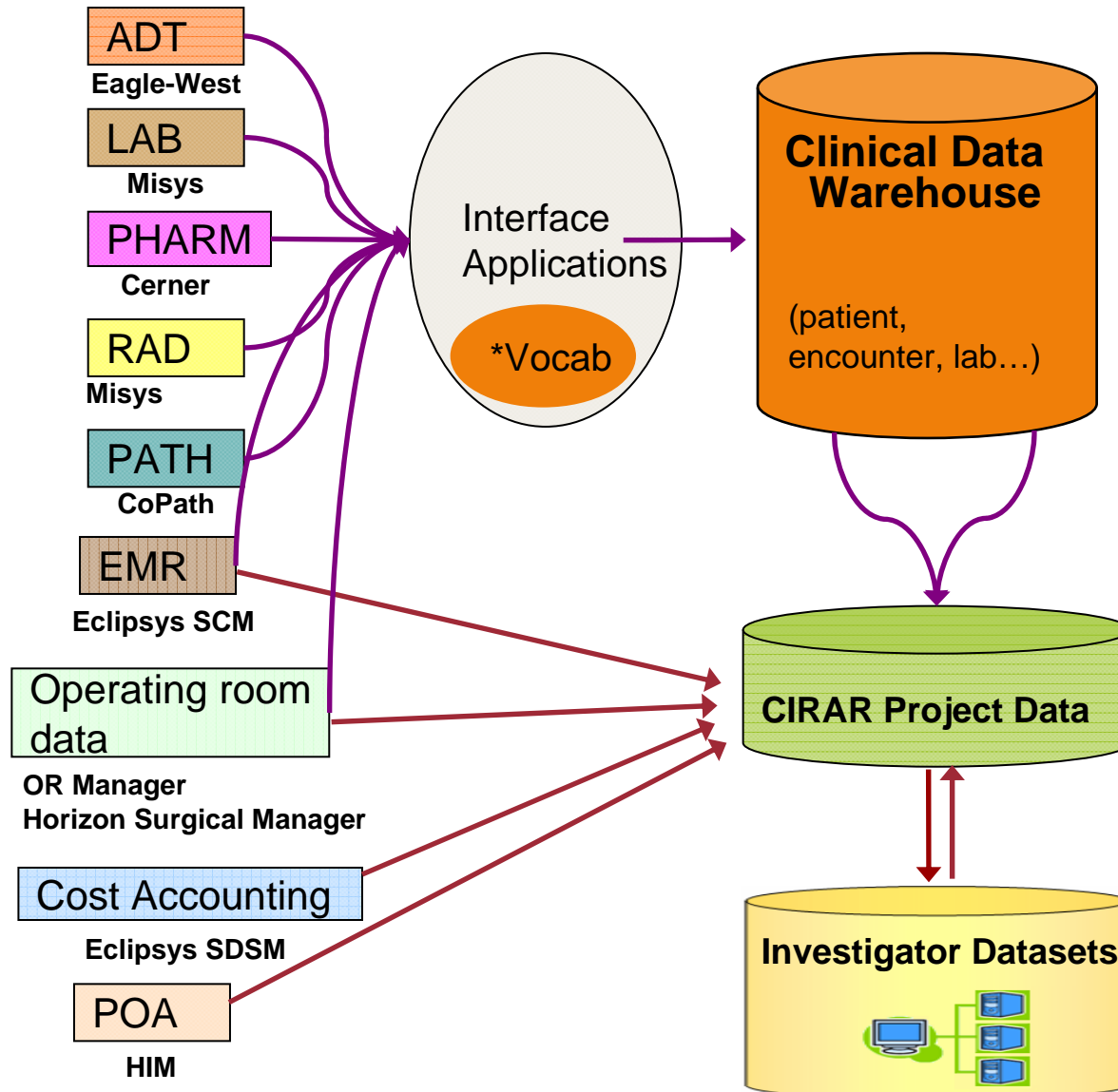
Methods

- Data linkage
- Identify cases resistant infection 2005-2009
- Match cases to susceptible and non-infected controls
- **Aim 1** – compute length of stay, costs, breakdown of costs
- **Aim 3** – determine primary payer expenditures; incidence of costs

Data Needs

- **Organism** – Enterococci, Staphylococcus aureus, Klebsiella pneumoniae, Acinetobacter baumannii, Pseudomonas aeruginosa, Streptococcus pneumoniae
- **Sensitivity/resistance pattern**
- **Infection Site** – surgical, bloodstream, urinary tract, pneumonia
- **Infection Source** – hospital vs. community acquired
- **Risk factors for infection** – age, trauma, burns, diabetes, renal failure, malignancies, endotracheal intubation, central venous catheters, urinary catheters, mechanical ventilation, invasive procedures, immunosuppressive agents, length of hospital stay, stay in ICU
- **Risk factors for Resistant Infection** - prior hospitalization, exposure to long-term/broad-spectrum antibiotics transplantation, gastrointestinal surgery

High Level Data Architecture: Columbia (MHB, MSCH, TAP)



* Medical Entity Dictionary (MED)

Obstacle #1: Cooperation from Institutions and Individual Departments

- Gaining access to a single department's data requires cooperation and approval from multiple parties
 - Data Owners
 - Data Custodians

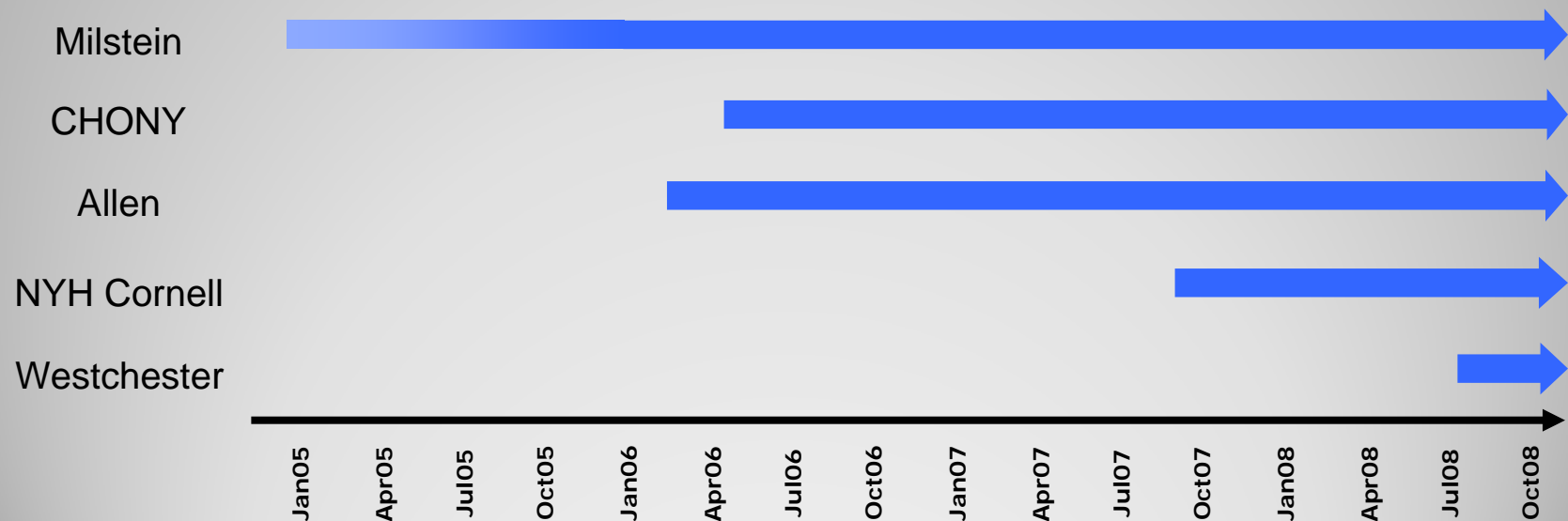


Mutual Benefit

Obstacle #2: Timing Issues

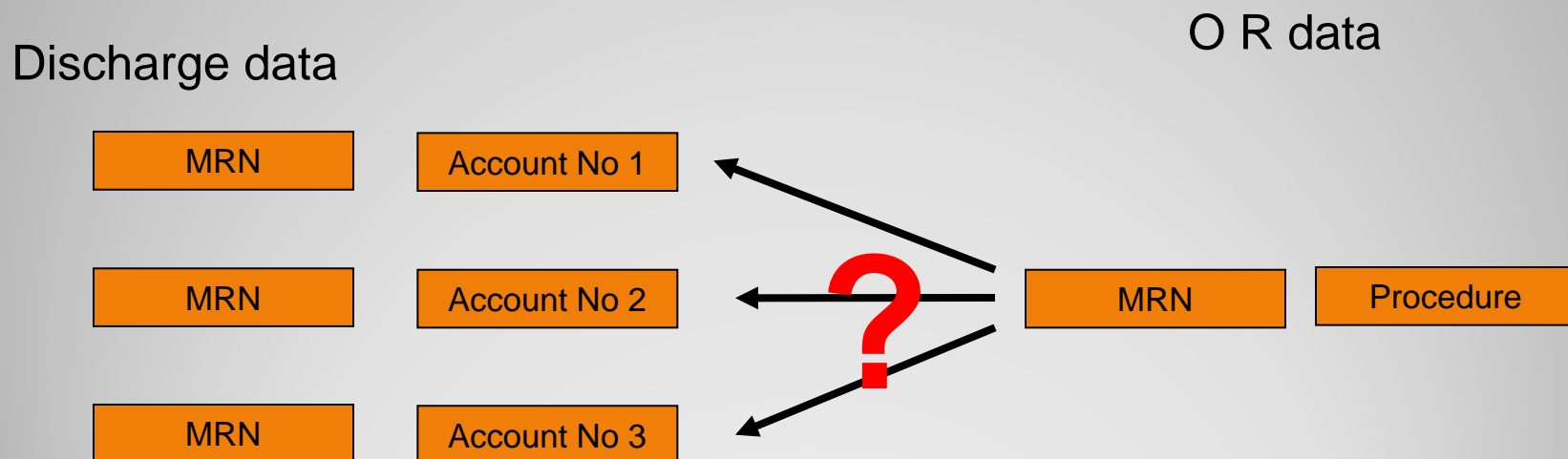
- Dates for which data are available vary between different data systems

Eclipsys XA rollout



Obstacle #3: Integrating Disparate Datasets

- Departments use different methods to identify patients and encounters



Obstacle #4: Limits of Electronic Logic

- How do the results of an automated case finding program using electronic data compare with the “gold standard” of a clinician’s or epidemiologist’s judgment?
- How should the validity of electronic data be assessed?

Obstacle #5: Definition Variation

- Variable definitions can vary between data sources
- Available variable definitions may not be appropriate for your project

EXAMPLE:

(1) Defining Infections

(2) Operating Room Procedure Time

Definition Variation

Defining MRSA Infection

Epi-portal

- Culture positive for MRSA
- Culture positive for Staph. aureus and resistant to Oxacillin or Methicillin

Project

- Positive blood culture for organism excluding common skin contaminants (i.e., diphtheroids { *Corynebacterium* spp.}, *Bacillus* {not *B. anthracis*} spp., *Propionibacterium* spp., coagulase-negative staphylococci, viridans group streptococci, *Aerococcus* spp., *Micrococcus* spp.)
- No positive culture of same organism from different infection site in past 30 days
- Two or more positive blood cultures collected within 2 days of each other drawn on separate occasions
- Fever $>38^{\circ}\text{C}$

Definition Variation

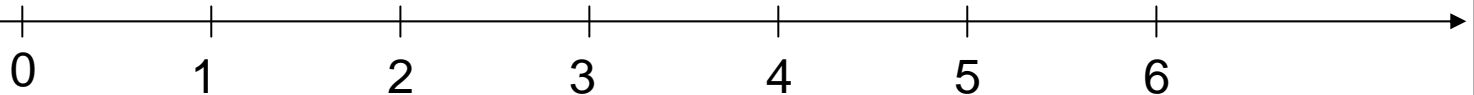
Operating Room Procedure Time

Procedure A



Procedure B

Procedure C



Total procedure time

$$A + B + C = 10?$$

$$\text{Start of A to end of B} = 5?$$

Obstacle #6: Construct Validity

- Do the data actually represent what you believe them to represent?

EXAMPLE: Has the patient received an antibiotic in the past six months?

Physician Order Written?	YES
Patient Charged for Drug?	YES
Drug Administered to Patient?	NO

Obstacle #7: Data Reliability

- A variable of interest may not be well coded
- Are missing/misclassified data systematic or random?

EXAMPLE: Urinary Catheterization

Discharge data (ICD 9)	Clinical data (Eclipsys XA)		
		Present	Absent
Present	11	1	12
Absent	7	5	12

Obstacle #8: Free Text

- Do free text queries reliably capture the variable of interest?

EXAMPLE: **Nursing Home Residency**

Structured Field: **Patient admitted from Nursing Home?**
*Yes, **No***

Free Text: **Patient's Home Address**
1060 Amsterdam Ave. Apt 8C
1060 Amsterdam Avenue #8C
1060 Amsteerdam Apt. 8C

1060 Amsterdam Avenue is a Nursing Home

Obstacle #9: Time Stamps

- Is it possible to determine the sequence of events that occurred during a patient's stay?
- Can your data establish that a supposed risk factor occurred before disease onset?
- Was a condition present on admission, or did it develop during the patient's hospital stay?

EXAMPLE: Renal Failure

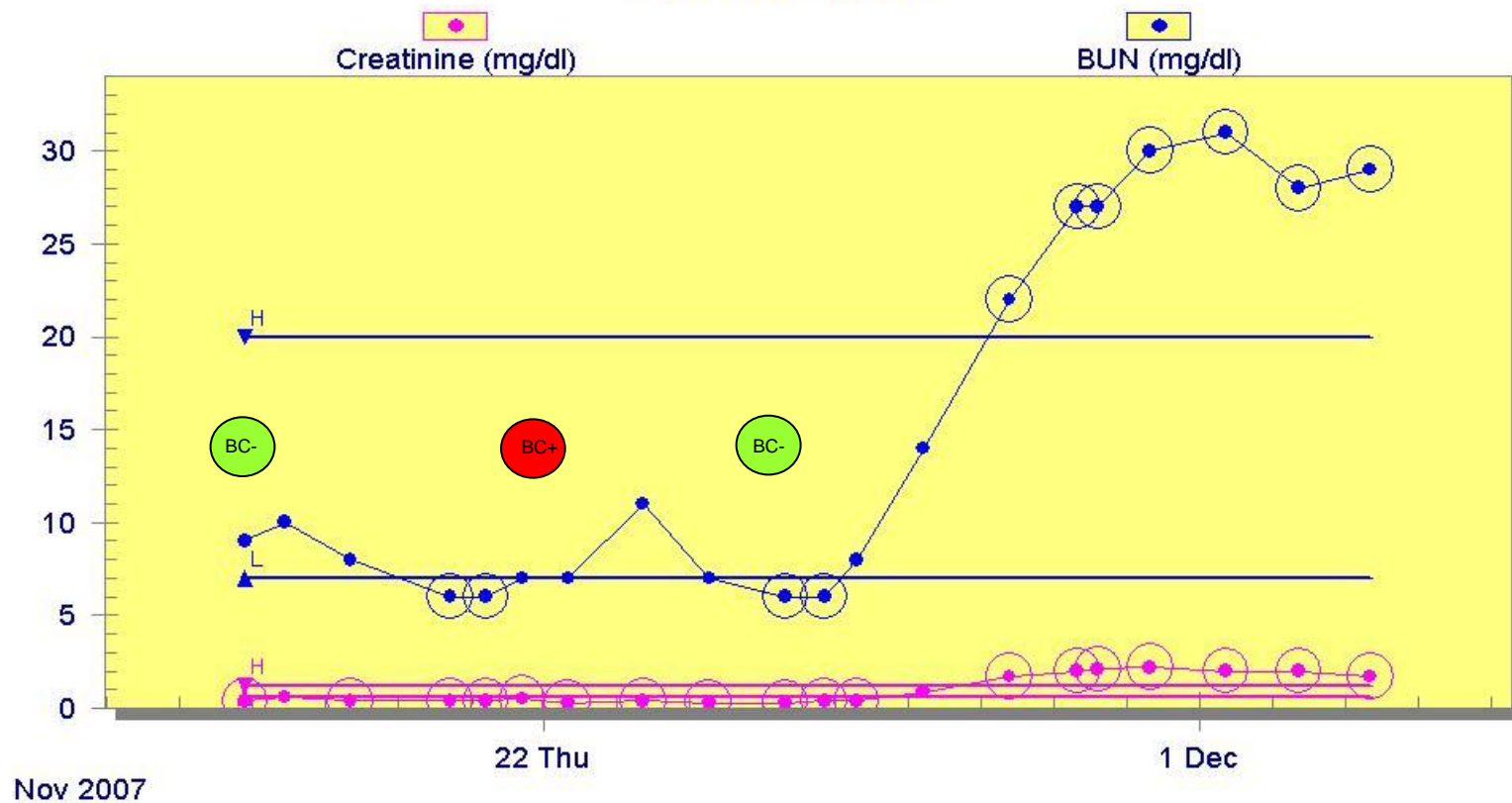
Time Stamps Renal Failure

Acute renal failure

Adm: 17 Nov, Disch: 4 Dec

POA flag = No

BUN/Creatinine



BC: Blood culture

National Issues: Comparisons Across Hospitals

- Overcoming obstacles 1-9 within a single hospital does not ensure that the data will be comparable to that of another hospital

EXAMPLE: Differences in ICD-9 coding practices across hospital

National Issues: ICD-9 CM Documentation and Coding Variation

“CMS Won’t pay for Medical Errors” Patient Example

<p>Admit Source:</p> <p>Admit Type: EMERGENCY/URGENT</p> <hr/> <p>MS DRG: 834 Acute leukemia w/o major O.R. procedure w MCC Revised = 835, “w CC”</p> <p>NYDRG: 576 ACUTE LEUKEMIA W MAJOR CC</p> <p>APDRG: 690 ACUTE LEUKEMIA</p> <p>Mort. Risk: 3 Sev. Level: 3</p>	<p>Discharge Disp: ROUTINE</p> <p>Financial Group: MCR FEE FOR SERVICE</p> <p>Coded Date: 4/1/2008</p> <p>Clin. Case Type: HEMATOLOGIC MALIGNANCIES</p> <p>Service Line: ONCOLOGY</p> <hr/> <p>CMI: 4.59</p> <p>Rev: 2.58</p> <p>LOS: 36</p>
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<p>Principal Diagnosis: 205.00 ACUTE MYELOID LEUK W/O REMISSN</p> <p>Secondary Diagnosis:</p> <table style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 15%;">ICD-9 Code</th> <th style="width: 45%;">Description</th> <th style="width: 15%;">POA</th> <th style="width: 25%;">MS CC:</th> </tr> </thead> <tbody> <tr><td>284.1</td><td>Pancytopenia</td><td>POA</td><td>CC</td></tr> <tr><td>285.9</td><td>Anemia, unspecified</td><td>POA</td><td>---</td></tr> <tr><td>287.4</td><td>Secondary thrombocytopenia</td><td>POA</td><td>---</td></tr> <tr><td>288.03</td><td>Drug induced neutropenia</td><td>POA</td><td>---</td></tr> <tr><td>311</td><td>Depressive disorder, not elsewhere</td><td>POA</td><td>---</td></tr> <tr><td>401.9</td><td>Essential hypertension; unspecified</td><td>POA</td><td>---</td></tr> <tr><td>414.01</td><td>Coronary atherosclerosis; of native</td><td>POA</td><td>---</td></tr> <tr style="border: 2px solid black;"><td>707.03</td><td>Decubitus ulcer; lower back</td><td>HAC</td><td>MCC</td></tr> <tr><td>V10.46</td><td>Personal history of malignant neo</td><td>POA</td><td>---</td></tr> <tr><td>V45.82</td><td>Percutaneous transluminal corona</td><td>POA</td><td>---</td></tr> </tbody> </table> <p># of Secondary Diagnoses: 10 # HACs: 1</p>	ICD-9 Code	Description	POA	MS CC:	284.1	Pancytopenia	POA	CC	285.9	Anemia, unspecified	POA	---	287.4	Secondary thrombocytopenia	POA	---	288.03	Drug induced neutropenia	POA	---	311	Depressive disorder, not elsewhere	POA	---	401.9	Essential hypertension; unspecified	POA	---	414.01	Coronary atherosclerosis; of native	POA	---	707.03	Decubitus ulcer; lower back	HAC	MCC	V10.46	Personal history of malignant neo	POA	---	V45.82	Percutaneous transluminal corona	POA	---	<p>Principal Procedure: 99.25 INJECT CA CHEMOTHER NEC</p> <p>Secondary Procedures:</p> <table style="width: 100%; border-collapse: collapse;"> <tbody> <tr><td>38.93</td><td>VENOUS CATHETER NEC</td></tr> <tr><td>41.31</td><td>BONE MARROW BIOPSY</td></tr> <tr><td>87.41</td><td>C.A.T. SCAN OF THORAX</td></tr> <tr><td>99.04</td><td>PACKED CELL TRANSFUSION</td></tr> <tr><td>99.05</td><td>PLATELET TRANSFUSION</td></tr> </tbody> </table> <p># of Secondary Procedures: 5</p>	38.93	VENOUS CATHETER NEC	41.31	BONE MARROW BIOPSY	87.41	C.A.T. SCAN OF THORAX	99.04	PACKED CELL TRANSFUSION	99.05	PLATELET TRANSFUSION
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Fictitious Data for Demonstration Purposes

National Issues: A Challenging Research Patient!

TRACHEOSTOMY W LONG TERM MECHANICAL VENTIL

Mort. Risk: 4 **Sev. Level:** 4

LOS: 192

VAR: 141.9

Principal Diagnosis: 710.0

SYST LUPUS ERYTHEMATOSIS

Principal Procedure: 84.11

TOE AMPUTATION

Secondary Diagnosis: **POA:** **: MS CC** **CMS Tgt:**

038.0	Streptococcal septicemia	POA	MCC	0
112.5	Candidiasis; disseminated	HAC	MCC	0
286.9	Other and unspecified coagulation	POA	CC	0
344.09	quadriplegia and quadripareis; ot	POA	MCC	0
403.91	Hypertensive renal disease; unspe	POA	CC	0
440.24	Atherosclerosis of the extremities	POA	CC	0
486	Pneumonia, organism unspecified	HAC	MCC	0
518.81	Acute respiratory failure	HAC	MCC	0
567.29	Other suppurative peritonitis	HAC	MCC	0
584.9	Acute renal failure, unspecified	HAC	MCC	0
585.6	End stage renal disease	POA	MCC	0
599.0	Urinary tract infection, site not sp	HAC	CC	4.
707.03	Decubitus ulcer; lower back	HAC	MCC	5.
707.14	Ulcer of heel and midfoot; chronic	POA	CC	0
785.52	Shock without mention of trauma;	HAC	MCC	0
995.92	Systemic inflammatory response s	POA	MCC	0
997.01	Cardiac complications	HAC	CC	0
997.03	Respiratory complications	HAC	CC	0
998.11	Hemorrhage complicating a proce	HAC	CC	0
998.12	Hematoma complicating a proced	HAC	CC	0
999.31	Infection due to central venous ca	HAC	CC	6.
999.39	Infection following other infusion,	HAC	CC	0
V46.11	Dependence on respirator, status	HAC	CC	0

of Secondary Diagnoses: 23

HACs: 15

- Medpar research file from claims (only 8 secondaries)
- Relationships to Clinical Databases (ie CLABSI, UTI, Pressure Ulcer)?
- UTI is a CMS “HAC” only when in combination with catheter-related UTI code (rarely documented)
- Acute Renal Failure: Risk or outcome depends on timing
- Variation across hospitals

Fictitious Data for Demonstration Purposes

National Issues

Acute Renal Failure ICD-9 Coding Variation Across AMCs (Major Chest Procedures DRG; Q1 2008 Medicare)

BARNES JEWISH	MASS GENERAL	NYPH CORNELL	UCSF
BRIGHAM & WOMENS	MAYO CLINIC	NYU	UNIV CHICAGO
CEDARS-SINAI	METHODIST HOUSTON	RUSH	UNIV MICHIGAN
DUKE	MOUNT SINAI NYC	ST LUKES HOUSTON	UNIV PENNSYLVANIA
EMORY	NORTHWESTERN	STANFORD	VANDERBILT
JOHNS HOPKINS	NYPH COLUMBIA	UCLA	WASHINGTON

← (HIGHER) Severity – Adjusted Payment Per Case (LOWER) →

AMC:	1	2	3	4	5	6	7	25	26	27	28	29	30
5845	9%		7%	5%			2%		4%	3%	2%		
5849	14%	15%	7%	10%		3%	4%	3%		3%	2%	8%	
Total	23%	15%	14%	15%	0%	3%	7%	3%	4%	5%	3%	8%	0%

5845 Acute Renal Failure with Tubular Nephrosis

5849 Acute Renal Failure Not Otherwise Specified

Summary

- Although data access obstacles and analytic considerations have slowed the pace of progress, we have overcome!
(Thanks to cooperation and goodwill from collaborating institutions and programmers!)
- Our team looks forward to the many research opportunities to come from routinely collected electronic data