

Documentation Impact and the Physician Advisor

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# Who Governs Our Coding Language?

There are four designated entities who control our coding language

These entities are the gate keeper for how words documented in the

medical record translate to medical codes and which diagnoses are

recognized in the coded record.



#### 4 Cooperating Parties

- CDC responsible for diagnoses (the government)
- CMS responsible for inpatient procedures (the government)
- American Hospital Association responsible for interpreting ICD-10

   Through Coding Clinic
- American Health Information Management Association (AHIMA)
  - $_{\rm \circ}$  Provides input from the coding community

#### Who is missing?



# **Coding Clinic**

- Quarterly newsletter published by the American Hospital Association
- Coding Clinic
  - American Hospital Association (AHA)
  - American Health Information Management Association (AHIMA)
  - Centers for Disease Control and Prevention (CDC)
  - Centers for Medicare and Medicaid Services (CMS)
- The Editorial <u>Advisory</u> Board consists of an expert panel of physicians representing the American Medical Association, the American College of Surgeons, the American Academy of Pediatrics and the American College of Physicians, as well as coding professionals representing healthcare facilities.



# A Few Concepts



#### Support Hospital Quality Reporting through Clinical Documentation Integrity

#### Medicare.gov Hospital Compare







#### Awards

Healthgrades awards tell you which hospitals deliver superior quality care. Healthgrades evaluates hospital performance using objective **quality measures including clinical outcomes and patient safety**, as well as patient experience.





U.S. News Announces 2018-19 Best Hospitals



| Hospital Acquired | Hosp. Readmission | Hospital Value-  |  |
|-------------------|-------------------|------------------|--|
| Conditions        | Reduction         | Based Purchasing |  |



#### Pay For Performance

#### Value-Based Programs

The left column—HAC Deficit Reduction Act—puts your CC or MCC at risk if the diagnosis is not POA = YES or W. The center column—HAC Reduction Program—uses PSIs and HAIs to put your hospital at risk of a 1% CMS penalty if your hospital is in the lowest 25%. The right column—Value Based Purchasing—uses mortality metrics O/E combined with HAIs to put your hospital at risk new to a 2% penalty (or bonus!). *Penalties and bonuses are based on your documentation! Get credit for the high-quality care you provide!* 

| Hospital Acquired Conditions (HAC)<br>Deficit Reduction Act (DRA)   | Hospital Acquired Condition (HACs) Reduction Program<br>(HACRP) |  |   | Hospi                                     | tal Value-Based Purchasing (HVBP)   | 14       |                   |            |
|---|---|--|---|---|---|----------|-------------------|------------|
|   | DOMAIN 1:   | CMS Recalibrated PSI-90 (Composite of below measures) 15%  |   | DOMAIN:                                   | MORTALITY 25  | 86       |                   |            |
| HAC-1_Foreign Object Retained After Surgery   | P5I 03  | Pressure Ulcer   |   | MEASURE ID                                | Measure Description   |          |                   |            |
| HAC-2_Air Embolism  | P51 06  | latrogenic Pneumothorax  |   |   |   |          | 2019              | 2020 20    |
| HAC-3 Blood Incompatibility   | P51 08  | In Hospital Fall with Hip Fracture   |   | MORT - 30-AMI                             | Acute AMI 30-day mortality rate   |          | yes               | yes ye     |
| HAC-4 Stage III and IV Pressure Ulcers  | PSI 09  | Perioperative Hemorrhage or Hematoma   |   | MORT-30-HF                                | Heart Failure (HF) 30-day mortality rate  |          | yes               | yes ye     |
| HAC-5_Falls and Treume  | PSI 10  | Postoperative Acute Kidney Injury Requiring Dialysis   |   | MORT-30-PN                                | Pneumonia 30-day mortality rate (Incl. se   | psis)    | yes               | yes n      |
| HAC-6 Catheter-Associated Uninary Tract Infection (CAUTI)   | P5I 11  | Postoperative Respiratory Failure  |   |   |   |          |                   |            |
| HAC-7 Vascular Catheter-Associated Infection  | PSI 12  | Perioperative Pulmonary Embolism or Deep Vein Thrombosis   |   |   |   |          |                   |            |
| HAC-8_Surgical Site Infection, Mediastinitis, Following CABG  | PSI 13  | Postoperative Sepsis   |   | Complications                             |   |          |                   |            |
| HAC-9 Manifestations of Poor Glycemic Control   | PSI 14  | Postoperative Wound Dehiscence   |   | тна/тка                                   | Total Hip/Knee Arthroplasty Complication  |          | yes               | yes ye     |
| HAC-10 DVT/PE Following Certain Orthopedic Procedures   | PSI 15  | Unrecognized Abdominopelvic Accidental Puncture/Laceration   | 1 |   |   |          |                   |            |
| HAC-11 Surgical Site Infection Following Bariatric Surgery for Obesity  |   | Charts abstracted and reported to NHSN   | 1 |   |   |          |                   |            |
| HAC-12 Surgical Site Infection Following Certain Ortho Procedures   |   | CDC National Healthcare Safety Network (NHSN) 85%  | 1 | DC NN:                                    | PATIENT SAFETY 25%  | 8        |                   |            |
| HAC-13 SSI Following Cardiac Implantable Electronic Device (CIED)   | DOMAIN 2:   | HEALTHCARE ASSOCIATED INFECTIONS   |   | MEASO ID                                  | Measure Description   | 1        | 2019              | 20 20      |
| HAC-14 latrogenic Pneumothorax with Venous Catheterization  | HAI-1   | Central Line-Associated Bloodstream Infection (CLABSI)   | + | HAJ                                       | CLABSI  |          | yes               | 195 yr     |
|   | HAI-2   | Catheter-Associated Urinary Tract Infection (CAUTI)  | + | HAI-2                                     | - UTI   |          |                   | yes ye     |
|   | HAI-3   | SSI Colon Surgery  | + | HAI-3                                     | SSI C. SPECY  |          | yes               | yes ye     |
|   | HAI-4   | SSI Abdominal Hysterectomy   | + | HAI-4                                     | SSI Abdominal Hystereccomy  |          | yes               | yes ye     |
|   | HAI-5   | Methicillin-resistant Staphylococcus aureus bacteremia (MRSA)  | + | HAI-5                                     | MRSA  |          | yes               | yes ye     |
|   | HAI-6   | Clostridium difficile Infection (C.diff)   | + | HAI-6                                     | CDI (C.diff)  |          | yes               | yes ye     |
|   |   | FINANCIAL IMPACT   |   | <u>80</u>                                 |   |          |                   | 117        |
| Ider the DRA HAC payment provision, hospitals no longer receive additional<br>system for cases in which one of the selected conditions occurred but was not<br>esent on admission (IPOA). That is, the case is paid as though the condition(s) were   | Reduces hos<br>25% (started                                     | pital payments by 1% for hospitals that rank among the lowest-performing<br>in 2015 - part of Affordable Care Act) |   | The Hospital VBP P<br>hospitals' base ope | rogram is funded by reducing participating<br>trating Medicare severity MS-DRG payments | s by     |                   |            |
| It prevent. The DRA HAC-POA payment provision is applicable for secondary<br>agnosis code reporting only, as the selected conditions are designated as a  | HVBP<br>(1) off ALL h   | ospital's base FY18 operating MS-DRG payment   |   | DRG                                       | DRG Description   | Relative | Base WL           | Reimbursem |
| explication or comorbidity (CC) or a major complication or comorbidity (MCC) when   | (2) incentive   | s distributed to top performers based on Total Performance Score (TPS)   |   | 280 ACUTE MYOCA                           | ARDIAL INFARCTION, DISCHARGED ALIVE W MCC   | 1.6577   | \$6,000.00        | \$9,946.   |
| ported as a secondary diagnosis. For the LMA HAL PUA payment provision, a<br>syment adjustment is only applicable if there are no other CC/MCC conditions   | (3) net resul   | t of (1) and (2) above applied to base operating MS-DRG payment (claim by  | 1 | 281 ACUTE MYOCA                           | ARDIAL INFARCTION, DISCHARGED ALIVE W CC  | 0.9848   | 56,000.00         | \$5,908    |
| and a second s | Lef mirm 1  |  |   |   |   |          | R.C. Annual Cont. | 64.00.0    |

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# Present On Admission (POA)

Present on admission is defined as present at the time the order for

inpatient admission occurs

Conditions that develop during an outpatient encounter are

considered as present on admission

- Emergency department
- Observation
- Outpatient surgery



### Present On Admission (POA)

- Principal Diagnosis
  - Must be present on admission POA
- CMS and Premier (Care Science QualityAdvisor) codes must be POA to risk adjust mortality calculation

| Indicator | ΡΟΑ                         |                |
|-----------|-----------------------------|----------------|
| Υ         | Yes                         |                |
| Ν         | No                          |                |
| U         | Unspecified                 | Designated NO  |
| W         | Clinically cannot determine | Designated YES |



#### **Uncertain Diagnoses**

If the diagnosis documented <u>at the time of discharge</u> is qualified as

"probable", "suspected", "likely", "questionable", "possible", or "still

to be ruled out", or other similar terms indicating uncertainty, code

the condition as if it existed or was established.



## 2 Midnight Rule

- On July 1, 2015, CMS released the updates to the "Two Midnight" rule.
- CMS emphasis on physician's medical judgment
- Physician or other practitioner must decide whether to admit as inpatient or treat as outpatient
- CMS observed a higher frequency of extended observation services
- Inpatient admissions will <u>generally</u> be payable under Part A if the admitting practitioner expected the patient to require a hospital stay that crossed two midnights and the medical record supports that reasonable expectation
- All treatment decisions for beneficiaries were based on the medical judgment of physicians
- CMS sought to respect the judgment of physicians



# **IPPS Ruling – 2 Midnights Rule**

#### Inpatient Status

- 1. Inpatient Order\*\*
- 2. Expectation of hospitalization crossing "2 Midnights" of time
- 3. Medical Necessity (what is that?)



#### **Medical Necessity**

- "...in order for payment to be provided under Medicare Part A, the care
  - must be reasonable and necessary."

- "The factors that lead a physician to admit a particular beneficiary based
  - on the physician's clinical expectation...must be clearly and completely

documented in the medical record."



### **Medical Necessity**

#### **Medical Necessity Buzz Words**

Support Inpatient Status

- Acute
- Acute on chronic
- Decompensated
- Exacerbation
- Worsening
- Failed outpatient treatment

- Patient is immunocompromised
- The CURB-65 Score is...
- The Pneumonia Severity Index is. . .
- The TIMI or HEART Score is . . .
- The SOFA Score is . . .



# **Myocardial Infarction**



# First Question: What's the Correct Status?



#### Medical Necessity – Should the patient be INPATIENT?

Inpatient admissions will <u>generally</u> be payable under Part A if the admitting practitioner expected the patient to require a hospital stay that crossed two midnights and the medical record supports that reasonable expectation

 All treatment decisions for beneficiaries were based on the medical judgment of physicians

CMS sought to respect the judgment of physicians



#### **Medical Necessity**

The care must be *reasonable and necessary* 

• Must be *clearly and completely documented* in the medical record



## Chest Pain isn't simply Chest Pain

| Documentation                           | MS-DRG | Title   | RW     | Example<br>Reimbursement |
|---|--------|---|--------|--------------------------|
| Angina pectoris due to<br>ASCAD         | 303    | Atherosclerosis w/o MCC                                 | 0.6656 | \$4568                   |
| Pleurisy                                | 195    | Simple Pneumonia w/o CC                                 | 0.6868 | \$4714                   |
| Angina pectoris NOS                     | 311    | Angina Pectoris   | 0.6872 | \$4716                   |
| Non-<br>cardiac/musculoskeletal<br>pain | 313    | Chest Pain  | 0.7073 | \$4854                   |
| Pericarditis                            | 316    | Other Circ System Dx w/o CC                             | 0.7513 | \$5156                   |
| Heartburn / GERD                        | 392    | Esophagitis, Gastroenteritis and Misc GI<br>d/o w/o MCC | 0.7554 | \$5184                   |
| Pleuritic (not chest wall)<br>pain      | 204    | Respiratory Signs and Symptoms                          | 0.7676 | \$5268                   |
| Biliary colic                           | 446    | Disorder of Biliary Tract w/o CC/MCC                    | 0.7950 | \$5456                   |
| Costochondritis or rib<br>fracture      | 206    | Other Resp Dx w/o CC/MCC                                | 0.8635 | \$5927                   |
| Thoracic radiculopathy                  | 552    | Medical Back d/o w/o MCC                                | 0.9010 | \$6184 <sup>19</sup>     |

#### Fourth Universal Definition of Acute Myocardial Infarction 2018

- Myocardial infarction (MI) is acute myocardial injury detected by abnormal cardiac biomarkers in the setting of evidence of acute myocardial ischemia
  - Detection of a rise and/or fall of troponin with at least one value above the 99<sup>th</sup> percentile upper reference limit (URL) and with at least one of the following:
    - Symptoms of acute ischemia
    - New ischemic EKG changes
    - Development of pathological Q waves
    - Imaging evidence of new loss of viable myocardium or new regional wall motion abnormality consistent with ischemia
    - Identification of an intracoronary thrombus by angiography or autopsy



Fourth Universal Definition of Acute Myocardial Infarction 2018

Acute myocardial injury

 $_{\circ}$  20% rise or fall of cardiac troponin over time

Chronic myocardial injury

 $_{\odot}$  <20% rise or fall of cardiac troponin over time

• CKD

• Structural Heart Disease



## **Clinical Myocardial Ischemia**

Symptoms

- Angina (chest pain, jaw pain, left shoulder/arm pain)
- Angina equivalents (SOB, fatigue)
- Syncope (often due to arrhythmia)
- Flash pulmonary edema (not gradual decompensation of chronic hear t failure)
- Palpitations / Cardiac arrest

#### "OR EVEN WITHOUT SYMPTOMS"



https://www.acc.org/latest-in-cardiology/articles/2018/11/16/09/06/fourth-universal-definition-of-mi

# Type 1 MI

 Atherothrombotic coronary artery disease (CAD) and usually precipitated by atherosclerotic plaque disruption (rupture or erosion) is designated as a type 1 MI

Myocardial Infarction Type 1





Plaque rupture/erosion with occlusive thrombus



Plaque rupture/erosion with non-occlusive thrombus



## NSTEMI is a Type 1?

Let the controversy begin!

Taken directly from the 4<sup>th</sup> Universal Definition

New ST-segment elevations in two contiguous leads or new bundle branch blocks with ischaemic repolarization as an ST-elevation MI (STEMI)

In contrast, patients without ST segment elevation at presentation are usually designated non-ST-elevation MI (NSTEMI)



# MI type 2

Detection of a rise and/or fall of troponin with at least one value above the 99<sup>th</sup> percentile upper reference limit (URL) and evidence of an **imbalance** between myocardial oxygen supply and demand unrelated to CAD requiring at least one of the following:

- Symptoms of acute ischemia
- New ischemic EKG changes
- Development of pathological Q waves
- Imaging evidence of new loss of viable myocardium or new regional wall motion abnormality consistent with ischemia
- Identification of an intracoronary thrombus by angiography or autopsy
- MI type 2 has a new code as of October 2017
  - MI due to Demand Ischemia
  - MI due to Ischemic Imbalance



# MI type 2

- The pathophysiological mechanism leading to ischaemic myocardial injury in the context of a mismatch between oxygen supply and demand has been classified as type 2 MI
- By definition, acute atherothrombotic plaque disruption is not a feature of type 2 MI
   Myocardial Infarction Type 2





Atherosclerosis and oxygen supply/demand imbalance



Vasospasm or coronary microvascular dysfunction



Non-atherosclerotic coronary dissection



Oxygen supply/demand imbalance alone



# MI type 2

#### Causes

- Fixed coronary atherosclerosis
- Coronary spasm
- Coronary embolism
- Coronary artery dissection
- Sustained tachyarrhythmia
- Severe hypertension / LV hypertrophy
- Severe Bradyarrhythmia
- Respiratory failure
- Severe anemia
- Hypotension / Shock



# Ok, what is "NSTEMI Type 2"?

Depends on who you ask

- Kennedy/Goyal (ACC)
  - NSTEMI type 2 is conflicting documentation because all NSTEMI are type 1
  - Remember the ACC is physician opinion which carries limited weight with the 4 cooperating parties

Huff/Huff

 $_{\circ}\,$  NSTEMI encompasses all MI that do not raise the ST segments

Coding NSTEMI type 2 codes to MI type 2 (I21.A1) -- Phew!



https://www.acc.org/latest-in-cardiology/articles/2018/11/16/09/06/fourth-universal-definition-of-mi

#### **4 Cooperating Parties**

- CDC responsible for diagnoses (the government)
- CMS responsible for inpatient procedures (the government)
- American Hospital Association responsible for interpreting ICD-10

   Through Coding Clinic
- American Health Information Management Association (AHIMA)
  - Provides input from the coding community

#### Do you see the American College of Cardiology on that list?

# "NSTEMI Type 2"

**Coding Clinic, Q4 2017**, page 12 "Types of Acute Myocardial Infarction":

**Question**: How should a type 2 NSTEMI due to demand ischemia be coded?

<u>Answer</u>: Assign code I21.A1, Myocardial infarction type 2. Do not assign code I24.8, Other forms of acute ischemic heart disease for the demand ischemia. Code also the underlying cause, if known. According to the *ICD-10-CM Official Guidelines for Coding and Reporting*, "When a type 2 AMI code is described as NSTEMI or STEMI, only assign code I21.A1. Codes I21.01-I21.4 should only be assigned for type 1 AMIs."



https://www.acc.org/latest-in-cardiology/articles/2018/11/16/09/06/fourth-universal-definition-of-mi

## **Physician Advisors**

#### • Time to change HATS!





#### **AMI Mortality Cohort**

| Measure:        | 30-Day Risk-Standardized Mortality Rate Following AMI   |    |  |  |  |  |
|-----------------|---|----|--|--|--|--|
| Dx Inclusion:   | <ul> <li>Principal diagnosis of AMI (Excluding Type 2 MI)</li> <li>Not transferred from another acute care facility</li> <li>Age 65 or over</li> <li>Enrolled in Medicare FFS 12 months prior to index admission or VA beneficiary</li> </ul>   |    |  |  |  |  |
| Exclusions:     | <ul> <li>Discharged alive same day/next day, not transferred to another acute care facility</li> <li>Enrolled in <u>Medicare hospice</u> program or used VA hospice services any time in the 12 months prior to the index admission (including first day of the index admission)</li> <li>Discharged AMA</li> </ul> |    |  |  |  |  |
| Risk Variables: | <ul> <li>Anterior myocardial infarction (index admission only)</li> <li>Other (non-anterior) location of myocardial infarction (index admission only)</li> <li>History of CABG surgery</li> <li>History of PTCA</li> <li>25 condition categories</li> </ul>   | 32 |  |  |  |  |

#### **Acute Myocardial Infarction Metrics**

| ICD-10 code | Description   | CC/MCC | HCC | CMS 30d<br>Mortality |
|-------------|---|--------|-----|----------------------|
| R79.89      | Other specified abnormal findings of blood chemistry (troponin elevation) |        |     | NO                   |
| 121.4       | NSTEMI (Type 1 MI)  | MCC    | 86  | YES                  |
| 121.3       | STEMI of unspecified site   | MCC    | 86  | YES                  |
| 121.9       | AMI, unspecified  | MCC    | 86  | YES                  |
| I21.A1      | MI type 2 (due to demand ischemia)  | MCC    | 86  | NO                   |
| I21.A9      | Other MI type (3,4,5)   | MCC    | 86  | NO                   |
| 124.8       | Demand ischemia   | CC     | 87  | NO                   |
| 124.9       | Acute ischemic heart disease (ACS)  |        | 87  | NO                   |
| 120.0       | 0 Angina, unstable  |        | 87  | NO                   |
| 151.81      | Takotsubo Syndrome  | CC     |     | NO                   |
| No code     | Non-traumatic Acute Myocardial Injury                                     |        |     |                      |

# Pneumonia



# First Question: What's the Correct Status?



#### Medical Necessity – Should the patient be INPATIENT?

- Inpatient admissions will <u>generally</u> be payable under Part A if the admitting practitioner expected the patient to require a hospital stay that crossed two midnights and the medical record supports that reasonable expectation
- All treatment decisions for beneficiaries were based on the medical judgment of physicians
- CMS sought to respect the judgment of physicians



#### **Medical Necessity**

- The care must be *reasonable and necessary*
- Must be *clearly and completely documented* in the medical record



#### Pneumonia

Should this patient be inpatient?

- Clinical indicators such as SOB, Fever, Cough
- Infiltrate on CXR
- Abnormalities on Physical Examination
  - o Did anyone do a physical examination? Did anyone document it?
    - Crackles
    - Egophony
    - Tactile fremitus
    - Bronchial breath sounds
- Failed outpatient antibiotics



#### Pneumonia

#### CURB 65

- Confusion
- Uremia (BUN >19)
- Respiratory Rate > 30
- Blood Pressure SBP < 90 or DBP < 50</p>
- Age > 65



#### Pneumonia Severity Index

Demographic: Age/Sex/Nursing

Home Resident

- Neoplastic disease/Liver
- disease/Heart Failure/CVA

hx/Renal disease



AMS

- Respiratory Rate > 30
- SBP < 90
- Temp <  $95^{\circ}$  or >  $103.8^{\circ}$
- Pulse > 125
- pH < 7.35
- BUN > 30
- Sodium < 130</li>
- Glucose > 250
- Hematocrit < 30</p>
- PaO2 < 60</p>
- Pleural Effusion

# Do you have the right diagnosis?



#### Pneumonia as Principal Diagnosis

#### Will bucket into DRG 195 Simple Pneumonia or DRG 177 Respiratory Diseases

| Clinical to Coding                       |   |   |   |  |   |  |
|--|---|---|---|--|---|--|
| <u>S</u>                                 | imple Pneumoni  | ia  | Complex Pneumonia   |  |   |  |
| Simple                                   | Pneumonia and   | Pleurisy  | Respiratory I   | nfections and In   | flammations   |  |
| 195                                      | 194   | 193   | 179   | 178  | 177   |  |
| No CC/MCC                                | w CC  | w MCC   | No CC/MCC   | w CC   | w MCC   |  |
| 0.6868                                   | 0.9002  | 1.3167  | 0.9215  | 1.2744   | 1.8408  |  |
| 2.6                                      | 3.3   | 4.2   | 3.2   | 4.3  | 5.5   |  |
| Influenza PNA Viral PNA like adenoviral, |   |   | Influenza PNA w/specified secondary PNA   |  |   |  |
| unspecified Strep, H. flu; Mycoplasma,   |   |   | Tuberculous I   | ungal Virulent c   | organisms like  |  |
| Chlamydial BronchoPNA; Lobar PNA J18.9   |   |   | CMV; RSV;   | K. pneumo; Sta   | ph; E. coli;  |  |
| Pneumonia, unspecified HCAP!             |   |   | Legionnaire   | s';Gram negativ  | e Aspiration  |  |
|  |   |   | P   | ulmonary Absce   | SS  |  |
|  | Simple<br>Simple<br>195<br>No CC/MCC<br>0.6868<br>2.6<br>Influenza PI<br>unspecified<br>Chlamydial B<br>Pneum | Simple PneumoniaSimple Pneumonia and195194No CC/MCCw CC0.68680.90022.63.3Influenza PNA Viral PNA likeunspecified Strep, H. flu; MChlamydial BronchoPNA; LotPneumonia, unspecified | Clinical tSimple Pneumonia and Pleurisy195194193No CC/MCCw CCw MCC0.68680.90021.31672.63.34.2Influenza PNA Viral PNA like adenoviral,<br>unspecified Strep, H. flu; Mycoplasma,<br>Chlamydial BronchoPNA; Lobar PNA J18.9<br>Pneumonia, unspecified HCAP! | Clinical to CodingSimple Pneumonia and PleurisyCodSimple Pneumonia and PleurisyRespiratory I195194193195194193No CC/MCCw CCw MCCNo CC/MCCw CCw MCC0.68680.90021.31672.63.34.21nfluenza PNA Viral PNA like adenoviral,<br>unspecified Strep, H. flu; M∠coplasma,<br>Chlamydial BronchoPNA; Lobar PNA J18.9<br>Pneumonia, unspecified HCAP!Influenza PNA<br>Legionnaire<br>PNA | Clinical to CodingSimple Pneumonia and PleurisyComplex PneumonSimple Neumonia and PleurisyRespiratory Infections and Inf195194193179195194193179No CC/MCCw CCw MCCNo CC/MCC0.68680.90021.31670.92151.27442.63.34.23.24.3Influenza PNA Viral PNA like adenoviral,<br>unspecified Strep, H. flu; Mcoplasma,<br>Chlamydial BronchoPNA; Lobr PNA J18.9<br>Pneumonia, unspecified HCAP!Influenza PNA w/specified set<br> |  |

#### **Uncertain Diagnoses**

If the diagnosis documented <u>at the time of discharge</u> is qualified as

"probable", "suspected", "likely", "questionable", "possible", or "still

to be ruled out", or other similar terms indicating uncertainty, code

the condition as if it existed or was established.



#### Pneumonia

- Probable gram-negative pneumonia, Rx Zosyn
- Probable MRSA pneumonia, Rx Vancomycin
- Suspected Aspiration pneumonia
  - Clindamycin or Flagyl Rx
- All below Map to the DRG for Simple Pneumonia
  - Community Acquired Pneumonia
  - Healthcare Associated Pneumonia (HCAP)
  - Nosocomial Pneumonia



#### Pneumonia for ICD-10

| ICD-10 code  | Description   | CC or MCC |  |  |  |
|--|---|-----------|--|--|--|
| J15.9  | Unspecified bacterial pneumonia                                       | MCC       |  |  |  |
| J18.9  | Pneumonia, unspecified organism<br>(includes CAP & HCAP & Nosocomial) | MCC       |  |  |  |
|  |   |           |  |  |  |
| J69.0  | Aspiration Pneumonia  | MCC       |  |  |  |
| J15.6  | Pneumonia due to gram negative bacteria                               | MCC       |  |  |  |
| J15.212  | Pneumonia due to MRSA   | MCC       |  |  |  |
| J15.8 Pneumonia due to specified bacteria (anaerobic) MCC      |   |           |  |  |  |
| <ul> <li>Simple pneumonia maps to DRG 195 Pneumonia</li> </ul> |   |           |  |  |  |
| Specified codes map to DRG 177 Resp Diseases                   |   |           |  |  |  |

# Do you have the right diagnosis?



# How many criteria for Sepsis are there?



# How many criteria for Sepsis are there?





### Sepsis: If "Some" are due to infection

#### SIRS criteria

- Altered Mental Status
- Significant edema or positive fluid balance
- Hyperglycemia in the absence of diabetes
- CRP more than two SD above the normal value
- Procalcitonin more than 2 SD above the normal value
- Hypotension (SBP < 90 mmHg or SBP decrease > 40 mmHg)
   Brundage Group EXCELLENCE IN REVENUE CYCLE

- Hypoxemia (PaO2/FiO2 < 300)</p>
- Acute oliguria (urine output < 0.5mL/kg/hr for 2 hours)</li>
- Creatinine increase >0.5mg/dL
- INR >1.5
- Ileus
- Thromobocytopenia (PLT < 100,000)</p>
- Hyperbilirubinemia (> 4 mg/dL)
- Hyperlactatemia (> 1 mmol/L
- Decreased capillary refill or mottling

# Other criteria are for <u>Severe</u> Sepsis



#### **Severe Sepsis**

#### Table 2. Terminology and International Classification of Diseases Coding

| Current Guidelines<br>and Terminology                         | Sepsis  | Septic Shock   |
|---|---|--|
| 1991 and 2001<br>consensus<br>terminology <sup>9,10</sup>     | Severe sepsis<br>Sepsis-induced<br>hypoperfusion  | Septic shock <sup>13</sup>   |
| 2015 Definition   | Sepsis is<br>life-threatening organ<br>dysfunction caused by a<br>dysregulated host<br>response to infection  | Septic shock is a subset of<br>sepsis in which underlying<br>circulatory and<br>cellular/metabolic<br>abnormalities are profound<br>enough to substantially<br>increase mortality  |
| 2015 Clinical<br>criteria                                     | Suspected or<br>documented infection<br>and<br>an acute increase of ≥2<br>SOFA points (a proxy<br>for organ dysfunction)  | Sepsis <sup>a</sup><br>and<br>vasopressor therapy needed to<br>elevate MAP ≥65 mm Hg<br>and<br>lactate >2 mmol/L (18 mg/dL)<br>despite adequate fluid<br>resuscitation <sup>13</sup>   |
| Recommended<br>primary ICD<br>codes <sup>a</sup>              |   |  |
| ICD-9   | 995.92  | 785.52   |
| ICD-10 <sup>a</sup>   | R65.20  | R65.21   |
| Framework for<br>implementation<br>for coding and<br>research | Identify suspected infect<br>for blood cultures and an<br>specified period <sup>b</sup><br>Within specified period a<br>1. Identify sepsis by usin<br>life-threatening organ dy<br>2. Assess for shock criter<br>vasopressors, MAP <65 m<br>(18 mg/dL) <sup>d</sup> | ion by using concomitant orders<br>itibiotics (oral or parenteral) in a<br>round suspected infection <sup>c</sup> :<br>g a clinical criterion for<br>ysfunction<br>ia, using administration of<br>nm Hg, and lactate >2 mmol/L |



#### https://jamanetwork.com/journals/jama/fullarticle/2492881

#### **4 Cooperating Parties**

- CDC responsible for diagnoses (the government)
- CMS responsible for inpatient procedures (the government)
- American Hospital Association responsible for interpreting ICD-10

   Through Coding Clinic
- American Health Information Management Association (AHIMA)
  - Provides input from the coding community

#### Do you see the Society of Critical Care Medicine on that list?

#### Severe Sepsis-3

- New Terms and Definitions
- (Severe) Sepsis is defined as life-threatening organ dysfunction (not failure) caused by a dysregulated host response to infection.
- Organ dysfunction can be identified as an acute change in total SOFA score ≥2 points consequent to the infection.

 A SOFA score ≥2 reflects an overall mortality risk of approximately 10% in a general hospital population with suspected infection.

http://jama.jamanetwork.com/article.aspx?articleid=2492875



#### **Severe Sepsis Organ Failure Assessment**

|       |  | core – Acute change of 2 points due to the infectious process supports sepsis |                               |                                     |  |  |  |  |
|-------|--|---|-------------------------------|-------------------------------------|--|--|--|--|
|       | System   | 0   | 1                             | 2                                   | 3  | 4  |  |  |
|       | <b>Neurologic</b><br>GCS   | 15  | 13-14                         | 10-12                               | 6-9  | < 6  |  |  |
|       | <b>Respiratory</b><br>PaO <sub>2</sub> /FiO <sub>2</sub><br>RA PaO <sub>2</sub> , O <sub>2</sub> sat | <u>&gt;</u> 400<br><mark>84, 95%</mark>                                       | < 400<br><mark>84, 95%</mark> | < 300<br><mark>63,</mark> 91%       | < 200 with respiratory<br>support<br>42, 80%                                   | < 100 with respiratory<br>support<br>21, < 80%           |  |  |
| LYGII | Cardiovascular   | MAP <u>&gt;</u> 70 mmHg   | MAP < 70 mmHg                 | Dopamine < 5 or<br>Dobutamine (any) | Dopamine 5.1-15 or<br>Epinephrine <u>&lt;</u> 0.1 or<br>Norepi <u>&lt;</u> 0.1 | Dopamine > 15 or<br>epinephrine > 0.1 or<br>norepi > 0.1 |  |  |
|       | <b>Hepatic</b><br>Bilirubin, mg/dL   | < 1.2   | 1.2-1.9                       | 2.0-5.9                             | 6.0-11.9   | > 12.0   |  |  |
|       | <b>Coagulation</b><br>Platelets, x 1,000   | <u>&gt;</u> 150   | < 150                         | < 100                               | < 50   | < 20   |  |  |
|       | <b>Renal</b><br>Creatinine, mg/dL  | < 1.2   | 1.2-1.9                       | 2.0-3.4                             | 3.5-4.9  | > 5.0  |  |  |
| 200   | UOP, ml/d  |   |                               |                                     | < 500  | < 200  |  |  |

#### Abbreviations:

PaO<sub>2</sub>: partial pressure of oxygen; FiO<sub>2</sub>: fraction if inspired oxygen; MAP: Mean arterial pressure

Catecholamine doses are in mcg/kg/min for at least 1 hour.

## **Physician Advisors**

#### • Time to change HATS!





# Back to Pneumonia Do you have the right diagnosis?



### Pneumonia Cohort

| Measure:        | 30-Day Risk-Standardized Mortality Rate Following Pneumonia  |
|-----------------|--|
| Dx Inclusions:  | <ul> <li>Principal discharge dx of pneumonia <u>or</u></li> <li>Principal discharge dx of sepsis (Excluding severe sepsis)         <ul> <li>with a secondary dx of pneumonia POA (and NO secondary diagnosis of severe sepsis POA)</li> </ul> </li> <li>Not transferred from another acute care facility</li> <li>Age 65 or over</li> <li>Enrolled in Medicare FFS 12 months prior to index admission or VA beneficiary</li> </ul> |
| Exclusions:     | <ul> <li>Discharged alive same day/next day, not transferred to another acute care facility</li> <li>Enrolled in <u>Medicare hospice</u> program or used VA hospice services any time in the 12 months prior to the index admission (including first day of the index admission)</li> <li>Discharged AMA</li> </ul>  |
| Risk Variables: | <ul> <li>History of CABG surgery</li> <li>History of PTCA</li> <li>30 condition categories</li> </ul>  |

# If the patient is actually sick enough to meet medical necessity, then you meet "probably" Sepsis



# Timing of the Diagnosis

- H&P: "Probable Sepsis"
  - $_{\circ}$  Diagnosis made POA
    - ER
    - Hospitalist, especially Nocturnist
    - Resident
  - $_{\circ}$  Will likely be the Principal Dx and drive the DRG
  - $_{\circ}$  Will risk adjust to Sepsis
  - $_{\rm \circ}\,$  Will risk adjust to Severe Sepsis
    - IF YOU WANT TO AVOID A DENIAL



#### Severe Sepsis – SEP 1 Core Measure

The timing of the diagnosis is critical

SEP-1

- Severe Sepsis documented or ABSTRACTED
- Severe Sepsis ABSTRACTED all three within six hours of one another
- a) Documentation of any (bacterial) infection
- b) 2 or more SIRS Criteria
- c) Organ Dysfunction



## **Severe Sepsis**

| Sepsis-3 Publication     | <u>Severe Sepsis 2012</u>               |
|--------------------------|---|
| Sepsis = Severe Sepsis   | Sepsis induced hypotension              |
| All SOFA = Severe Sepsis | Lactic Acid > 2                         |
|                          | Urine Output < 0.5 mg/kg/hr for 2 hours |
|                          | w/ fluids                               |
|                          | ALI w PaO2/FiO2 < 250 w/o PNA           |
|                          | ALI w PaO2/FiO2 < 200 w PNA             |
|                          | Creatinine > 2                          |
|                          | Bilirubin > 2                           |
|                          | Platelets < 100k                        |
|                          | Coagulopathy INR > 1.5                  |

## Sepsis Clinical to Coding

| Clinical to Coding     |                      |
|------------------------|----------------------|
| Sepsis-3 Publication   | Sepsis ICD 10 Coding |
|                        |                      |
| Sepsis = Severe Sepsis | Sepsis               |
|                        | Severe Sepsis        |
|                        | Septic Shock         |



#### Summary: Sepsis Clinical to Quality

**Mortality Measures** 

Sepsis POA and Pneumonia POA bucket into Pneumonia Mortality

Severe Sepsis POA and Pneumonia POA bucket into Severe Sepsis

Mortality



# Physician Advisor Knowledge



#### **Physician Advisor Skill Set**





# Thank you. Questions?

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