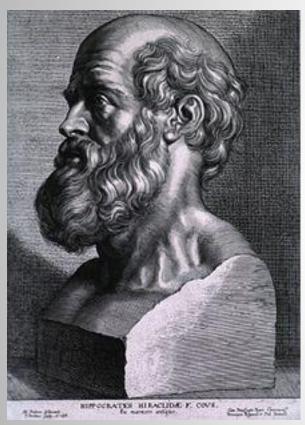
Nosocomial Infections

7/25/18 Noon Conference
Dan Van Aartsen
PGY3 Internal Medicine

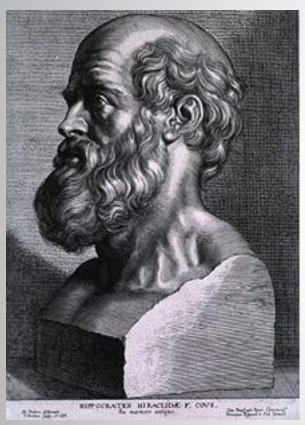
Learning Objectives

- Define Nosocomial Infections
- Identify common hospital acquired infections
- Know the common causes and understand basic pathophysiology of nosocomial infections
- Learn the fundamentals of prevention of hospital acquired infections and basic management



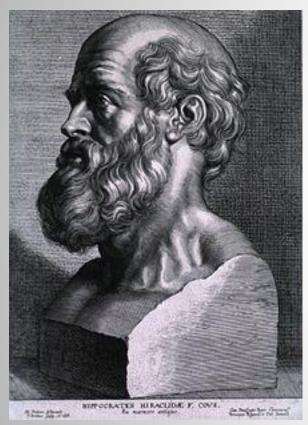
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 - nósos (disease, illness) + koméō ("to take care of") (nosokomeíon = "hospital")



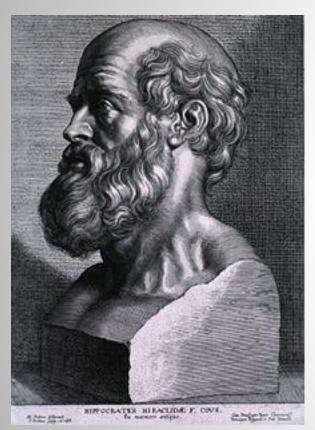
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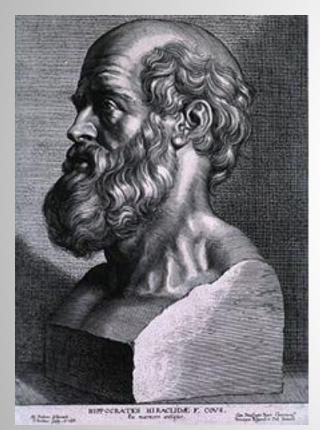
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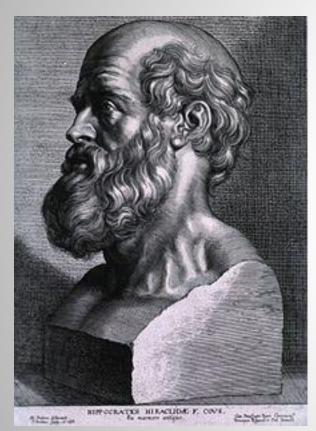
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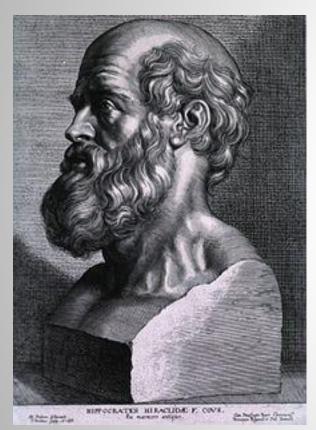
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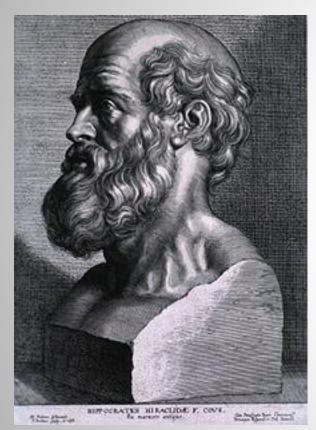
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 - This can be important! Some treatment guidelines are based on community-acquired vs. hospital acquired



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- Definition?? Not universally agreed upon
 - "An infection occurring in a patient during the process of care in a hospital or other health care facility which was not present or incubating at the time of admission." (WHO)



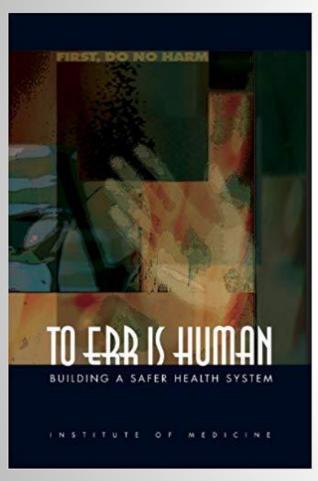
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 - "...This includes infections acquired in the hospital, but appearing after discharge, and also occupational infections among staff of the facility."

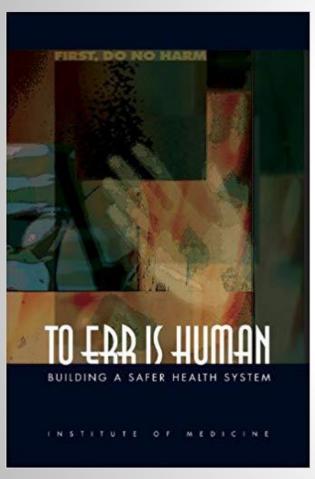
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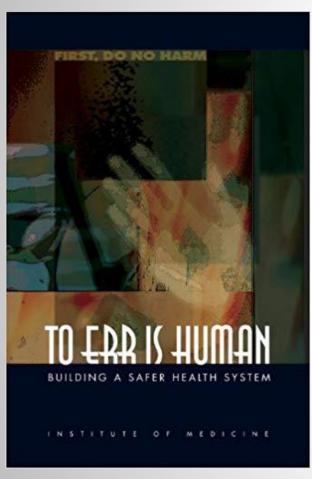
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- Three landmark events ignited the field of infection control...



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 - Focused on preventable medical errors of all types, but popularized the concept of *preventable* hospital-acquired infections



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 - "Deaths linked to hospital germs represent the fourth leading cause of mortality among Americans, behind heart disease, cancer and strokes, according to the federal Centers for Disease Control and Prevention. These infections kill more people each year than car accidents, fires and drowning combined."
 - Brought mainstream attention to HAI and forced action within the health care community

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 - Paradigm shift from infection control to infection prevention

The most commonly seen healthcare-associated infections?

Catheter-Associated Urinary Tract Infections

- Catheter-Associated Urinary Tract Infections
- Central Line-Associated Bloodstream Infections

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HAI

- CAUTI
- CLABSI
- HAP/HCAP
- VAP
- SSI
- Cdiff

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- CLABSI
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 - And don't check unless there's symptoms

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- Discontinue catheter as soon as it is not required
- Consider alternatives to indwelling catheters, (eg condom catheter, intermittent catheterization, suprapubic catheterization)

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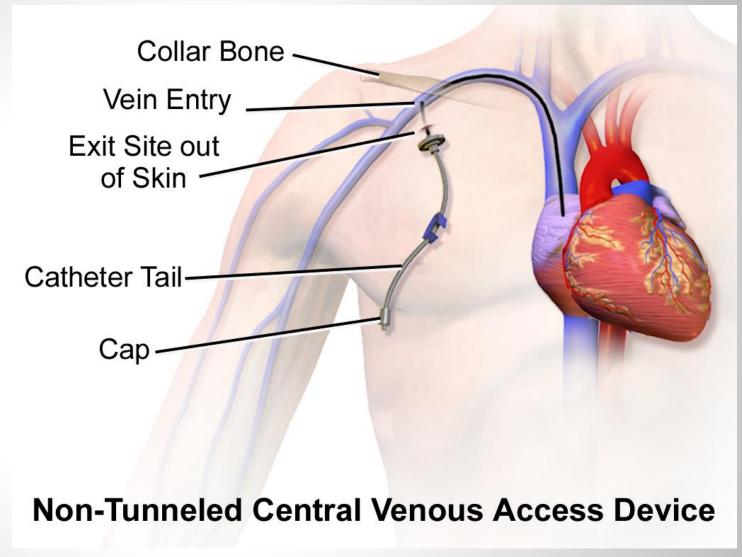
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 - Typically can be PO if tolerated

CLABSI



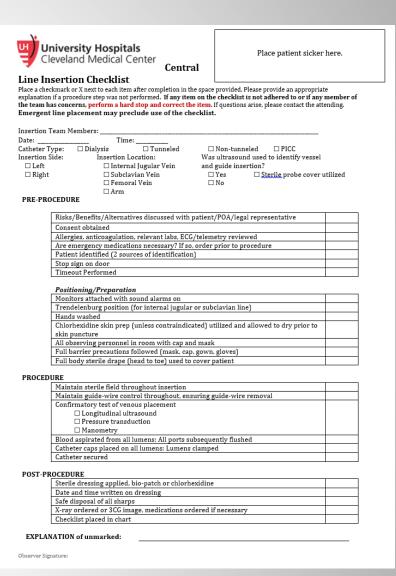
Blausen.com staff (2014). "Medical gallery of Blausen Medical 2014". WikiJournal of Medicine 1 (2). DOI:10.15347/wjm/2014.010. ISSN 2002-4436. -

CLABSI

- "CLABSI" only includes central lines, but infections can be also be associated with peripheral IVs
 - tunneled lines, PICCs, ports, etc
- Suspect when bloodstream infection occurs with no other apparent source
- Staph aureus, CONS, Candida bacteremia in the absence of other sources should raise suspicion
 - Increasingly multi-drug resistant gram negative bacteria

CLABSI – Prevention

- Many "Bundles" have been shown to reduce CLABSI rates
 - Standardized procedures, checklists, educational training, maximal sterile barriers
- MICU initiative to reduce CLABSI
- (remove lines when they're not needed!!)



CLABSI - Management

- Remove the line!
 - Possible exceptions include CONS, +/- enterococcus and GNBs if stable
- Obtain catheter cultures (but only if infection is suspected)
- Also get peripheral cultures
- If CONS grow keep checking
- Antibiotics
 - Vancomycin (high MRSA rates)
 - Empiric Gram negative coverage IF critically ill, neutropenic, femoral catheter, or known GN infection
 - Tailor to patient and local susceptibilities (here Zosyn)
 - Also cover Candida species if femoral line is suspected source

HCAP/HAP/Nosocomial PNA, VAP

HAP/VAP

- HAP = pneumonia that occurs ≥48 hours after admission
- VAP = pneumonia that occurs ≥48 hours after intubation
- Suspect in patients with new infiltrates, increasing O2 or ventilator requirements, fevers, chills, etc
- MRSA, Pseudomonas, Acinetobacter, Klebsiella
- Resistance is common
 - Risk factors for MDR organisms: recent antibiotics, prolonged hospitalization, poor functional status, hemodialysis, severe illness

HAP/VAP - Management

- Establish microbiologic diagnosis!
 - Blood and respiratory cultures indicated in all patients
 - Consider Pneumococcal and Legionella urine ag testing
 - Viruses can account for up to 1/3 of severe pneumonia even hospital acquired
 - Deep respiratory cultures may be necessary

Antibiotics

 Depends MRSA and MDR risk factors – empiric coverage may entail simple broad spectrum respiratory fluorquinoline up to double antipseudomonal + MRSA coverage

The recommendations in this algorithm are generally in keeping with the 2016 IDSA/ATS guidelines for the management of HAP and VAP. These regimens are intended for the initial treatment of patients in whom the microbiologic cause has not yet been identified. The doses below are intended for patients with normal renal function; dosing will need to be adjusted for patients with renal dysfunction. Empiric treatment choices should be influenced by the local distribution of pathogens causing VAP and their antimicrobial susceptibility patterns (ideally using an antibiogram that is specific to the hospital's ICU population). Antimicrobial selection should also be based upon the patient's risk factors for MDR pathogens, including recent antibiotic therapy, the presence of underlying diseases, and available culture data (including prior microbiology data). Additional considerations include potential toxicities, potential drug interactions, cost, availability, and clinician familiarity with the medications. Once the results of pretherapy cultures are available, therapy should be narrowed based upon the susceptibility pattern of the pathogens identified and the potential toxicities of the regimens. Are any of the following risk factors for MDR VAP present? ■ IV antibiotic use within the previous 90 days? Septic shock at the time of VAP ARDS preceding VAP ■ ≥5 days of hospitalization prior to the occurrence of VAP Acute renal replacement therapy prior to VAP onset Yes Does the patient have either of the following risk factors for resistant gram-negative bacilli? ■ Treatment in a unit in which >10% of gram-negative bacilli are resistant to an agent being considered for monotherapy Treatment in a unit in which the prevalence of resistance among gram-negative bacilli is unknown Yes Does the patient have either of the following risk factors for MRSA? Does the patient have either of the following risk factors for MRSA? ■ Treatment in a unit in which >10 to 20% of Staphylococcus aureus ■ Treatment in a unit in which >10 to 20% of S. aureus isolates are isolates are methicillin-resistant methicillin-resistant ■ Treatment in a unit in which the prevalence of MRSA is not known ■ Treatment in a unit in which the prevalence of MRSA is not known Nο Nο Yes One of the following: A One of the following: One of the following: One of the following: * ■ Piperacillin-tazobactam 4.5 g IV every 6 hours¶ ■ Piperacillin-tazobactam 4.5 g IV every 6 hours¶◊ ■ Piperacillin-tazobactam 4.5 g IV every 6 hours¶ ■ Piperacillin-tazobactam 4.5 g IV every 6 hours¶◊ ■ Cefepime 2 g IV every 8 hours¶ ■ Cefepime 2 g IV every 8 hours¶ Cefepime 2 q IV every 8 hours ¶ ■ Cefepime 2 g IV every 8 hours¶ ■ Imipenem 500 mg IV every 6 hours¶ ■ Ceftazidime 2 g IV every 8 hours ■ Levofloxacin 750 mg IV every 24 hours ■ Ceftazidime 2 g IV every 8 hours ■ Meropenem 1 g IV every 8 hours¶ ■ Imipenem 500 mg IV every 6 hours¶ ■ Levofloxacin 750 mg IV every 24 hours ■ Meropenem 1 q IV every 8 hours¶ ■ Ciprofloxacin 400 mg IV every 8 hours Plus one of the following: * † ■ Aztreonam 2 g IV every 8 hours ΔΔ Aztreonam 2 g IV every 8 hours Amikacin 15 to 20 mg/kg IV daily ¥** ■ Gentamicin 5 to 7 mg/kg IV daily ¥** Plus one of the following: # † Plus one of the following: § ■ Tobramycin 5 to 7 mg/kg IV daily ¥** ■ Amikacin 15 to 20 mg/kg IV daily ¥** ■ Vancomycin 15 mg/kg IV (maximum 2 g per dose initially) ■ Gentamicin 5 to 7 mg/kg IV daily¥** every 8 to 12 hours with goal to target 15 to 20 mcg/mL Levofloxacin 750 mg IV every 24 hours trough level; consider a loading dose of 25 to 30 mg/kg ■ Tobramycin 5 to 7 mg/kg IV daily¥** Ciprofloxacin 400 mg IV every 8 hours (maximum 3 g) x 1 for severe illness ♦¥ Aztreonam 2 g IV every 8 hours ¶¶ Levofloxacin 750 mg IV every 24 hours Linezolid 600 mg IV every 12 hours ■ Ciprofloxacin 400 mg IV every 8 hours Aztreonam 2 g IV every 8 hours (if not selected for the first agent above) ¶¶ Plus one of the following: 5 Vancomycin 15 mg/kg IV (maximum 2 g per dose initially) every 8 to 12 hours with goal to target 15 to 20 mcg/mL trough level; consider a loading dose of 25 to 30 mg/kg

(maximum 3 g) x 1 for severe illness ♦¥

Linezolid 600 mg IV every 12 hours

Prevention of HAI

- "Standard Precautions"
 - HAND HYGIENE HAND HYGIENE HAND HYGIENE
 - Gloves when touching blood, body fluids, secretions, etc,
 - Injection safely
- Sterile technique
- Avoid unnecessary medical devices
- Remove unneeded medical devices
- Isolation Precautions

Who, why, when, how??

- Who, why, when, how??
- 7 types of transmission based precautions at UH:
 - Contact
 - Contact plus
 - Airborne
 - Droplet
 - Protective
 - Special Alert Precautions
 - Special precautions: Droplet/Contact (peds only)
 - Combos of above

- Contact
 - Gown/gloves
 - MRSA
 - Highly resistant organisms
 - HSV, disseminated or severe
 - Major wounds not contained by dressing
 - Others, eg head lice, viral conjunctivitis, etc

Contact Precautions

WASH OR SANITIZE HANDS TO ENTER AND EXIT ROOM:





EVERYONE MUST WEAR GLOVES TO ENTER:



EVERYONE MUST WEAR GOWN & GLOVES FOR PATIENT CONTACT:





FOR PATIENT TO LEAVE ROOM:

Body fluids must be containedWounds/rash must be covered

- Contact
- Contact plus
 - Gown/gloves + hand-washing
 - C diff
 - (also norovirus, cryptosporidium)



WASH OR SANITIZE HANDS TO ENTER



MUST WASH HANDS WITH SOAP AND WATER TO EXIT:

EVERYONE MUST WEAR GOWN & GLOVES TO ENTER:





FOR PATIENT TO LEAVE ROOM:

Check with nurse

- Contact
- Contact plus
- Airborne
 - N-95 Respirator
 - TB, Avian flu, others

Airborne Precautions

Keep Door(s) Closed

Negative Pressure Room



DO NOT ENTER

BEFORE CHECKING WITH NURSE



mask

WASH OR SANITIZE HANDS TO **ENTER AND EXIT ROOM:**





To ENTER this room Health Care Staff MUST wear:

Visitors: MUST wear surgical mask when in patient's room



FOR PATIENT TO LEAVE ROOM: Must wear surgical mask

· Cover rash if present

- Contact
- Contact plus
- Airborne
- Droplet
 - Mask
 - Influenza, H flu, bacterial meningitis

Droplet Precautions

WASH OR SANITIZE HANDS TO ENTER AND EXIT ROOM:





EVERYONE <u>MUST</u> WEAR MASK TO ENTER:



FOR PATIENT TO LEAVE ROOM:

Must wear mask

- Contact
- Contact plus
- Airborne
- Droplet
- Protective
 - For neutropenic patients
 - Mask only if respiratory symptoms
 - For patient protection, not prevention of transmission

Protective Precautions

WASH OR SANITIZE HANDS TO ENTER AND EXIT ROOM:





ANYONE WITH RESPIRATORY SYMPTOMS

MUST WEAR MASK TO ENTER:



FOR PATIENT TO LEAVE ROOM:

Must wear mask

- Contact
- Contact plus
- Airborne
- Droplet
- Protective
- Special Alert Precautions
 - Localized zoster shingles in an immunocompetent patient
 - Only providers with immunity can care for patient



- Contact
- Contact plus
- Airborne
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- Special precautions: Droplet/Contact (peds only)

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- Special precautions: Droplet/Contact (peds only)
- Many combos of above

Questions on Isolation? See UH Isolation policies (on intranet)

Viral Hemorrhagic Fevers (due to Lassa, Ebola, Marburg, Crimean-Congo fever viruses)	Contact and Airborne	Airborne Isolation requires negative pressure; HCW must wear N-95 respirator. Single patient room. Appropriate waste handling. Largest viral load in final stages of illness when hemorrhage may occur; additional PPE, including double gloves, leg and shoe covers may be used. Face and eye protection required. See www.cdc.gov for most current recommendations. Use "High Alert Precautions" sign (brown)
VRE (Vancomycin Resistant Enterococcus); NOT labelled as highly resistant. See also MDRO.	Standard	
Viral Respiratory Diseases (Adult)	Standard	
West Nile – See Arthropodborne viral encephalides		
Whooping cough – See Pertussis		
Wound infections		
Major – no dressing or not contained by dressing	Contact	Duration of illness
Minor – contained by dressing	Standard	
Yersinia – See Gastroenteritis		
Zika	Standard	
Zygomycosis (phycomycosis, mucormycosis)	Standard	Not transmitted person to person

Define Nosocomial Infections

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- Learn the fundamentals of prevention of hospital acquired infections and basic management

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Thanks!