

NHL-VS CAMPUS PROGRAM 2017

Dr. Juzar Ali. LSUHSC, New Orleans, LA, USA Dr. Sanjay Bhatt, LAC/USC, Los Angeles, CA, USA

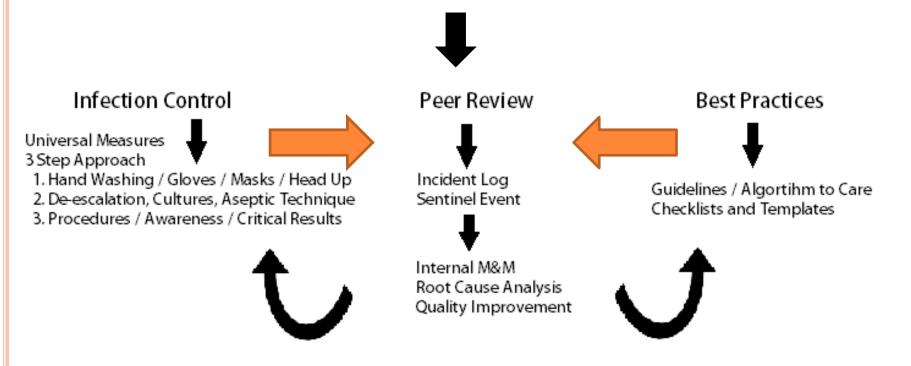
#### **OVERVIEW**

- Broad Outline & Recap to date
- Team Members / Team Work
- Departmental/Champions Updates
- Infection Control
- Peer Review
- Best Practices
- Closing

# BROAD OUTLINE TEAM MEMBERS

#### EQuIPS (Enhanced Quality Improvement and Patient Safety)

A Team Effort Between Doctors / Nurses / Allied Health Leadership Who Believe High quality safe patient-care can be provided in spite of a high demand, challenging environment and limited resources



Goals: Increase Awareness, Enhanced Patient Safety, Education, Process Improvement





#### TEAM

- DEAN / MS / HODs LEADERSHIP QUALITY COUNCIL
- Collaboration with:
  - Nursing
  - Allied Health
  - Dept. of Microbiology
  - Infection Control
  - Department of PSM
  - Others on invitation

- CQM Faculty Champions
  - Medicine: Dr. Palat
  - Medicine: Dr. Suthar
  - Surgery: Dr. Kushwala
  - Surgery: Dr. Vyas
  - OBG: Dr. Jani
  - EMD: Dr. Jarwani
  - Others on invitation

# CQM CHAMPIONS DEPARTMENTAL INPUT UPDATES MARCH 2017

#### FOCUS ONE

## INFECTION CONTROL

Introduction
Antibiogram
Antibiogram Observations
3 Step Approach

#### ANTIBIOGRAM

Dr. Parul Shah – Professor / Head of Microbiology Dr. Tanmay Mehta – Assistant Professor

#### ANTIBIOGRAM

- Information: Specimen Culture & Sensitivity Reports
  - % of samples for a given organism which were sensitive to certain antibiotics
- o Dates: 1st January 2016 31st December 2016

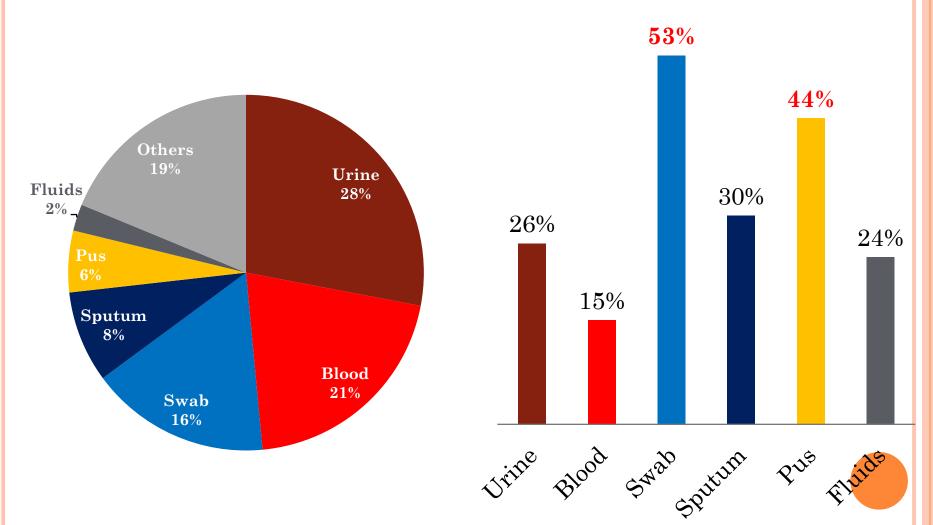
#### USE OF AN ANTIBIOGRAM

- Prepare an antibiotic policy
- To Initiate empirical treatment
- Detection of emergence of new hospital isolates
- Detection of changes in resistance patterns

Total Samples for Culture = **21631**Total No. of Isolates obtained = 6816 (**31.51**%)

**Samples for Culture** 

% Isolates from sample



## WHO'S "DIRTY DOZEN" JUST ANNOUNCED

SEE WHO Feb 2017 released report with regional data

#### WHO PRIORITY PATHOGENS

CRITICAL: Acinetobacter Baumanni – Carbp res

Pseudomonas Aeruginosa – Carbp res

Enterobacteriaceae Carbp' res ESBL

HIGH: Enterococcus, Staph MRSA, QR-

Salmonella, Hb/Cb/NG

MEDIUM: Strep /Hemoph/Shigella

## URINE

												U	Irine	isola	tes -	% St	ıscep	otible																				
Organism	% of isolates	Amikacin	Amoxicillin/Clavulinic acid	Ampicillin/Sulbactam	Azithromycin	Aztreonam	Cefepime	<b>Cefoperazone/Sulbactam</b>	Сеfotaxime	Ceftazidime	Ceftriaxone	Cefuroxime	Chloramphenicol	Ciprofloxacin	Clindamycin	Colistin	Doxycycline	Ertapenem	Erythromycin	Gatifloxacin	Gentamicin	Imipenem	Levofloxacin	Linezolid	Meropenem	Minocycline	Nitrofurantoin	Ofloxacin	Okacillin	PenicillinG	Piperacillin	Piperacillin/Tazobactam	Polymixin B	Teicoplanin	Tetracycline	Tobramycin	Trimethoprim/Sulfamethox	Vancomycin
GRAM-NEGATIVE ORGANISMS																																						
Escherichia coli	55%	77	-	45	-	1	36	68	-	-	20	-	-	21	-	100	25	74	-	49	56	73	22	-	73	25	77	-	-	-	-	55	100	-	-	-	28	-
Klebsiella pneumoniae	12%	58	-	30	-	-	32	60	-	-	21	-	-	28	-	100	29	41	-	48	53	56	28	-	56	29	46	-	-	-	-	79	100	-	-	-	29	-
Pseudomonas aeruginosa	7%	30	-	-	-	31	32	30	-	24	-	-	-	23	-	100	-	-	1	ı	33	35	25	-	35	-	-	33	-	-	25	40	99	-	-	26	-	-
Klebsiella sp.	4%	53	-	30	-	1	30	40	-	-	25	-	-	31	-	100	28	48	ı	39	48	48	34	-	48	28	44	-	-	-	-	-	100	-	-	-	33	-
GRAM-POSITIVE ORGANISMS																																						
Staphyloccocus aureus	2%	-	60	-	0	-	-	-	-	-	-	-	100	55	100	-	65	-	0	-	77	-	61	100	-	65	-	50	58	0	-	-	-	-	-	66	45	100

# Urine

Total = $6055$ with $1581$ (26.11%)	Isola	ites	100% Sensitive
Organism	No.	%	
Gram negative bacilli	Ì	77%	
Escherichia coli	870	<b>55%</b>	
		11.55	
Klebsiella pneumoniae	189	%	
Pseudomonas aeruginosa	104	6.57%	Colistin,
Klebsiella sp.	61	3.85%	Polymyxin B
Gram Positive cocci		2%	Linezolid,
			Vancomycin,
			Clindamycin,
			Chlorampheni
Staphylococcus aureus	31	1.96%	col

# BLOOD

												B	lood	isol	ates	- % S	usce	eptibl	e																			
	% of isolates	Amikacin	Amoxicillin/Clavulinic acid	Ampicillin/Sulbactam	Azithromycin	Aztreonam	Cefepime	Cefoperazone/Sulbactam	Cefotaxime	Ceftazidime	Ceftriaxone	Cefuroxime	Chloramphenicol	Ciprofloxacin	Clindamycin	Colistin	Doxycycline	Ertapenem	Erythromycin	Gatifloxacin	Gentamicin	Imipenem	Levofloxacin	Linezolid	Meropenem	Minocycline	Nitrofurantoin	Oflosacin	Oxacillin	PenicillinG	Piperacillin	Piperacillin/Tazobactam	Polymixin B	Teicoplanin	Tetracycline	Tobramycin	Trimethoprim/Sulfametho xazole	Vancomycin
GRAM-POSITIVE ORGANISMS																																						
Coagulase-negative staphylococci	21%	-	42	-	31	ı	-	1	-		ı	-	88	62	68	-	64	-	30	-	76	-	66	100	-	64	-	55	43	13	-	-	-	100	ı	-	35	100
Staphyloccocus aureus	21%	-	60	-	36	-	-	-	-	-	1	-	88	76	82	-	72	-	36	-	80	-	77	100	-	72	-	68	60	16	-	-	-	99	-	-	50	100
												G	RAM	I-NE	GATI	VE O	RGAI	NISM	S																			
Klebsiella pneumoniae	18%	40	-	19	-	-	15	32	11	-	11	10	50	22	T -	100	16	51	-	39	24	37	22	-	37	16	-	50	T -	-	17	50	100	-	0	-	18	-
Acinetobacter baumanii	9%	28	-	40	-	-	26	71	7	-	11	0	0	26	-	100	28	33	-	39	28	32	28	-	32	28	-	-	-	-	-	-	100	-	-	29	26	-
Escherichia coli	7%	74	-	46	-	-	42	72	0	-	26	25	71	42	-	100	15	68	-	61	58	68	42	-	68	15	-	0	-	-	0	0	100	-	0	-	39	-
Klebsiella sp.	5%	41	-	11	-	-	11	30	-	-	11	11	70	26	-	100	19	54	-	50	19	48	26	-	54	19	-	-	-	-	-	-	100	-	-	-	19	-
																																					П	

# Blood

Blood (Total = 4439; no. of is 675 (15.2%))	solat		100% Sensitive antibiotics
Organism	No.	%	
Gram Positive cocc	i	41%	
Staphylococcus,		20.5	
coagulase negative	139	9%	
		20.59	Linezolid,
Staphylococcus aureus	137	%	Vancomycin
		38.35	
Gram negative bacil	li	%	
		17.62	
Klebsiella pneumoniae	119	%	
		8.59	
Acinetobacter baumannii	58	%	
		6.81	
Escherichia coli	46	%	
		5.33	Colistin,
Klebsiella sp.	36	%	Polymyxin B

# SWAB

												S	wab	sola	tes –	- % S	iusce	ptible	2																			
Organism	% of isolates	Amikacin	Amoxicillin/Clavulinic acid	Ampicillin/Sulbactam	Azithromycin	Aztreonam	Cefepime	Cefoperazone/Sulbactam	Cefotaxime	Ceftazidime	Ceftriaxone	Cefuroxime	Chloramphenicol	Ciprofloxacin	Clindamycin	Colistin	Doxycycline	Ertapenem	Erythromycin	Gatifloxacin	Gentamicin	Imipenem	Levofloxacin	Linezolid	Meropenem	Minocycline	Nitrofurantoin	Offoxacin	Oxacillin	PenicillinG	Piperacillin	Piperacillin/Tazobactam	Polymixin B	Teicoplanin	Tetracycline	Tobramycin	Trimethoprim/Sulfametho xazole	Vancomycin
GRAM-NEGATIVE ORGANISMS																																						
Pseudomonas aeruginosa	26%	24	-	0	-	32	32	22	-	21	0	0	0	19	-	100	0	0	-	0	22	42	19	-	43	0	-	23	-	-	20	37	100	-	-	21	0	-
Klebsiella pneumoniae	20%	30	-	14	-	-	15	32	21	0	8	6	39	14	T -	100	15	32	-	41	22	31	13	-	31	15	-	18	T -	-	18	60	100	-	10	-	14	-
Escherichia coli	14%	64	-	24	-	-	18	48	10	-	4	3	66	6	T -	100	17	66	-	38	46	56	7	-	56	17	-	67	T -	-	33	64	100	-	67	-	16	-
Acinetobacter baumanii	10%	7	-	21	-	-	5	46	1	-	2	0	0	6	-	100	10	0	-	18	5	7	6	-	7	10	-	-	T -	-	-	-	100	-	-	6	8	-
Klebsiella sp	5%	20	-	9	-	-	8	27	-	-	5	3	47	9	-	100	13	41	-	26	18	37	12	-	38	13	-	-	T-	-	-	-	100	-	-	- 1	13	-
GRAM-POSITIVE ORGANISMS																					•																	
Staphyloccocus aureus	15%	-	50	-	30	-	-	-	-	-	-	-	91	60	70	T -	63	-	30	-	68	-	60	100	-	64	-	42	50	4	-	-	-	99	-	-	39	100

# Swab

Crevale			1000/ Caracitizes
Swab	_		100% Sensitive
(Total = 3543; No. of is	solat	ces =	antibiotics
1890 (53.34%))			
Organism	No.	%	
Gram Negative bacil	li	75%	Colistin,
Pseudomonas	483	25.5	Polymyxin B
aeruginosa 💮 💮 💮		<b>5</b> %	
Klebsiella pneumoniae	373	19.73	
•		%	
Escherichia coli	268	14.17	
		%	
Acinetobacter baumannii	192	10.15	
		%	
Klebsiella sp.	101	5.34	
1		%	
Gram positive cocci	i	15.13	Linezolid,
		%	Vancomycin,
Staphylococcus aureus	286	15.13	Teicoplanin,
		%	Chloramphen
			icol

## Pus

													us i	solat	tes -	% Su	iscep	tible																				
Organism	% of isolates	Amikacin	Amoxicillin/Clavulinic acid	Ampicillin/Sulbactam	Azithromycin	Aztreonam	Cefepime	Cefoperazone/Sulbactam	Cefotaxime	Ceftazidime	Ceftriaxone	Cefuroxime	Chloramphenicol	Ciprofloxacin	Clindamycin	Colistin	Doxycycline	Ertapenem	Erythromycin	Gatifloxacin	Gentamicin	Imipenem	Levofloxacin	Linezolid	Meropenem	Minocycline	Nitrofurantoin	Oflosacin	Okacillin	PenicillinG	Piperacillin	Piperacillin/Tazobactam	Polymixin B	Teicoplanin	Tetracycline	Tobramycin	Trimethoprim/Sulfamethosaz	Vancomycin
GRAM-POSITIVE ORGANISMS																																						
Staphyloccocus aureus	43%	-	62	-	48	-	-	-	-	-	-	-	96	76	91	-	82	-	49	-	83	-	78	100	-	83	-	59	61	5	-	-	-	100	-	-	41	100
GRAM-NEGATIVE ORGANISMS																																						
Escherichia coli	21%	78	-	38	-	-	28	74	50	-	14	11	70	14	-	100	15	87	-	46	58	78	14	-	80	15	-	100	-	-	100	100	100	-	100	-	22	-
Klebsiella pneumoniae	14%	47	-	28	- 1	-	30	49	23	-	13	11	44	24	-	100	21	52	-	50	43	40	24	-	40	21	1	71	-	-	29	60	100	-	71	-	21	-
Pseudomonas aeruginosa	8%	46	-	-	-	49	54	38	-	42	-	-	-	32	-	100	-	-	-	-	30	68	32	-	68	-	ı	35	-	-	31	56	98	-	-	29	-	-
Klebsiella sp	4%	26	-	16	-	-	11	21	-	-	11	5	37	11	-	100	5	22	-	16	21	22	16	-	22	5	ı	-	-	-	-	-	100	-	-	-	21	-
Acinetobacter baumanii	3%	20	-	27	-	-	0	57	0	-	0	-	-	13	-	100	7	-	-	27	13	7	13	-	7	7	ı	-	ı	-	-	ı	100	-	ı	8	7	-
·																																						

## Pus

Pus (Total = 1211 ; Number of = 534 (44.09%))	of iso	lates	100% Sensitive antibiotics
Organism	No.	%	
Gram positive cocci		42.50	Linezolid,
		%	Vancomycin,
Staphylococcus aureus	227	42.50	Teicoplanin
ss. aureus		%	
Gram negative bacill	i	48.00	
		%	
Escherichia coli	111	20.78	Colistin,
		%	Polymyxin B,
			Piperacillin,
			Piperacillin-
			tazobactam,
			Tetracycline
Klebsiella pneumoniae	76	14.23	Colistin,
		%	Polymyxin B
Pseudomonas aeruginosa	41	7.67	
		%	
Klebsiella sp.	19	3.55	
		%	
Acinetobacter baumannii	15	2.80	
		%	

# SPUTUM

							Sput	um Is	solat	es —	% St	ısce	ptible	,														
Organism	%of isolates	Amikacin	Ampicillin/Sulbactam	Aztreonam	Cefepime	Cefoperazone/Sulbactam	Cefotaxime	Ceftazidime	Ceftriaxone	Cefuroxime	Chloramphenicol	Ciprofloxacin	Colistin	Doxycycline	Ertapenem	Gatifloxacin	Gentamicin	Imipenem	Levofloxacin	Meropenem	Minocycline	Ofloxacin	Piperacillin	Piperacillin/Tazobactam	Polymixin B	Tetracycline	Tobramycin	I rimethoprim/Sulfamethoxaz ole
GRAM-NEGATIVE ORGANISMS																												
Klebsiella pneumoniae	22%	64	31	<u> </u>	39	56	-		31	25	61	36	100	31	62	51	56	61	39	61	31	-		-	100			42
Pseudomonas aeruginosa	16%	75	- )	76	81	81	-	68	-			73	100	-		-	72	81	73	82	-	80	71	87	100		61	-
Escherichia coli	11%	87	28		29	67	0		5	4	73	15	98	15	76	39	67	73	15	74	15	17	0	100	98	40		27
Acinetobacter baumanii	9%	13	26	ı	10	50	0	-	0			21	100	12		29	10	13	21	13	12	-	-	-	100		11	13

# Sputum

Sputum (Total = $1800$ ; no of iso $534(29.66\%)$ )	olate	es=	100% Sensitive antibiotics
Organism	No	%	
Gram negative bacil	li	56%	Colistin,
Klebsiella	115	21.5	PolymyxinB
pneumoniae		3%	(*Piperacillin
Pseudomonas aeruginosa	84	15.73	-tazobactam)
		%	
Escherichia coli*	60	11.23	
		%	
Acinetobacter baumannii	48	8.98	
		%	
Klebsiella sp.	36	6.74	
		%	

# FLUID

												F	luid i	sola	tes -	% Su	scep	tible																				
Organism	% of isolates	Amikacin	Amosicillin/Clavulinic acid	Ampicillin/Sulbactam	Azithromycin	Aztreonam	Cefepime	Cefoperazone/Sulbactam	Cefotaxime	Ceftazidime	Ceftriaxone	Cefuroxime	Chloramphenicol	Сіргоflокасіп	Clindamycin	Colistin	Doxycycline	Ertapenem	Erythromycin	Gatifloxacin	Gentamicin	Imipenem	Levofloxacin	Linezolid	Meropenem	Minocycline	Nitrofurantoin	Ofloxacin	Oxacillin	PenicillinG	Piperacillin	Piperacillin/Tazobactam	Polymixin B	Teicoplanin	Tetracycline	Tobramycin	Trimethoprim/Sulfametho xazole	Vancomycin
GRAM-NEGATIVE ORGANISMS																																						
Escherichia coli	33%	68	-	15	-	-	13	58	0	-	5	5	65	13	-	100	13	72	-	35	58	65	13	-	65	13	-	-	-	-	-	0	100	-	-	-	25	-
Klebsiella pneumoniae	16%	35	-	6	-	-	15	37	0	-	0	0	35	5	-	100	6	0	-	30	15	32	5	-	30	6	-	0	-	-	0	100	100	-	0	-	25	-
Acinetobacter baumanii	14%	12	-	12	-	-	12	35	20	-	12	-	-	18	-	100	13	-	-	24	12	12	18	-	12	13	-	-	-	-	-	-	100	-	-	10	6	-
Pseudomonas aeruginosa	8%	44	-	-	-	56	56	-	-	-	-	-	-	44	-	100	-	-	-	-	56	56	44	-	56	-	-	20	-	-	33	56	100	-	-	44	-	-
GRAM-POSITIVE ORGANISMS																																						
Staphyloccocus aureus	15%	-	68	-	42	-	-	-	-	-	-	-	100	68	90	-	79	-	42	-	88	-	63	100	-	79	-	43	68	11	-	-	-	100	-	-	47	100
				-			$\sim$		-																											-		

# Fluid

Fluid			100% Sensitive
(Total = 523; isolates 1)	<b>24 (23.</b> <sup>1</sup>	<mark>7%</mark> ))	antibiotics
	Numb		
	er of		
	isolate		
Organism	S		
Gram negative baci	lli	<b>72</b> %	
Escherichia coli	41	33%	
		16.12	
Klebsiella pneumoniae*	20	%	Colistin,
		13.70	
Acinetobacter baumannii	17	%	
Pseudomonas aeruginosa	10	8.06%	
		15.32	
Gram positive cocci		%	Linezolid,
			Vancomycin,T
Staphylococcus aureus ss.		15.32	eicoplanin,Chl
aureus	19	%	oramphenicol

## GRAM POSITIVE COCCI

- Coagulase negative staphylococci isolates in blood
  - Treat the infection, Not contamination
- MRSA isolates 44%
- Emergence of VISA & VRSA

#### GRAM NEGATIVE BACILLI

- Resistance to Fluoroquinolones 80%
- $\circ$  ESBL isolates = 90%
- $\circ$  CRE isolates = 30-70%
- MDR Acinetobacter baumanii & Pseudomonas aeruginosa
- Only 100% Sensitive drugs are Colistin & Polymyxin B
- Emergence of resistance to colistin & Polymyxin B

#### **OBSERVATIONS**

- MRSA / TB / Swine Flu Presentations
- High Incidence of S. Epidermidis in Blood Cultures
- Limited N95 Masks
- Limited De-Escalation of Antibiotic Therapy
- Increasing Resistance Pattern
  - Will and Should be followed over several years

## DR. SANJAY BHATT

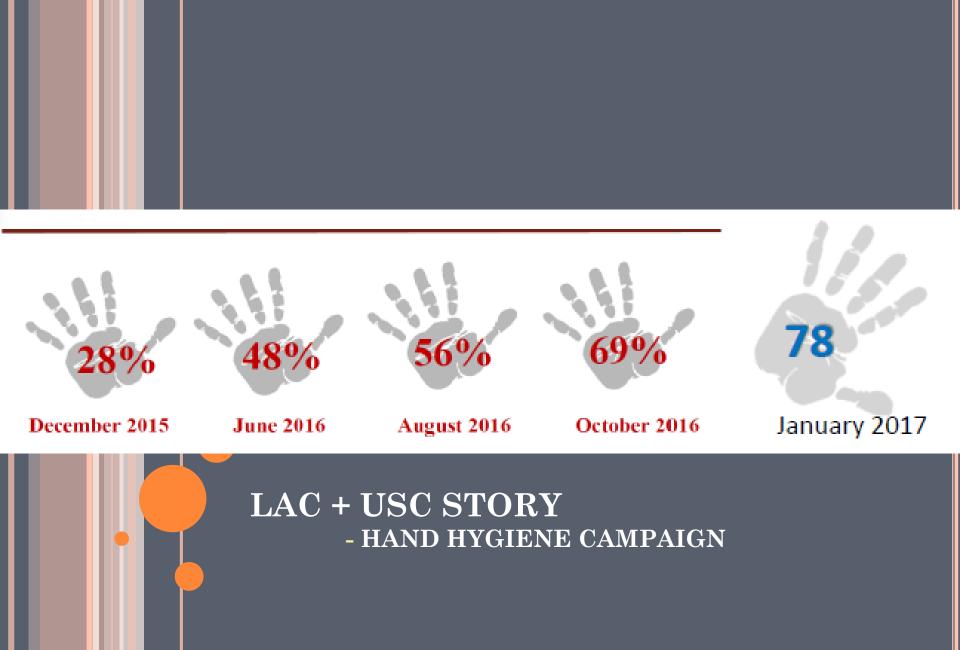
Presentation and Discussion

#### INFECTION CONTROL

- o STEP 1
- Hand Washing / Gloves
- Masks
- $\circ$  Bed Up > 30°

- o STEP 2
- De-escalation of Antibiotics
- Draw Cultures
- Aseptic Technique

- STEP 3
- Sterile Procedures
- Awareness though Identifiers / Poster
- Attention to critical labs / cultures



**SISTER:** 48%

PHYSICIAN: 11%

OTHER: 81% !!

16/1 - 11/2N = 620

VS HOSPITAL



#### VS HOSPITAL CULTURES

- 2016 Data (~1000 beds total)
  - Urine Cultures
    - 6055 total
    - 17 Urine Cultures / Day
  - Blood Cultures
    - 4439 total
    - 12 Blood Cultures / Day
    - Without NICU (1580):

**o**8 / **Day** 

## ASEPTIC TECHNIQUE (BLOOD CULTURE)

- o 21% S. Epidermidis
  - Normal Human Flora (not pathogenic except in IC)



#### OTHER INFECTION CONTROL MEASURES

QUARTERLY Infection Control & Microbiology

Newsletter ..... "The Flora"

- Web Access with updates
- Regular Infection Control Meetings
  - Every 3 months at least or more if needed
- Intensify Environmental Sanitation and Biomedical Waste Management within the hospital UNIT BY UNIT
- Ensure Availability of Supplies

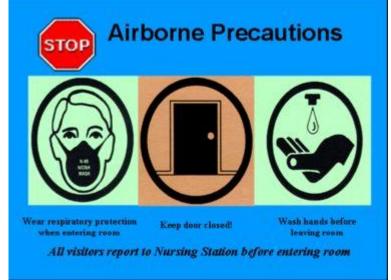
# INTEGRITY OF INFECTION CONTROL AT BEDSIDE PROCEDURES

- Ensure Sterile Precautions
  - Pre/Peri/Post Preparation
  - Checklist
  - Procedure Tray
  - Proper Documentation
  - Follow up
- Each Unit / Ward / Department is Responsible for Adherence
- Awareness through Posters/
- STUDENT/RESIDENT TEAM RESPONSIBLE AND ACCOUNTABLE TO FACULTY
- Recognize with Star Ratings by CQM

# AWARENESS THOUGH IDENTIFIERS / POSTERS: Examples







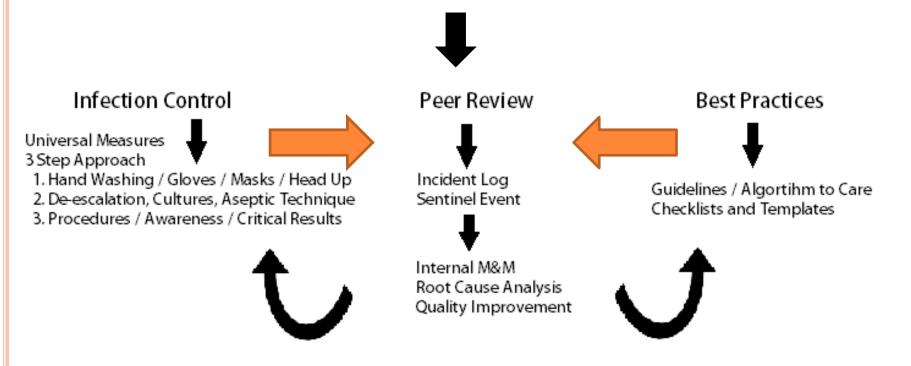
Juzar ali Presentation and Discussion

A PIVOTAL ARM IN THE IMPLEMENTATION OF EQUIPS PROGRAM IS.....

# INTERNAL PEER REVIEW PROCESSES

#### EQuIPS (Enhanced Quality Improvement and Patient Safety)

A Team Effort Between Doctors / Nurses / Allied Health Leadership Who Believe High quality safe patient-care can be provided in spite of a high demand, challenging environment and limited resources



Goals: Increase Awareness, Enhanced Patient Safety, Education, Process Improvement





#### IDENTIFY SENTINEL EVENT OR ANY DEVIATION IN PATIENT SAFETY OR QUALITY OF CARE IRRESPECTIVE OF LEVEL AT WHICH IT OCCURRED

- Review and identify potential critical clinical or operational problems
- If and What went wrong detected ...
- What could have been avoided......
- What could be corrected......

Method: Examples each Unit can adopt

- INCIDENT NOTIFICATION through LOG/ HOT LINE
- "Missed Call" /SMS Number
- Create a ... NHL SAFE ... 645 7233
- Or use or develop an App like "Magpi"

## BEST PRACTICES

INDWNTIFICATION OF NEED ,IMPLEMENTATION AND MONITORING

PUBLICATION OF DATA WITH ANALYSIS & RESULTS WEB SITE , NEWS LETTER, JOURNALS

## FOCUS 3 OF EQUIPS

## BEST PRACTICES FOCI TO WORK AT

#### CHOOSE A TARGET BASED ON DATA & NEED & PRIORITY

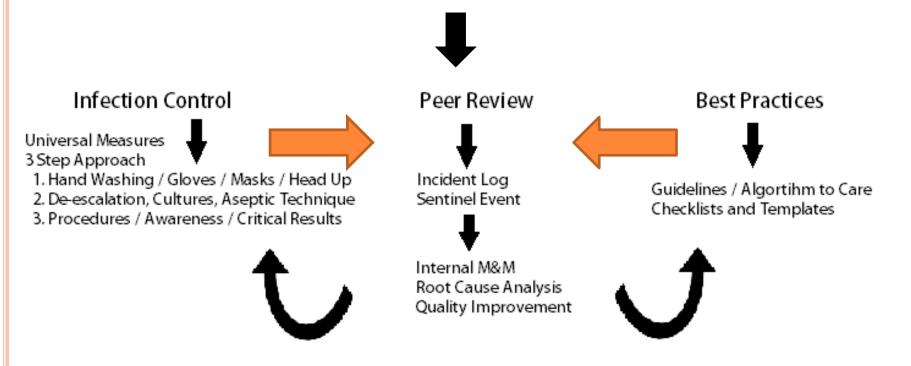
- Ventilator Associated Pneumonia 2.Catheter Associated UTI or Blood stream Infections 3. Blood Reactions 4. Bed Sores 5. Documentation Deficiencies 6. Timeliness of PCI or Thrombolytic 7. Proper Discharge Processes 8.Pneumonia Follow up 9. In Patient Surgical Infection prevention 10. Vascular Ortho Injuries 11. Delayed Dx in Oncology 12. Surgical site infection 13 OR Efficiency and turnover methods process 14. Pre peri and post partum issues
- 1. PICK YOUR STARS FROM THE GALAXY

 \EACH DEPT TO DECIDE and PICK A FOCUS EVERY QUARTER

• CREATE GUIDELINES/CHECKLISTS/TEMPLATES etc.

#### EQuIPS (Enhanced Quality Improvement and Patient Safety)

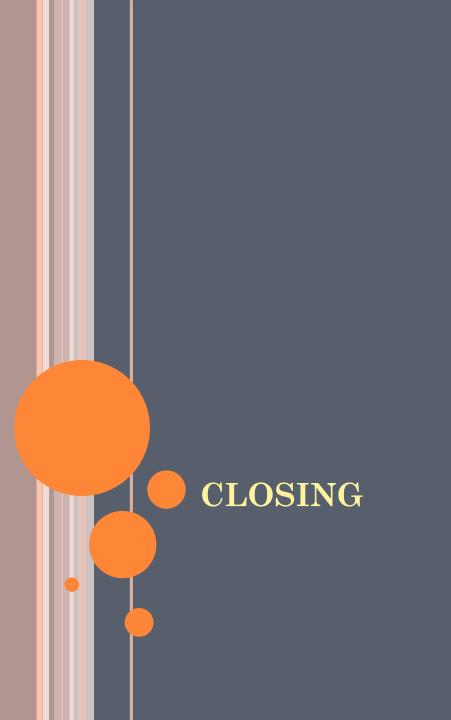
A Team Effort Between Doctors / Nurses / Allied Health Leadership Who Believe High quality safe patient-care can be provided in spite of a high demand, challenging environment and limited resources



Goals: Increase Awareness, Enhanced Patient Safety, Education, Process Improvement







# ACTION WHEN WHO AND WHERE?

#### • NURSING /ALLIED HEALTH TEAM

- INFECTION CONTROL MEASURES
- BEDSIDE PROCEDURE CHECK LIST

#### SENIOR STUDENTS / JUNIOR DOCTORS

INCIDENT LOGS / PATIENT SAFETY STEPS CHECK LIST



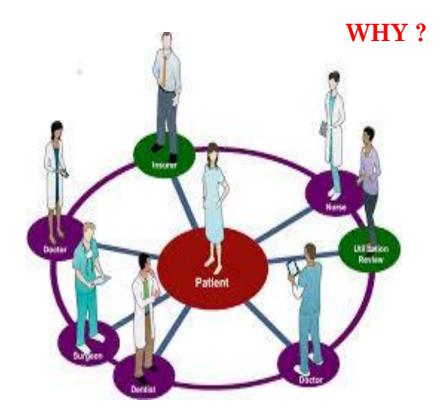
#### FACULTY

- INTERNAL PEER REVIEW AND BEST PRACTICES DEVELOPMENT IN CONCENTRIC CIRCLES WITH RIPPLE EFFECT WITH ONE STEP AT A TIME, ONE BED AT A TIME, ONE UNIT AT A TIME

  DEPARTMENTS
- DEVELOP OR RESURRECT & ENERGIZE INFECTION CONTROL COMMITTEE / SUPPORT CQM

#### ADMINISTRATION :

- SUPPORT / RESOURCES / INCENTIVES/ RECOGNITION
- PREP FOR INAUGRATION, ACCREDITATION







#### SUMMARY

- Teamwork: Top Down /Bottom Up
- Leadership at all levels
- Patient Care: It is the right thing to do
- HealthCare Team : It deserves this
- Possible: It can be done
- Act as a role model
  - Follow a procedure, rules and guidelines as a DEMAND
  - To create and then COMMAND a team following
  - Don't worry about naysayers and initial non followers
  - THEY WILL FOLLOW....BELIEVE IT
- It has been shown to be Evidence Based Practice
- Long Term Effect:
  - A few extra minutes today pays of in the long term in Safety ,Quality, Resources & Recognition

## PRIOR TO OPENING OF THE NEW HOSPITAL

- Have Processes in Place (Require starting now)
- 2017 Inauguration
  - Officials and Bureaucrats
  - Health Teamwork
    - We can showcase our work thus far
    - Main Presentation will be part of the inauguration
    - Poster Show in New Hospital during Inauguration



#### RESPECTFUL PLEA

- You have the Manpower / Material / Means / Monies
  - All you need is the Motivation
- Danger in the Status Quo
  - Please don't use the limitations of the system, environmental challenges, and the patient population you serve as a CRUTCH to maintain Status Quo
- Keep the Passion to Cynicism ratio HIGH
- Thank you

# RESOURCES / REFERENCES

Also in Folder submitted to CQM Lead

### INFECTION CONTROL

• Resource



Patient Safety

A World Alliance for Safer Health Care

SAVE LIVES
Clean Your Hands

Glove Use Information Leaflet

Outline of the evidence and considerations on medical glove use to prevent germ transmission

Definitions

The impact of wearing gloves on adherence to hand hygiene policies has not been definitively established, since published studies have yielded contradictory results. However, the recommendation to wear gloves during an entire episode of care for a patient who requires contact precautions, without considering indications for their removal, such as an indication for hand hygiene, could actually lead to the

### GLOVES

- Worn to:
  - To reduce the risk of contamination of health-care workers hands
  - To reduce the risk of germ dissemination
    - To the environment
    - Transmission from the health-care worker to/from the patient
    - From one patient to another.
- Gloves should therefore be used during all patient-care activities that may involve exposure to blood and all other body fluid (including contact with mucous membrane and non-intact skin), during contact precautions and outbreak situations.

### KEY POINTS

- Glove use does NOT replace alcohol-based product or hand-washing
- Wear gloves when anticipated contact with blood or other body fluids, mucous membranes, nonintact skin or potentially infectious material
- Do not wear the same pair of gloves for more than one patient.
- Change gloves if moving from a contaminated body site to another body site
- Do not reuse gloves

#### STERILE GLOVES INDICATED

Any surgical procedure; vaginal delivery; invasive radiological procedures; performing vascular access and procedures (central lines); preparing total parental nutrition and chemotherapeutic agents.

#### EXAMINATION GLOVES INDICATED IN CLINICAL SITUATIONS

Potential for touching blood, body fluids, secretions, excretions and items visibly soiled by body fluids.

DIRECT PATIENT EXPOSURE: Contact with blood; contact with mucous membrane and with non-intact skin; potential presence of highly infectious and dangerous organism; epidemic or emergency situations; IV insertion and removal; drawing blood; discontinuation of venous line; pelvic and vaginal examination; suctioning non-closed systems of endotrcheal tubes.

INDIRECT PATIENT EXPOSURE: Emptying emesis basins; handling/cleaning instruments; handling waste; cleaning up splits of body fluids.

#### GLOVES NOT INDICATED (except for CONTACT precautions)

No potential for exposure to blood or body fluids, or contaminated environment

DIRECT PATIENT EXPOSURE: Taking blood pressure, temperature and pulse; performing SC and IM injections; bathing and dressing the patient; transporting patient; caring for eyes and ears (without secretions); any vascular line manipulation in absence of blood leakage.

INDIRECT PATIENT EXPOSURE: Using the telephone; writing in the patient chart; giving oral medications; distributing or collecting patinet dietary trays; removing and replacing linen for patient bed; placing non-invasive ventilation equipment and oxygen cannula; moving patient furniture.

	Indication
Gloves on	Before a sterile procedure
	When anticipating contact with blood or another body fluid, regardless of the existence of sterile conditions and including contact with non-intact skin and mucous membrane
	Contact with a patient (and his/her immediate surroundings) during contact precautions.
Gloves off	As soon as gloves are damaged     (or non-integrity suspected)
	When contact with blood, another body fluid, non-intact skin and mucous membrane has occurred and has ended
	When contact with a single patient and his/her surroundings, or a contaminated body site on a patient has ended
	4) When there is an indication for hand hygiene.

#### STERILIUM

- Pathogens may gain access to the caregivers' hands via small defects in gloves or by hand contamination during glove removal.
- Hand hygiene by rubbing or washing ensures hand decontamination

# Contact Precautions संपर्ध सावयेती

o Gloves / Sterilium

- Norovirus
- Rotavirus
- Generalized rash
- Draining wounds
- Uncontrolled secretions
- Pressure ulcers
- Ostomy tubes
- Clostridium difficile
- Acidobacteria
- ESBL
- o MRSA (44%)

#### DROPLET PRECAUTIONS

- Standard Mask
- Travel 1 meter from the patient
  - Deposited on the host's nasal mucosa, conjunctivae or mouth

- Pertussis
- Influenza
- Diphtheria
- NeisseriaMeningitidis

## AIRBORNE PRECAUTIONS

o N95 Mask

- Measles
- Severe Acute Respiratory Syndrome (SARS)
- Varicella (chickenpox)
- o Swine Flu
- Mycobacterium tuberculosis

### BED UP $> 30^{\circ}$

- Decrease the incidence of aspiration pneumonia and pressure ulcers
- o Ann Intern Med 1992; 116:540-543
  - Bed Up decreased rates of aspiration of gastric contents four-fold
- Lancet 1999; 354:1851-1858
  - 34% in supine position developed VAP compared with 8% of patients in the head up group

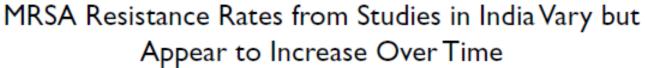
## BED UP $> 30^{\circ}$

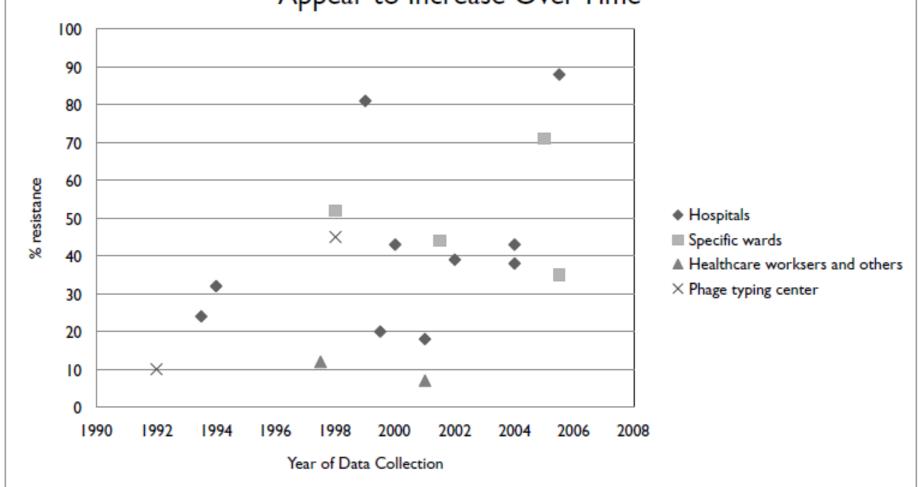
#### • Indications:

- Altered Sensorium / Overdose
- Nausea / Vomiting
- Intubated / Peri-Intubation

### Contraindications

- Neurosurgical Approaches
- Ischemic Stroke (First 24 hours if tolerated)





## WHO'S "DIRTY DOZEN"

SEE WHO Feb 2017 released report with regional data

#### WHO PRIORITY PATHOGENS

CRITICAL: Acinetobacter Baumanni – Carbp res

Pseudomonas Aeruginosa – Carbp res

Enterobacteriaceae Carbp' res ESBL

HIGH: Enterococcus, Staph MRSA, QR-

Salmonella, Hb/Cb/NG

MEDIUM: Strep /Hemoph/Shigella

# CRIT CARE CLIN. 2011 JAN;27(1):149-62

- Antibiotic de-escalation
  - Mechanism whereby the provision of effective initial antibiotic treatment is achieved while avoiding unnecessary antibiotic use that would promote the development of resistance
  - Based on microbiology results around the day 3 therapy point
  - The empiric antibiotics that were started are stopped or reduced in number and/or narrowed in spectrum
  - Clinically effective and appropriate

## DE-ESCALATION OF ANTIBIOTICS (43%)

Crit Care Med. 2012 May;40(5):1404-9. doi: 10.1097/CCM.0b013e3182416ecf.

#### Antibiotic strategies in severe nosocomial sepsis: why do we not de-escalate more often?

Heenen S1, Jacobs F, Vincent JL.

Author information

#### Abstract

OBJECTIVES: To assess the use of antibiotic de-escalation in patients with hospital-acquired severe sepsis in an academic setting.

**DESIGN:** We reviewed all episodes of severe sepsis treated over a 1-yr period in the department of intensive care. Antimicrobial therapy was considered as appropriate when the antimicrobial had in vitro activity against the causative microorganisms. According to the therapeutic strategy in the 5 days after the start of antimicrobial therapy, we classified patients into four groups: de-escalation (interruption of an antimicrobial agent or change of antibiotic to one with a narrower spectrum); no change in antibiotherapy; escalation (addition of a new antimicrobial agent or change in antibiotic to one with a broader spectrum); and mixed changes.

**SETTING:** A 35-bed medico-surgical intensive care department in which antibiotic strategies are reviewed by infectious disease specialists three times per week.

PATIENTS: One hundred sixty-nine patients with 216 episodes of severe sepsis attributable to a hospital-acquired infection who required broad-spectrum β-lactam antibiotics alone or in association with other anti-infectious agents.

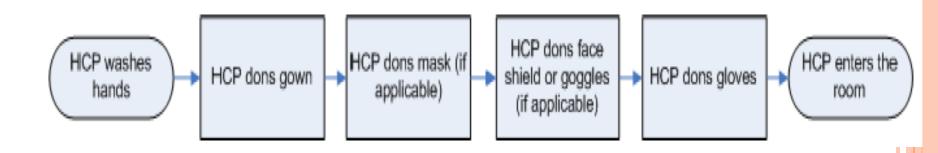
MEASUREMENTS AND MAIN RESULTS: The major sources of infection were the lungs (44%) and abdomen (38%). Microbiological data were available in 167 of the 216 episodes (77%). Initial antimicrobial therapy was inappropriate in 27 episodes (16% of culture-positive episodes). De-escalation was applied in 93 episodes (43%), escalation was applied in 22 episodes (10%), mixed changes were applied in 24 (11%) episodes, and there was no change in empirical antibiotic therapy in 77 (36%) episodes. In these 77 episodes, the reasons given for maintaining the initial antimicrobial therapy included the sensitivity pattern of the causative organisms and previous antibiotic therapy. The number of episodes when the chance to de-escalate may have been missed was small (4 episodes [5%]).

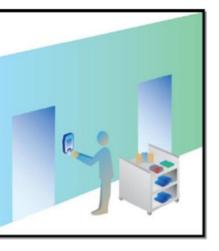
#### DRAW CULTURES

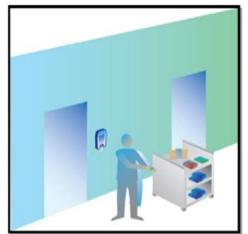
- Indications for Blood Cultures
- Balance between wasteful and useful
  - Based on Pre-Test Probability (Cellulitis 2%+ to Shock 69%+)
  - Suspicion of bacteremia or fungemia
    - Especially Important: Sepsis, meningitis, osteomyelitis, arthritis, endocarditis, peritonitis, pneumonia, and fever of unknown origin

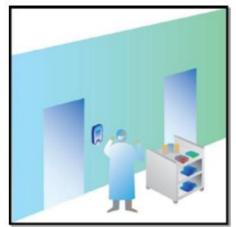
Does this adult patient with suspected bacteremia require blood cultures? AU Coburn B, Morris AM, Tomlinson G, Detsky AS SO JAMA. 2012 Aug;308(5):502-11.

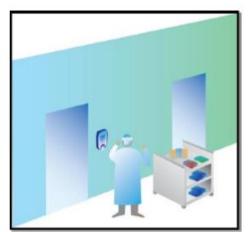
## PROCEDURES AND PPE



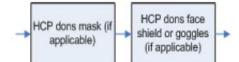






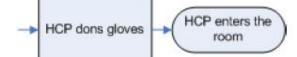












#### संक्रमण नियंत्रण INFECTION CONTROL ચેપ નિયંત્રણ પગલાં

बिस्तर स्वच्छता

BEDSIDE

**HYGIENE** 

से संपर्क करें

एहतियात

CONTACT

**PRECAUTION** 

સંપર્ક દ્વારા ચેપ અટકાવવા

एहतियात

**DROPLET** 

PRECAUTION

हवाई

एहतियात

**AIRBORNE** 

**PRECAUTION** 

अलगाव एहतियात **ISOLATION**  **SEPARATION** 



#### We're starting a major operation.

Mercy Health Partners has broken ground on one of its most important operations: A new atate-of-the-art hospital designed to serve Cincinnati's west side.

The new facility will offer comprehensive care in nearly every specialty from cardiac and women's services to oncology and orthopedics. And, like other Mercy hospitals, it will deliver advanced medicine within a few miles of where patients and their families live.

This new hospital comprements our growing network of primary care and specialty physician's groups, sister hospitals, urgent care facilities, diagnostic centers and senior communities that provide a system of care for our entire community.

The Sisters of Mercy and The Franciscan Sisters of the Poor defined our mission in 1858, and for the past. 152 years we have been dedicated to their vision. Today, we know they are proud.

e-mercy.com

MERCY



www.cdc.gov/HandHygiene





## RECOMMENDATIONS

- Blood culture
  - In duplicate
  - Before starting antibiotic
- Urine
  - Midstream Early Morning urine sample
  - Mention in the form if catheterized sample is sent
- Sputum
  - Preferable Early morning sample
  - If report does not correlate clinically repeat sample
- Fluid
  - Leak proof sterile container

# Recommendations (cont..)

- Avoid or minimize delay in transportation
- Shift to Automation
  - For culture & sensitivity
- Critical reports on personal contact
- **Descalation** of antibiotics
  - After getting Culture and sensitivity report