











NQF's National Quality Partners...

...maximize the impact of high-leverage drivers—payment, public reporting, consumer engagement, and accreditation and certification—that each of us brings to bear

NATIONAL QUALITY FORUM







National Targets For 2020

By 2020, the United S	tates will:
For CDC Recognized l	Jrgent Threats:
Reduce by 50% the incide	ence of overall Clostridium difficile infection compared to estimates from 2011.
Reduce by 60% carbapen	em-resistant Enterobacteriaceae infections acquired during hospitalization compared to estimates.
Maintain the prevalence o	of ceftriaxone-resistant Neisseria gonorrhoeae below 2% compared to estimates from 2013.
For CDC Recognized	Serious Threats:
Reduce by 35% multidrug from 2011.	g-resistant Pseudomonas spp. infections acquired during hospitalization compared to estimates
Reduce by at least 50% or compared to 2011.*	verall methicillin-resistant Staphylococcus aureus (MRSA) bloodstream infections by 2020 as
Reduce by 25% multidrug	g-resistant non-typhoidal Salmonella infections compared to estimates from 2010-2012.
Reduce by 15% the numb	per of multidrug-resistant TB infections. ¹
Reduce by at least 25% the stimates from 2008.	ne rate of antibiotic-resistant invasive pneumococcal disease among <5 year-olds compared to
Reduce by at least 25%th estimates from 2008.	e rate of antibiotic-resistant invasive pneumococcal disease among >65 year-olds compared to





NQF's Commitments to Advancing Antibiotic Stewardship

- ...work with federal agencies and private-sector stakeholders to encourage development and submission performance measures for use in quality improvement and accountability programs to improve appropriate antibiotic use
- ...expedite the process so that, if endorsed, the measures are positioned for rapid implementation in public and private programs
- ...encourage and support the use of antibiotic resistance and utilization measures through our diverse membership and ongoing measure selection activities
- ...convene a multistakeholder Action Team to activate our 425+ organizational members and promote a shared agenda aligned with the National Action Plan
- …foster partnerships that advance antibiotic stewardship programs, education, and engagement of our members to create the "burning platform"



NATIONAL QUALITY FORUM







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ASP Phase 1 Foundational

- MD/ PharmD Champion
- Multidisciplinary team
- Gap Assessment

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- Assess staff resources
- Competency/Training Planning
- Communication Plan for facility
- CEO support for AMP by
- approval of gap and action plan

- Selecting Physician
 Champion
- Complete gap assessment and action plan as a team
- Determine staffing needs to adequately resource AMP activities
- Create competency/training plan for all disciplines based on current knowledge and involvement
- Invite CEO to AMP team meeting to discuss plan, resources, and support



Gap Analysis

- Executive and Medical Staff Ownership
- Information technology
- · Staff development and competencies
- IV to PO conversion
- Renal dosing
- Criteria for antimicrobial use
- Formulary review
- Antibiogram and culture and sensitivity reporting(CLSI)
- Empiric antimicrobial treatment guidelines
- Antimicrobial streamlining (de-escalation)
- Duration
- Utilization reviews
- Measurement

http://www.cdc.gov/getsmart/healthcare/improveefforts/resources/doc/AMP-GapAssessment.doc



Clinical	Process		
 Length of stay 	Dose optimization		
Clinical cure/failure rates	 Adherence to hospital specific guidelines 		
 Readmission rates (30 days) 	 Appropriate de-escalation/streamlining 		
Resistance rates	 Appropriateness of therapy 		
Infection-related mortality	Cultures before antibiotics		
	Outcomes		
Humanistic	Economic		
 Adverse drug events avoided 	Antimicrobial utilization (DDD or DOT)		
• Time to receipt of appropriate antimicrobials	 Hospital wide antimicrobial expenditures 		
 Duration of antimicrobial therapy 	Relative consumption use		
IV/PO conversion rates	 Rate of intravenous antimicrobial use 		
Outpatient intravenous therapy rates	 Nonformulary agents avoided 		

ASP Phase 2

- Antimicrobial formulary review
- Review metrics (DDD Spreadsheet)
- Review CAP and SCIP core measure
- Microbiology
 - CLSI susceptibility reporting
 - Review new CLSI breakpoints
 - Appropriate use of microbiology document with emphasis on obtaining appropriate cultures <u>before</u> starting antimicrobial therapy for new septic episodes

Dose optimization

- Weight-based dosing
- Renal dosing
- IV to PO

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Associated with clinical and economic benefit

- Review facility by-laws and state scope of practice for automatic interchanges
- P&T and Med Exec approval
- Many institutions have these activities implemented (CAP/SCIP)
- Routine IV to PO/ Renal Dosing

ASP Phase 3

- Kinetic dosing
 - Vanc and AG
 - Prolonged infusion for pipercillin/tazobactam and carbapenems
- Approve institutional guidelines
 - CAP, HAP, UTI and ABU, MRSA, VAP, intra-abdominal, surviving sepsis, C. difficile
- Timely and appropriate use of antibiotics based on
- approved institutional guidelines and local antibiograms • Optimize duration based on evidence-based peer review publications
- Evaluate use agents based on local needs (front/back-end approach)
 - Suggested drugs: daptomycin, linezolid, echinocandins, tigecycline, and carbapenems
- Clinical pharmacy rounding with team

- Involve nursing staff early in extended infusion work
- Meropenem and P/T prolonged infusion recommendation can be combined with formulary change to P&T
- Evaluate ability of pharmacist to cover drug regimen reviews and rounding consistently

ASP Phase 4

• De-escalation "72 hour time-out"

• Suggestions: review charts with positive blood cultures, 3 or more antibiotics for ≥72 hours, drug-bug mismatches, or antibiotics without a positive culture, duration

• Review and/or implement rapid diagnostics, point of care testing, and biomarkers (PCT)for appropriate use

- Ongoing antibiogram development (e.g. unit specific)
- Report approved metrics to all stakeholders on a regular basis

•Clinical decision support/CPOE

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- Overlap will occur with various phases.
- Expectation is for all facilities to complete elements of each phase in a timely manner.
- Advanced programs can start on other phases before the suggested timelines.
- Synergize AMP with Core Measures and Sepsis Programs



Key Elements for Successful ASP/AMP

- Establish compelling need and goals for ASP
- Senior leadership support
- Effective local physician champion
- Adequate resources and competencies (pharmacy, infection preventionist [IP], microbiology, information technology [IT])
- Primary objectives: optimize clinical outcomes and reduce adverse events, not reduce costs
- Good teamwork
- Agreed upon process and outcome measures

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Providence Health and Services

- Born 160 years ago
- 34 acute care hospitals in 5 western states (Alaska, Washington, Oregon, Montana, and California)
- Total net operating revenue, 2014: 12 billion
- Total net income in 2014: 771 million
- System committed to Antimicrobial
 Stewardship

Providence Portland Medical Center (PPMC)

- Licensed beds = 483
- Average daily census = 281
- Average LOS = 4.8 days
- Admissions =21,351
- Infectious Diseases AMS Team :
 - 6 consultants and one teacher/researcher
 - Pharmacists

8/13/2015

- Infection control
- Clinical microbiology





Procalcitonin: A biomarker of activation of the innate immune system

- Serum procalcitonin levels do not increase, or increase minimally, in response to either viral colonization or viral invasive disease
- Potential bacterial pathogens that are present but not invading do not stimulate synthesis of Procalcitonin
- Bacterial pathogens causing pneumonia stimulate an increase in serum procalcitonin levels
- PCT levels increase if viral infection with bacterial superinfection
- Hence, a "normal" serum level of procalcitonin has a very high negative predictive value for invasive bacterial infection but does not distinguish viral colonization from viral invasion.

PCT Can Assist AMS

- Differentiate viral from bacterial infection, especially for RTIs
- Provide evidence of efficacy of selected antimicrobial regimen ("source control")
- Provide guide to the duration of antimicrobial therapy

2008: Launched Procalcitonin at PPMC

- PCT and AMS:
 - Separation of pure viral infection from bacterial infection
 - Data indicating the that duration of antibiotic therapy be determined by time to normalization of PCT level
 - In 2014, 18,000 PCT levels in 4 Portland hospitals
 - 2015, using EMR, system evaluated influence of PCT levels on Days of Antibiotic Therapy

PCT Levels Correlate with LOS/DOT







Microbial Etiology of CAP in 59 Evaluable Patients

	Detection of Potential Pathogen(s)
Sputum and blood cultures; Urine antigens (<i>S.pneu.; Legionella pneu</i> .)	16 of 59 (27%)
As above plus nasal swab PCR for <i>S.pneumoniae and</i> <i>S.aureus</i>	33 of 59 (56%)
All of above plus multiplex PCR panels for respiratory viruses plus PCT level	46 of 59 (78 %)
5.1 × 1.5	

Turn Around Times by Test Type

Test	TATs			
Serum Procalcitonin	1 hr			
Film Array multiplex PCR panel for 17 viruses	1.8 +/- 0.3 hrs			
Urine antigens	5- 7 hrs			
Nasal PCRs: <i>S.aureus and S. pneumo.</i>	12-30 hrs			
Nasal PCR: LDT* for 5 viruses	12-36 hrs			
Sputum Culture	2-3 days			
Blood culture	Up to 5 days			
*LDT = Lab Developed Test				



RESPONSE of Providers

Etiology of Pneumonia (No. of Patients	Days of Antibiotic Therapy per 1000 Patient Days
Bacterial (15)	1484 +/- 252*
Bacterial + Viral (13)	1661 +/-387*
Viral (18)###	1188 +/- 641 **

+ vs ++: p = 0.003

###:Antibiotics discontinued in only 4 of 18 within 48 hrs of MD receipt of test results showing virus detection by PCR and low serum PCT



Critical Care Research and Practice

Volume 2014 (2014), Article ID 307817, 4 pages http://dx.doi.org/10.1155/2014/307817

Research Article

*P

Table 1: Influence of the presence and absence of an infectious diseases specialist (IDS) on antibiotic use decisions by the ICU Multidisciplinary Team at Providence Portland Medical Center.

	In the presence of IDS	In the absence of IDS
Number of rounding sessions	61	51
Total number of patients receiving antimicrobials pre-rounds	384	352
Total days of therapy (DOT)		
Prerounds	669	593
Postrounds	511	572
Difference	-158	-21
Decrease DOT/patient	-0.41*	-0.06*
Total cost of antimicrobial regimen, \$		
Prerounds	17,521	15,153
Postrounds	13,749	14,426
Difference	-3,772	-727
Decrease cost/patient	-9.82*	-2.10*
Number of multidisciplinary rounds where DOT		
Increased	0	14
Did not change	15	14
Decreased	46*	23*

PPMC AMS Success: Pneumonia, Sepsis, and SSTIs

Days of antibiotic therapy per 1000 patient days							
	2013	2014 target	1 st Q	2 nd Q	3 rd Q	4 th Q	1sr Q 2015
Total PHS System	781	NA	858	873	927	865	794
PPMC	780	708	598	572	577	599	483
Pneu.	282	254	168****	192	141	186	102****
Seps.	362	326	353	272	309	254	269
SSTI	136	129	77	108	126	159	111

**** CAP Diagnostic Study in Progress

Physician Education

- The dynamic duo: identification of the pathogen and serum PCT levels
- How to help MDs use the information in a correct and timely fashion ?
 - Need faster communication
 - Require physicians to indicate a diagnosis for each antibiotic order; 48 hr. reassessment
 - Embed ID MD in ICU rounds

How To Be More Successful ?

- CMS et al: Provide reimbursement for expense of molecular diagnostics that favorably impact antimicrobial stewardship
- Rapid diagnostics requires rapid help in interpretation via direct communication with the patient's physician provider
- In short, achievement of full potential of emerging biomarkers and pathogen detection will require TIME, PEOPLE, AND MONEY







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Promoting Antibiotic Stewardship

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Q-HIP® Hospital Value-Based Payment Program

Rewards hospitals for meeting quality, patient safety, outcomes, and patient satisfaction measures

Patient Safety	Health Outcomes			
 CMS Meaningful Use IHI - Surgical Safety and Med Rec NQF Safe Practice 15 (Discharge Systems) NQF, AHRQ and ACEP ED Measures IHI Post Hospital Care Follow Up NQF Perinatal Care CDC CRE Toolkit 	 JC Stroke (STK) JC Influenza Immunization CMS/JC Perinatal NHSN Measures CMS Readmission Measures ACC-NCDR PCI Measures STS CABG Measures 			
Patient Satisfaction	Bonus			
H-CAHPS Survey Results	 JC Perinatal Measure Set IHI Patient Teach Back Registry/Patient Safety Collaboration AHRQ Patient Safety Culture Survey ACP Care Coordination Agreement 			





Antibiotic Stewardship Program Measure

<u>New Metric</u>: We are proposing the addition of a measure based on the CDC's new *Core Elements of Hospital Antibiotic Stewardship Programs*. The measure would adopt the following structure:

- Description: Hospital has implemented the Core Elements described in the CDC Core Elements of Hospital Antibiotic Stewardship Programs guide
- Compliance Criteria: To answer "Yes", the program must incorporate:
- Leadership Commitment: The hospital has designated a leader (e.g., physician, pharmacist, etc.) responsible for program outcomes
 of antibiotic stewardship activities at the hospital
- Accountability: There is a single leader responsible for program outcomes
- Drug Expertise: There is a single pharmacist leader responsible for working to improve antibiotic use
- Action: At least one recommended action has been implemented, such as systemic evaluation of ongoing treatment need after a set
 period of initial treatment (i.e. "antibiotic time out" after 48 hours)
- Tracking: Antibiotic prescribing and resistance patterns are monitored
- · Reporting: Regular reporting information on antibiotic use and resistance is provided to doctors, nurses and relevant staff
- · Education: Education of clinicians about resistance and optimal prescribing is ongoing
- Compliance Example:
- Written policies and procedures showing that an Antibiotic Stewardship program *is in place during the Measurement Year* to ensure optimal antibiotic prescribing and limit overuse and misuse of antibiotics. The *program includes* all the Core Elements from the *CDC Core Elements of Hospital Antibiotic Stewardship Programs guide.*

Anthem.

http://www.cdc.gov/getsmart/healthcare/implementation/core-elements.html











