

How to develop, implement and evaluate prescribing guidelines and post-prescription review

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American Board in Internal Medicine, Geriatric and Infectious Diseases

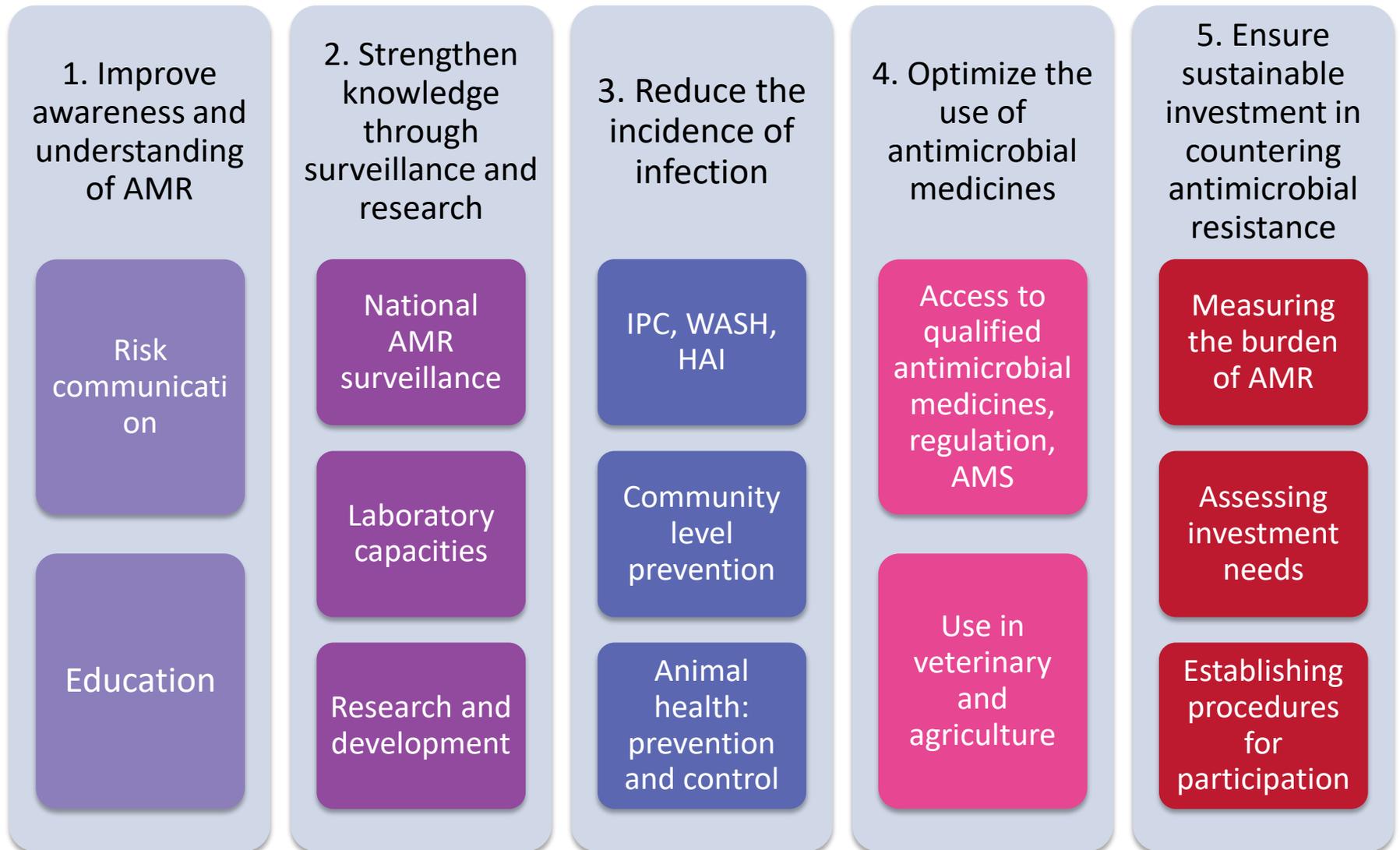
Chairperson of HIV management team in MOH

Chairperson of Antibiotic committee in MOH, Kingdom of Bahrain

Chairperson of the geriatric services in secondary care

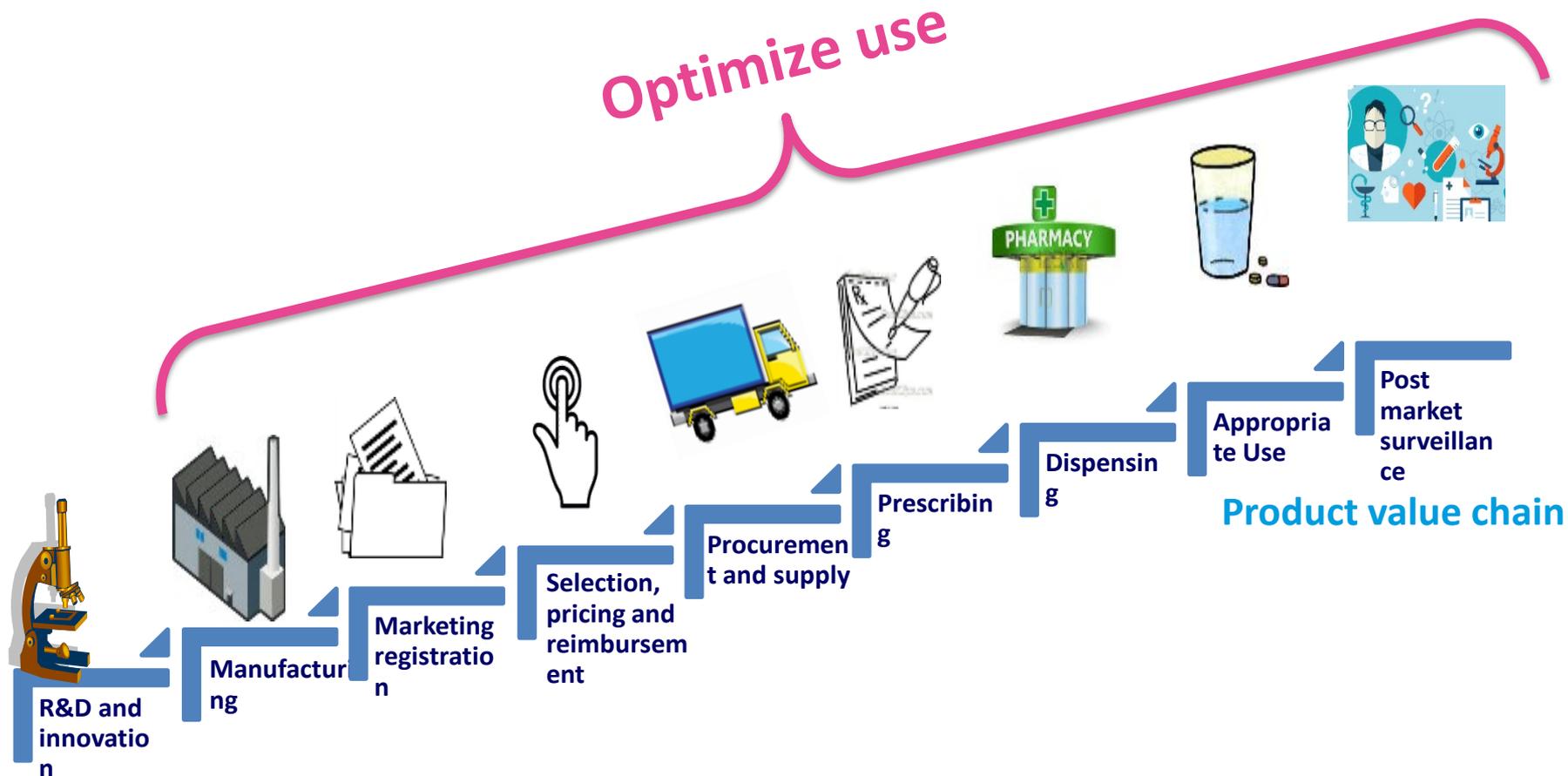
Associate Professor in Arabian Gulf University

Global Action Plan: 5 strategic objectives



Optimize the use of antimicrobials

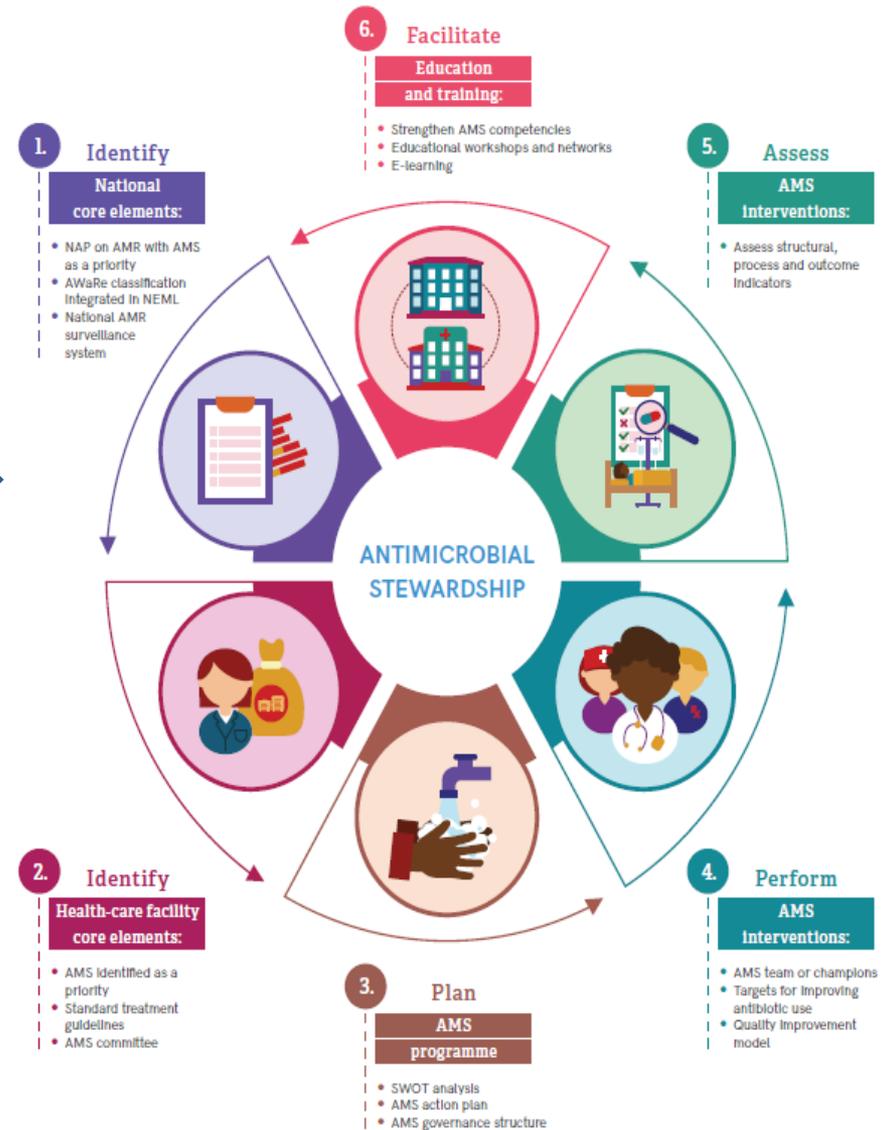
Integrated approach to Stewardship, AWaRe & Surveillance



Optimize use: Integrated approach to Stewardship, AWaRe & Surveillance

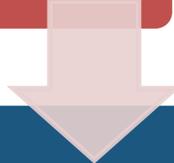
- National Action Plan to Combat Antimicrobial Resistance in Hashemite Kingdom of Jordan (2018-2022):
 - **Vision:** Reduction of mortality, morbidity and economic impact of AMR in Jordan.
- By the year 2022:
 1. **10% reduction in AMR morbidity**
 2. **20% reduction in antimicrobial consumption in humans**
 3. 30% reduction in antimicrobial consumption in animals
 4. **30% reduction in diseases due to multidrug resistant organisms**
 5. **40% increase of public knowledge on AMR and awareness of appropriate use of antimicrobials**

WHO practical toolkit: AMS in health-care facilities (2019)



Antimicrobial stewardship (AMS) – definition, objective, action & outcome

Definition: A strategy & set of actions to promote using antimicrobials responsibly



Objective: To ensure effective antibiotic treatment for patients today & tomorrow



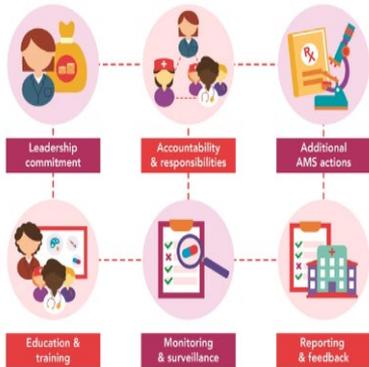
Action: Change prescribing practices and Abx use:
No ABx when not needed; old ABx when sufficient;
new ABx only when necessary



Outcome: Improve patient outcomes, prolong use of existing Abx, save lives & health-care costs, reduce emergence & spread of AMR

Hospital AMS Programme

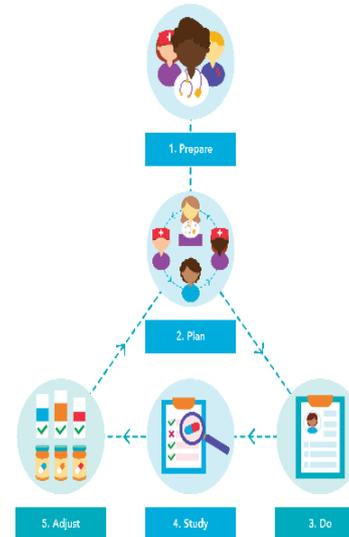
2. Hospital core elements



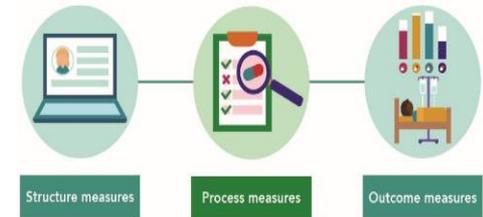
3. Planning AMS programmes

	HELPFUL	HARMFUL
INTERNAL/PRESENT FACTORS	Strengths Core elements: • AMR and AMS are a leadership priority. • IPC programme/committee is active. Human resources: • There is enthusiasm for AMS in the facility/wards. • There is clinical knowledge of AMS. Antimicrobial use and resistance data: • Prescription audit is conducted in one ward. • Facility aggregate antibiogram is available. AMS activities: • A pharmacist is involved in some AMS activities in one ward.	Weaknesses Core elements: • No medical record or prescription pad is available. Human resources: • No dedicated health care professional is available to lead the AMS team. Antimicrobial use and resistance data: • The supply of microbiology reagents is poor. • The supply of antibiotics is poor. AMS activities: • Health-care professionals have competing priorities and little time for AMS work.
	Opportunities Core elements: • Active implementation of the NAP on AMR • Increasing national awareness of AMR and its consequences for health Human resources: • Incorporating AMS responsibility into the IPC committee Antimicrobial use and resistance data: • Funds for conducting a facility PPS AMS activities: • Presenting findings from AMS activities to other wards/health-care professionals	Threats Core elements: • Unstable access to essential antibiotics • Increased costs for antibiotics • Prioritisation of issues other than AMS in the facility • Low facility budget Human resources: • Too many nonfunctional committees in the health-care facility Antimicrobial use and resistance data: • Increasing AMR rates, including carbapenem-resistant Enterobacteriaceae (CRE) AMS activities: • Opposition from clinical leaders
EXTERNAL/FUTURE FACTORS	SWOT	

4. Performing AMS Interventions



5. Assessing AMS Interventions



Health-care facility core elements



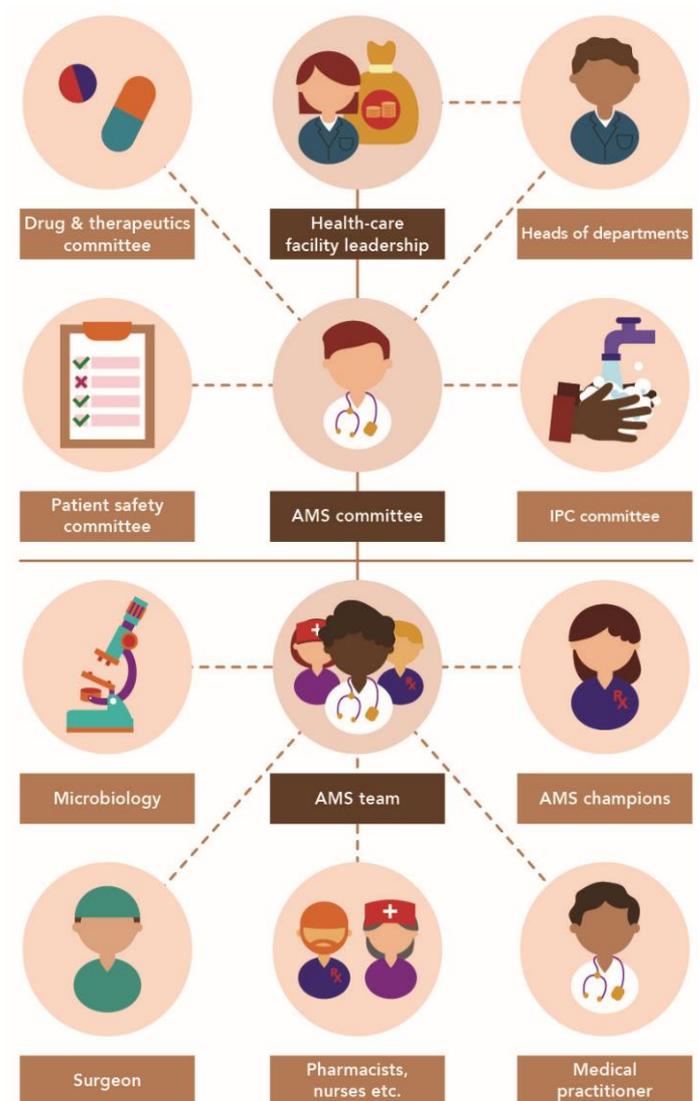
- ✓ Leadership – prioritize AMS
- ✓ Human resources
- ✓ Education and training
- ✓ Treatment guideline
- ✓ AMS interventions
- ✓ Monitoring ABx
- ✓ Reporting and feedback
- ✓ Links to IPC and WASH

Planning AMS programmes

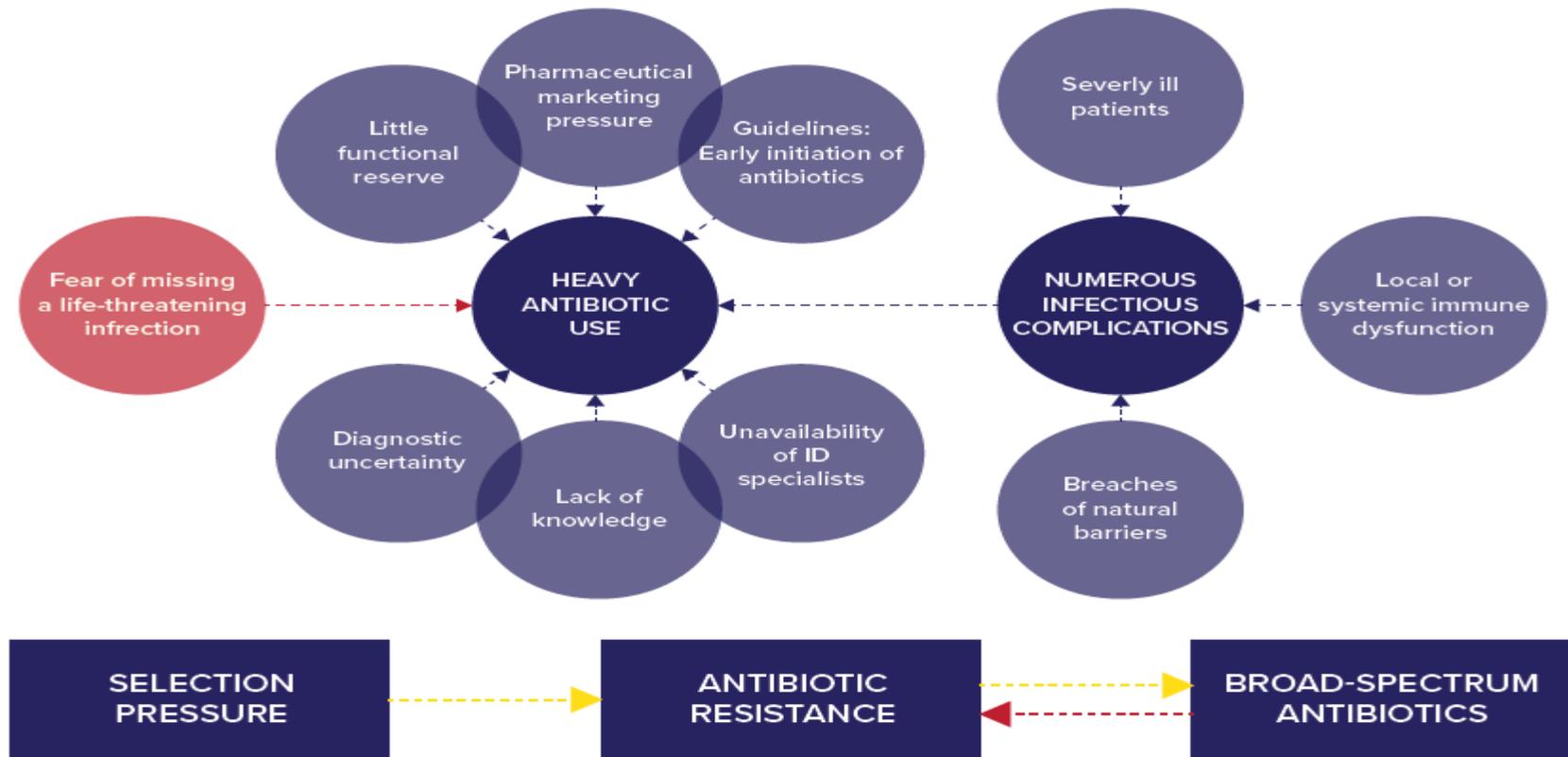
Situational or SWOT analysis	Conduct a SWOT analysis: <ul style="list-style-type: none"> Structures, policies and guidelines Human resources Data: antimicrobials, resistance AMS activities
Facility AMS action plan	To ensure accountability, prioritize activities and measure progress

Governance

- ✓ Responsibilities and accountability
- ✓ AMS team and/or AMS champions
- ✓ Link to other programmes/ committees

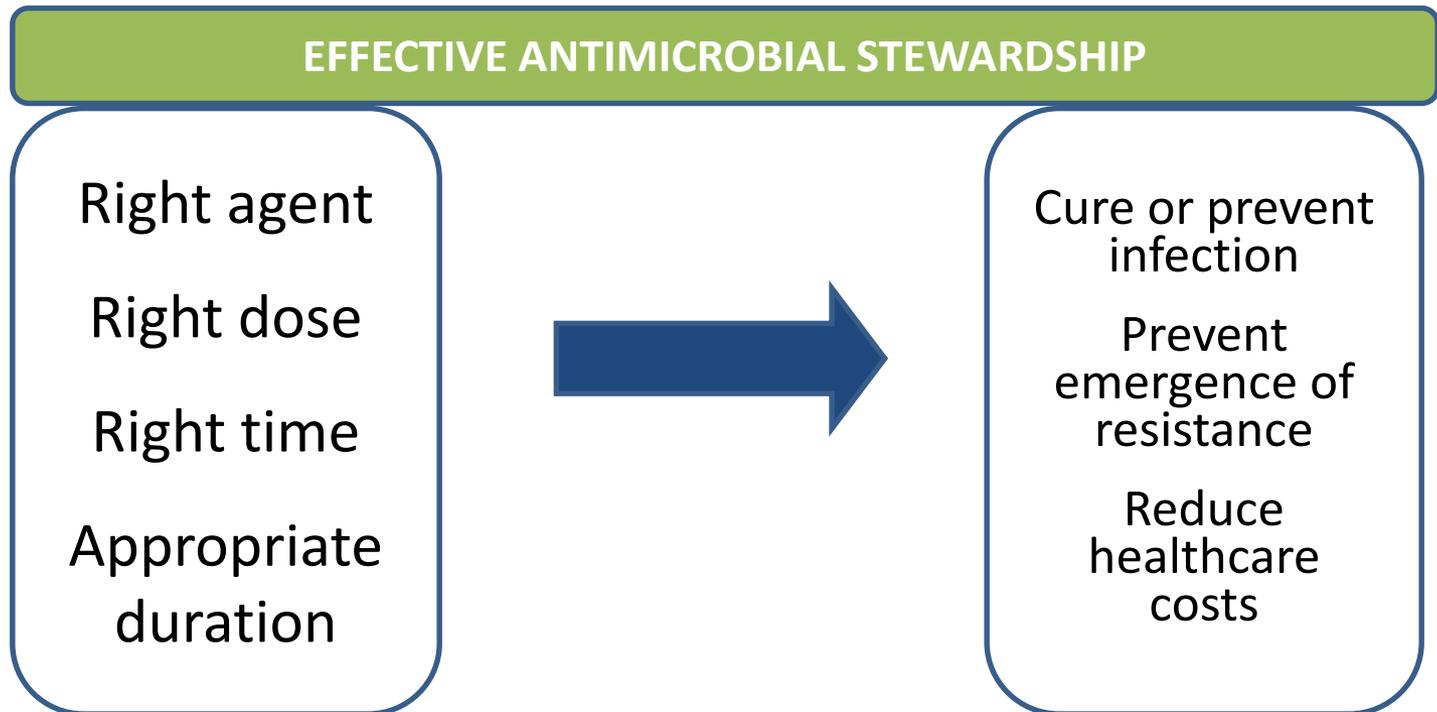


WHAT ARE THE DRIVERS FOR THE USE AND MISUSE OF ANTIBIOTICS?



What are the Goals of Antimicrobial Stewardship?

- Antimicrobial stewardship is an organizational or healthcare system-wide approach to promoting and monitoring judicious use of antimicrobials to preserve their future effectiveness^{1,2}
- Antimicrobial stewardship is a key strategy to overcome resistance by the careful and responsible management of antimicrobial use



- **1.** British Society for Antimicrobial Chemotherapy. A Practical Guide to Antimicrobial Stewardship in Hospitals. Available at: <http://bsac.org.uk/news/practical-guide-to-antimicrobial-stewardship-in-hospitals/>. Last accessed September 29, 2015.

Key Components of an Antimicrobial Stewardship Program

- A multidisciplinary antimicrobial stewardship program should include:
 - Infectious diseases physician and pharmacist and other key stakeholders as determined by the institution
 - Policy statement
 - Physician-directed or supervised multidisciplinary program with a minimum of one or more members trained in antimicrobial stewardship
- **Two core strategies were recommended**
 - **Prospective audit with intervention and feedback**
 - **Formulary restriction and authorization**
- Other recommended strategies
 - Education
 - Guidelines and clinical pathways
 - Order forms
 - Streamlining/de-escalation
 - Dose optimization
 - Intravenous-to-oral conversion

Implementing an Antibiotic Stewardship Program: Guidelines by the Infectious Diseases Society of America and the Society for Healthcare Epidemiology of America

Tamar F. Barlam,^{1,a} Sara E. Cosgrove,^{2,a} Lilian M. Abbo,³ Conan MacDougall,⁴ Audrey N. Schuetz,⁵ Edward J. Septimus,⁶ Arjun Srinivasan,⁷ Timothy H. Dellit,⁸ Yngve T. Falck-Ytter,⁹ Neil O. Fishman,¹⁰ Cindy W. Hamilton,¹¹ Timothy C. Jenkins,¹² Pamela A. Lipsett,¹³ Preeti N. Malani,¹⁴ Larissa S. May,¹⁵ Gregory J. Moran,¹⁶ Melinda M. Neuhauser,¹⁷ Jason G. Newland,¹⁸ Christopher A. Ohl,¹⁹ Matthew H. Samore,²⁰ Susan K. Seo,²¹ and Kavita K. Trivedi²²

Goal of the 2016 guidelines

- Provide a guideline that diverse stakeholders find useful
- More detailed, implementation-oriented focus compared with prior guidelines
- Expand scope
 - e.g. pharmacologic optimization, the role of microbiology-relevant interventions, and metrics by which to assess programs
 - Reference special populations, settings
- Use the GRADE system to rank the guideline's recommendations and the level of evidence

Select Examples

- Does the Use of Preauthorization and/or Prospective Audit and Feedback Interventions by ASPs Improve Antibiotic Utilization and Patient Outcomes?
We recommend preauthorization and/or prospective audit and feedback over no such interventions (strong recommendation, moderate-quality evidence).
- Should ASPs Develop and Implement Facility-Specific Clinical Practice Guidelines for Common Infectious Diseases Syndromes to Improve Antibiotic Utilization and Patient Outcomes?
We suggest ASPs develop facility-specific clinical practice guidelines coupled with a dissemination and implementation strategy (weak recommendation, low-quality evidence)
- Should ASPs Implement Interventions to Improve Antibiotic Use and Clinical Outcomes That Target Patients With Specific Infectious Diseases Syndromes?
We suggest ASPs implement interventions to improve antibiotic use and clinical outcomes that target patients with specific infectious diseases syndromes (weak recommendation, low-quality evidence)

Select examples continued

- Should ASPs Implement Interventions Designed to Reduce the Use of Antibiotics Associated With a High Risk of CDI?

We recommend antibiotic stewardship interventions designed to reduce the use of antibiotics associated with a high risk of CDI compared with no such intervention (strong recommendation, moderate-quality evidence)

- In Hospitalized Patients Requiring Intravenous (IV) Antibiotics, Does a Dedicated Pharmacokinetic (PK) Monitoring and Adjustment Program Lead to Improved Clinical Outcomes and Reduced Costs?

We recommend that hospitals implement PK monitoring and adjustment programs for aminoglycosides (strong recommendation, moderate-quality evidence).

We suggest that hospitals implement PK monitoring and adjustment programs for vancomycin (weak recommendation, low-quality evidence).

Select examples continued

- Should ASPs Implement Interventions to Increase Use of Oral Antibiotics as a Strategy to Improve Outcomes or Decrease Costs?

We recommend ASPs implement programs to increase both appropriate use of oral antibiotics for initial therapy and the timely transition of patients from IV to oral antibiotics (strong recommendation, moderate-quality evidence)

- Should ASPs Advocate for Rapid Diagnostic Testing on Blood Specimens to Optimize Antibiotic Therapy and Improve Clinical Outcomes?

We suggest rapid diagnostic testing in addition to conventional culture and routine reporting on blood specimens if combined with active ASP support and interpretation (weak recommendation, moderate-quality evidence)

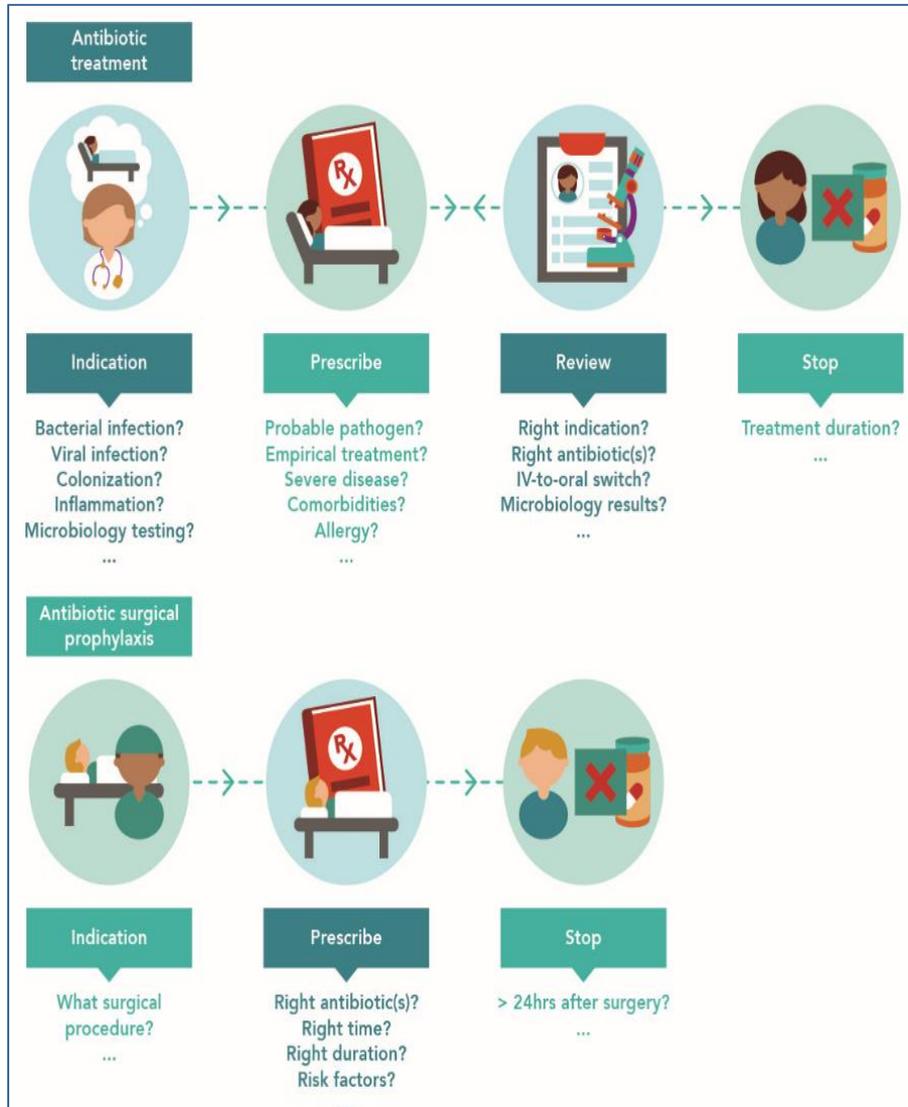
- Should ASPs Implement Interventions to Reduce Antibiotic Therapy to the Shortest Effective Duration?

We recommend that ASPs implement guidelines and strategies to reduce antibiotic therapy to the shortest effective duration (strong recommendation, moderate-quality evidence)

Current evidence on hospital antimicrobial stewardship objectives: a systematic review and meta-analysis

- Overall quality of evidence was low, but they concluded there was enough support for some interventions:
 - Following guidelines in administering empiric antibiotics
 - IV to PO
 - Antibiotic restrictions
 - ID consultations
 - therapeutic drug monitoring
 - De-escalation of therapy
- Conclusion: The overall evidence for these interventions shows significant benefits for clinical outcomes, adverse events, costs, resistance rates, or combinations of these. However, the included studies were generally of low quality.

Performing AMS interventions



Evidence based AMS interventions

Interventions (examples)	
Education	<ul style="list-style-type: none"> Formal/ informal Treatment guidelines
Feedback	<ul style="list-style-type: none"> Audit with feedback Ward rounds
Structure	<ul style="list-style-type: none"> Self-revision by prescriber Computerized order entry
Restriction	<ul style="list-style-type: none"> Pre-authorization Automatic stop orders

Davey P. Interventions to improve antibiotic prescribing practices for hospital inpatients. Cochrane Database Syst Review 2013 Apr 30;4: CD003543.

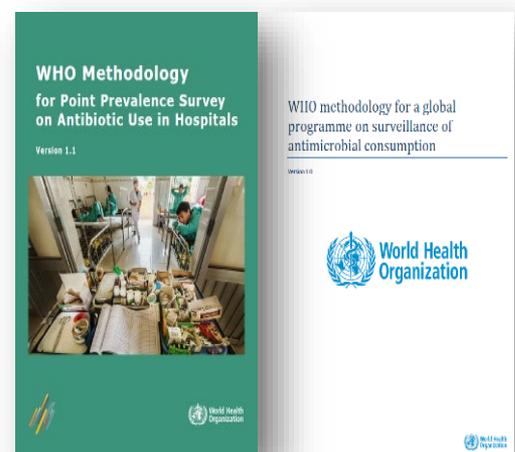
How to do it

- 1. what are the most common infection in your institution ?
- 2. What are the steps to be followed ?
- 3. What are the elements to be addressed ?
- 4. How to analyze ? The goals
- 5. How to follow up ?

Assessing AMS programmes



- **Baseline:** Measure the **quantity** and **quality** of **antibiotic prescribing and use**, to identify priority areas for AMS interventions
- **Goal:** to **compare** results within a hospital, department or ward **over time**; **AB prescribing and use, patient outcomes etc**



- ✓ **Structure measures:**
core elements
- ✓ **Outcome measures:**
ABX use, patient
outcomes
- ✓ **Process measures:**
proportions e.g. of
pneumonia patients
receiving appropriate
antibiotic treatment

Education & Training

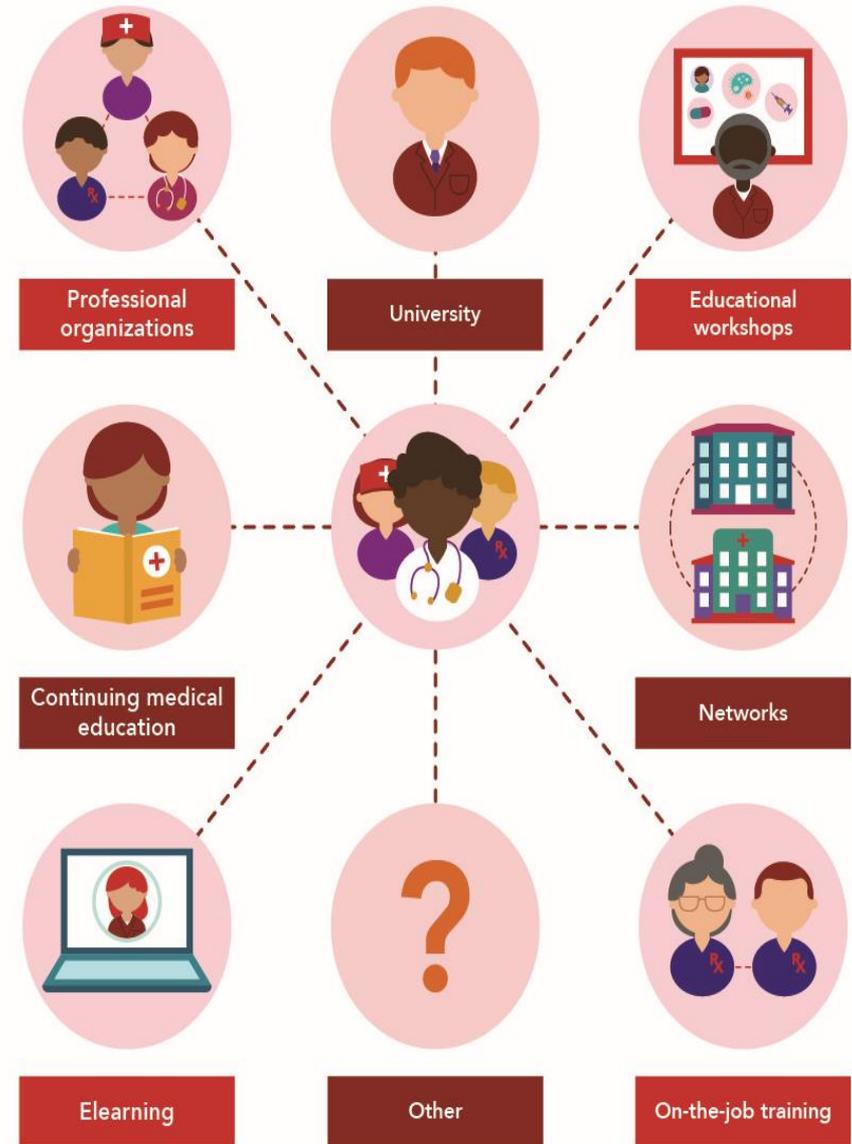
AMS competencies

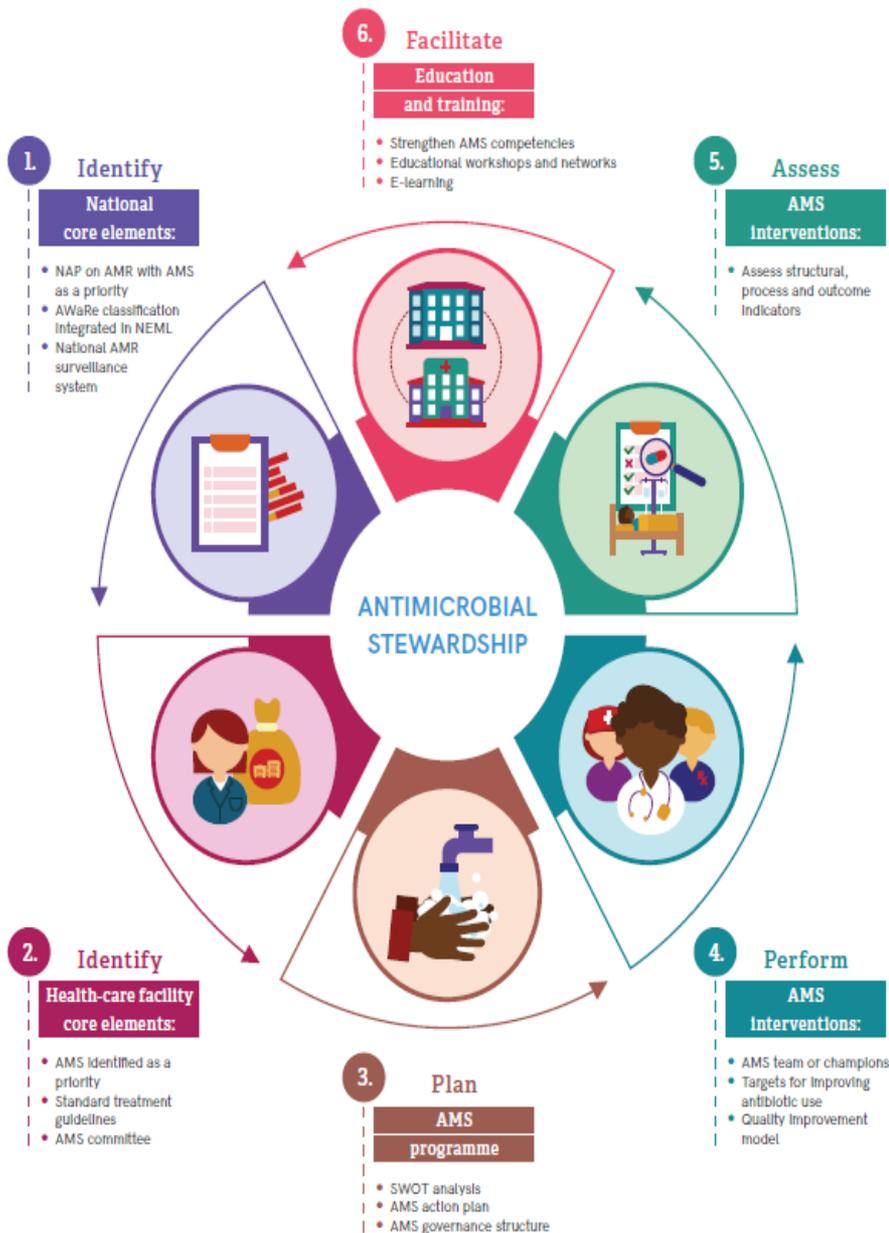
- ✓ Antibiotics
- ✓ Microbiology
- ✓ Infection management
- ✓ Plan and perform AMS interventions
- ✓ Monitor AMS interventions/ ABx use

Face to face workshops
Online e-learning resources

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04/02/2020





Summary

- 6 core components on AMS for health-care facilities
- Step-wise approach
- Build on what is existing, start small
- Build competency and team
- Quality improvement cycle

Elements to Consider Before Intervening

- Assess organizational culture
- Address organizational barriers
- Tailor education based on observed practice deficits
- Develop a usable intervention
- Solicit feedback from nurses and identify a nurse champion to develop education on proposed intervention
- Consider train the trainer or other team-based modalities
- Secure nursing and physician stakeholder support



Stewardship Goals

Reducing length of stay

Reducing duration of surgical prophylaxis

Restricting or limiting specific antimicrobials to reduce inappropriate use

Possible Unintended Consequences

Increasing rates of readmission

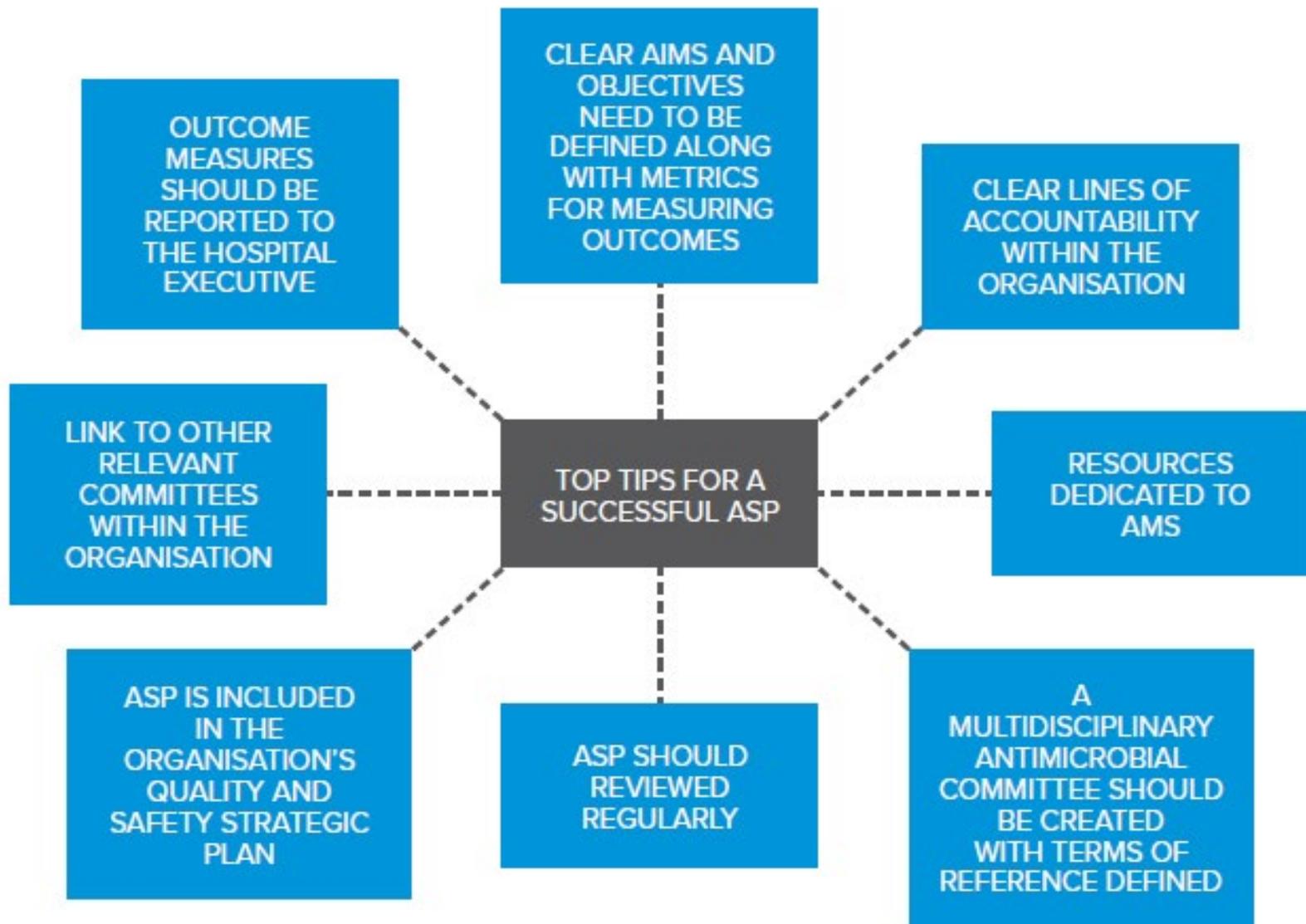
Increasing rates of surgical site infections

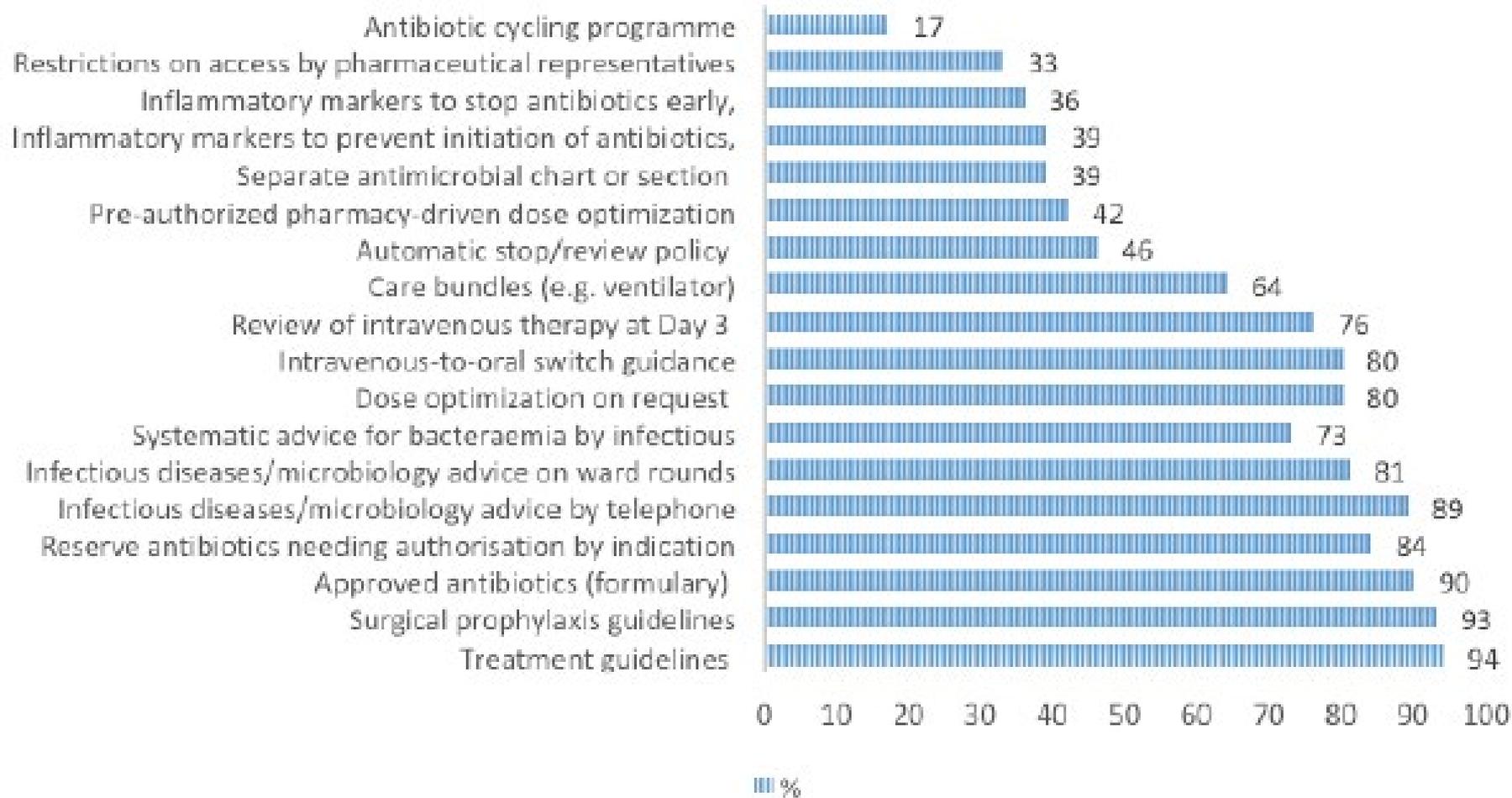
Increasing use of non-restricted antimicrobials (e.g. "squeezing the balloon")

Delaying doses of antimicrobials due to restriction processes

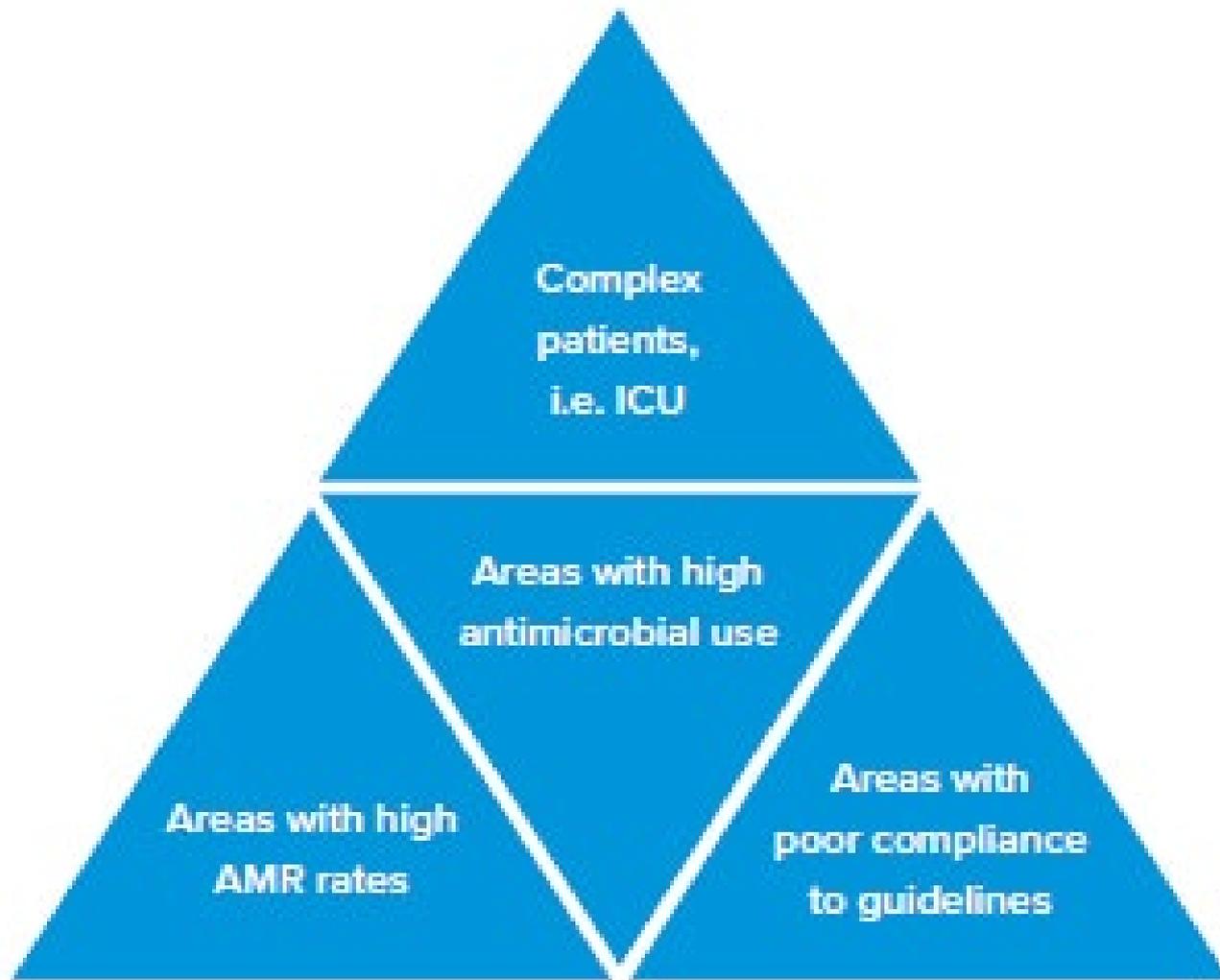
SOURCE

http://www.idsociety.org/Guidelines/Patient_Care/IDSA_Practice_Guidelines/Antimicrobial_Agent_Use/Implementing_an_Antibiotic_Stewardship_Program/





Results of a global survey on types of AMS interventions employed as part of an ASP



Suggested priority areas for targeting ASP

HOW DO I ASSESS THE EFFECTIVENESS OF AN ANTIMICROBIAL STEWARDSHIP PROGRAMME?

- There are many ways to assess the effectiveness of an ASP, including:
 - • Audit of compliance with guidelines
 - • Audit documentation – e.g. indication, stop/review date, 48-72 hour review
 - • Audit time to 1st dose of antibiotic in sepsis
 - • Monitor antibiotic consumption data, including benchmarking to similar institutions
 - • Monitor antibiotic expenditure data
 - • Monitor stewardship interventions and acceptance rates
 - • Review adverse events in relation to antimicrobials
- Defined outcome measures should be defined as part of an organizations ASP strategy.

Suggested Measures

Measurement Area	Measure
Antibiotic consumption	<ul style="list-style-type: none">• Days of therapy (DOT) per 1,000 patient days—overall and for specific agents or groups of agents• Defined daily dose (DDD) per 1,000 patient days (if DOT not available)• Standardized Antibiotic Administration Ratio*
Process measures	<ul style="list-style-type: none">• Provision of indication with each antibiotic start• Percentage of cases where therapy is appropriate (especially for serious infections, such as sepsis)• Appropriate Treatment of Methicillin-Sensitive Staphylococcus aureus (MSSA) Bacteremia• Frequency at which de-escalation occurs• Timely cessation of antibiotics given for surgical prophylaxis• Antibiotics not prescribed to treat asymptomatic bacteria• Appropriate cultures obtained before starting antibiotics• Adherence to hospital-specific guidelines• Acceptance of ASP recommendations• Frequency of performance of antibiotic time outs or reviews• Timely administration of appropriate antibiotics in cases of suspected sepsis

Suggested Measures continued

Measurement Area	Measure
Outcome measures	<ul style="list-style-type: none">• Length of stay• Cure of infection• Risk-adjusted mortality• Hospital readmissions for select infections• Hospital-onset <i>C. difficile</i> infections*• Adverse drug reactions (number/percentage/rate)• Antimicrobial resistance- focusing on hospital onset cases would most likely best reflect the impact of ASPs• Provider-level measures if available (e.g., treatment of <i>S. aureus</i> and bloodstream infections)
Financial	<ul style="list-style-type: none">• Antibiotic cost per patient day• Antibiotic cost per admission• Total hospital cost per admission

*NQF-endorsed measure

Shorter is Better

Syndrome	Short Course Studied (days)	Long Course Studied (days)	Result	Updated IDSA guidelines?
Acute bacterial sinusitis	5	10	Equal	2013
COPD exacerbation	≤5	≥7	Equal	2018 Gold guidelines
Intra-abdominal infection	4	10	Equal	2010, in development
CAP	3-5	7-10	Equal	2007, in development
HAP/VAP	≤5	10-15	Equal	2016
Cystitis/Pyelo	3-5/5-7	10-14	Equal	2011
SSTI	5-6	10-14	Equal	2014

Adapted from B. Spellberg. J Hosp Med 2018

And Yet...

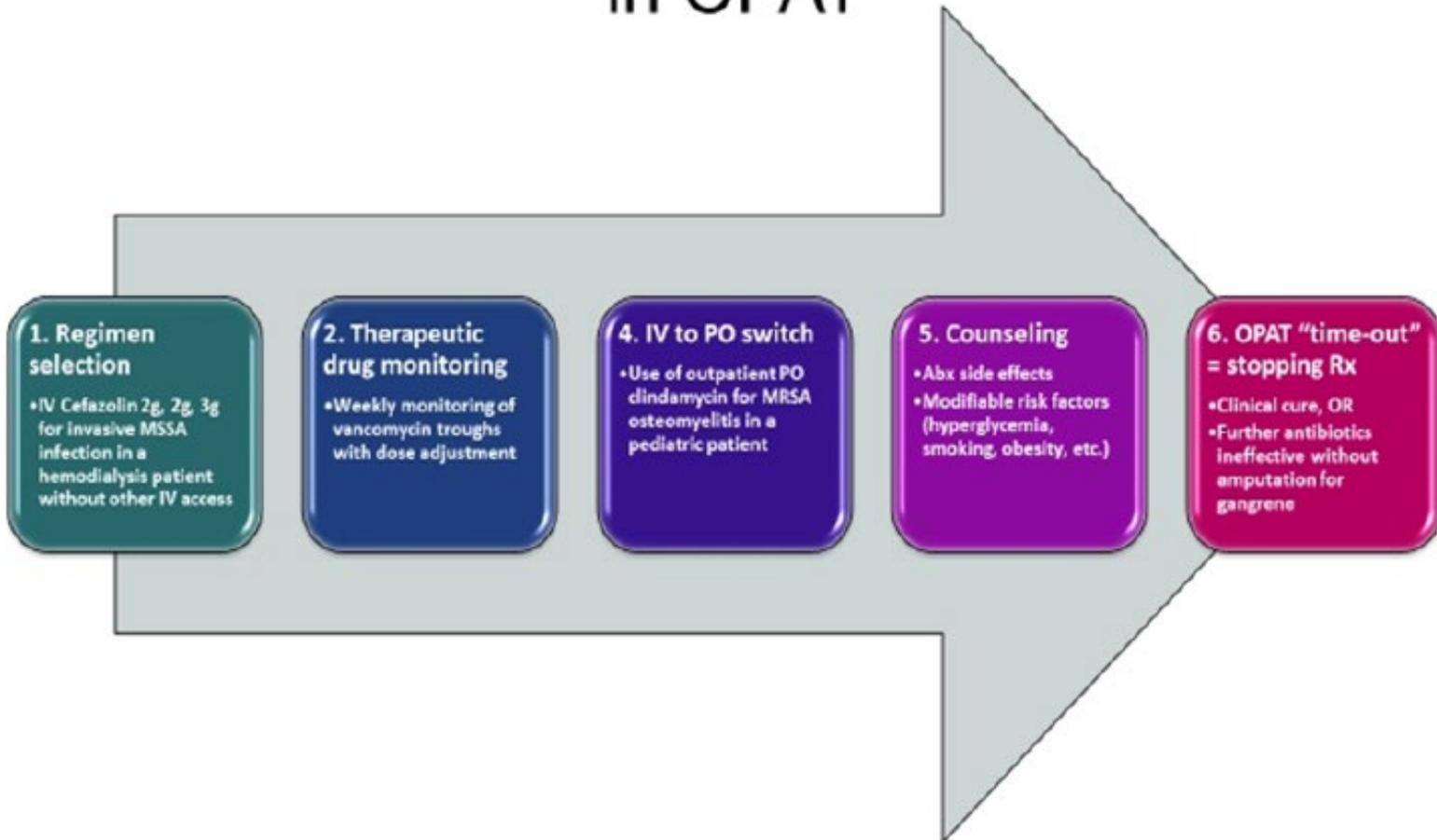
Syndrome	Admissions with durations >7 days*
PNA	761 (76%)
UTI	1324 (79%)
Intra-abdominal	594 (89%)
SSTI	828 (90%)

*admissions with both inpatient AND post-discharge antimicrobials

Discharge Best Practices

- ✓ Establish institutional guidelines with evidence-based, shortest effective durations
 - “shorter is better”
- ✓ Education
- ✓ Discharge audit/feedback
- ✓ Integrate data to capture both inpatient DOT and discharge prescriptions
- ✓ Document appropriately to reduce excess duration at d/c (especially ID consultants)

Continuum of Stewardship Activities in OPAT



IV to PO – Benefits

- ↓ LOS^{1, 3, 5, 6}
- ↑ early ambulation, patient comfort
- ↓ Antibiotic duration¹
- ↓ 30-day readmission¹
- ↓ *C. difficile* rates¹
- ↓ Cost^{2, 3, 6, 7}
- ↓ Incidence of line infections⁴
- ↓ Nursing workload



¹ Kutzhalts KE et al. Clin Ther 2016;38:1750-8

² Davis S, et al. Clin Infect Dis 2005;41:139-143

³ Onidvari K et al. Respir Med 1998;92:1032-9

⁴ Izing RB et al. J Antimicrob Chemother 1998;42:107-11

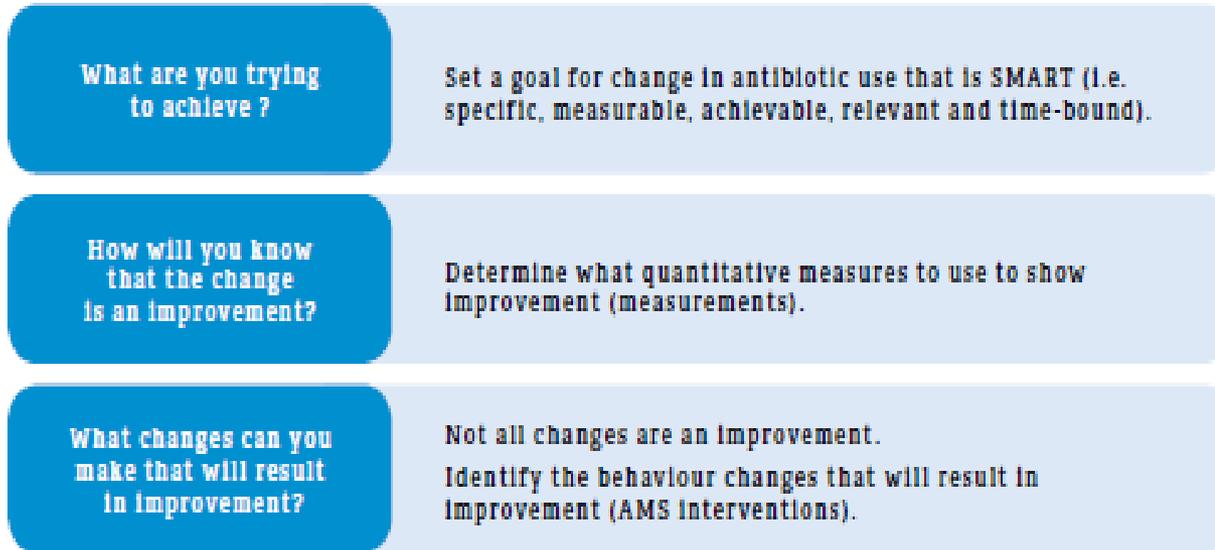
⁵ Park SM et al. Infect Chemother 2017;31-7

⁶ Kuri J, et al. Am J Health Syst Pharm 2001;58:2109-15

⁷ Falabella A et al. Am J med 1991;91:662-70

Implementation – a systematic approach

Questions to address when applying the quality improvement model for AMS interventions



Implementation: Measure baseline, improve, measure again, report back

PERFORMING AMS INTERVENTIONS



Adapting Evidence-based guidelines to the local setting

Local guidelines

HEALTH-CARE FACILITY CORE ELEMENTS

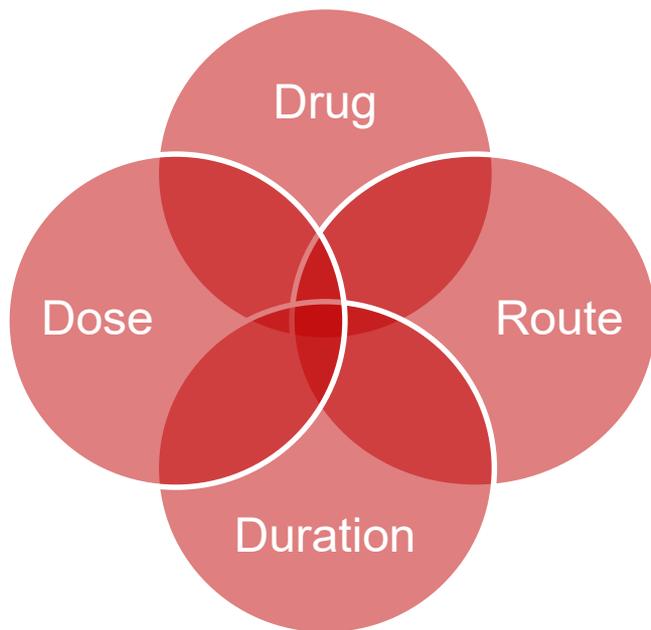
10. Up-to-date standard treatment guidelines

The health-care facility has available, up-to-date recommendations for infection management based on international/national evidence-based guidelines and local/national susceptibility patterns (where possible), to assist with antibiotic selection for common clinical conditions (indication, agent, dose, route, interval, duration). A process is in place for regular review and updating of the guidelines based on new evidence or other external input.

Yes

No

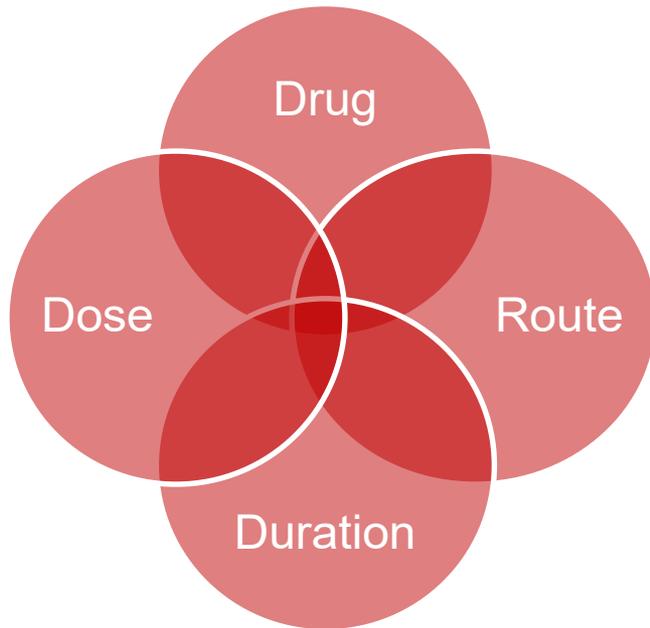
Why develop local guidelines for common infections?



Principles of Antimicrobial Stewardship

- Reduce variability in clinical practice
- Ensure use of best available evidence in patient care
- Educational tool
- Used in metrics
- Empowers other team members (pharmacists, nurses, etc.)
- Cost-savings, reduced length of stay

Why develop **local** guidelines for common infections?



Principles of Antimicrobial Stewardship

Why **Local**?

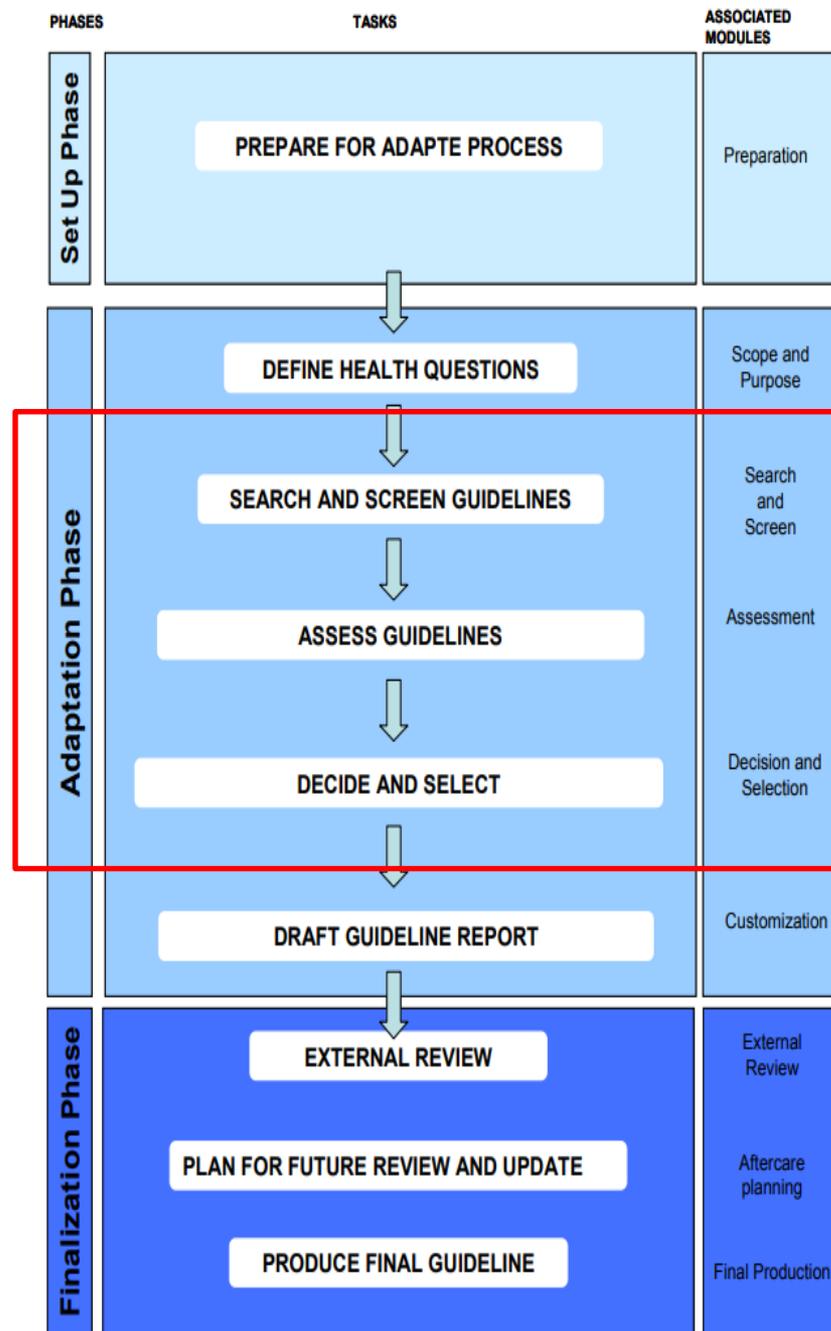
- Local epidemiology
- Drug availability/formulary
- Relevant diagnostics
- Designed with implementation in mind
- Team-building

ADAPTE PROCESS

- Other frameworks include: PGEAC, SGR, AAP, CAN-IMPLEMENT, and Adapted ADAPTE
- One group used evidence-based guidelines developed using GRADE to simplify this process.
CHEST2014; 146(3): 727 - 734

The ADAPTE Collaboration (2009). The ADAPTE Process: Resource Toolkit for Guideline Adaptation
<http://www.g-i-n.net>

Summary of the ADAPTE process



One approach

Adaptation

- Identify guidelines to adapt
- Develop writing group
 - Include stakeholders
 - Iterative process

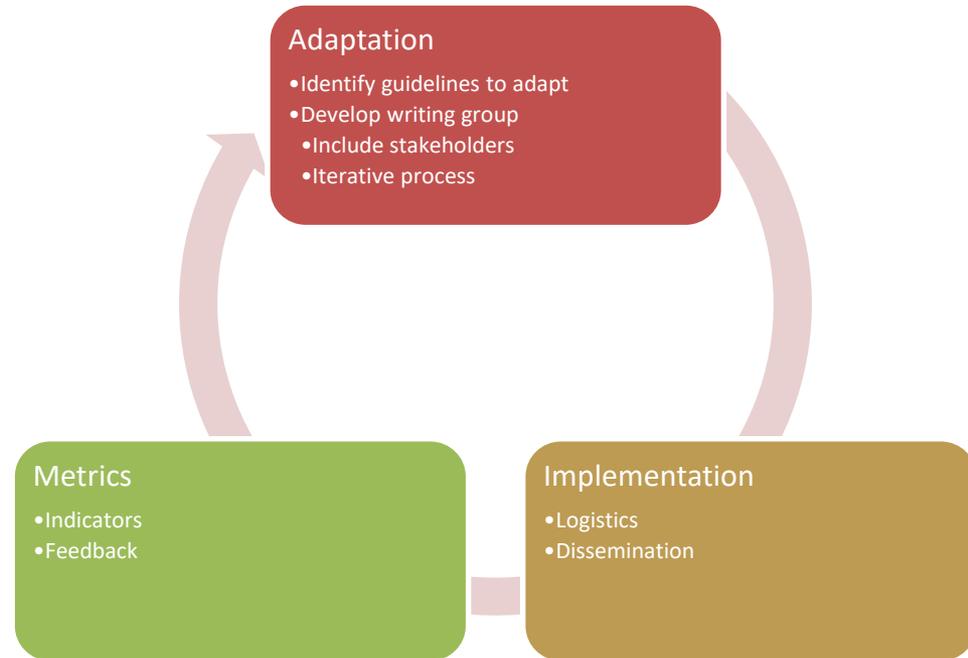
Implementation

- Logistics
- Dissemination
- Integration into existing workflow

Metrics

- Indicators
- Feedback

One approach



Metrics: Local guideline

Process Metrics	Outcome Metrics
Adherence to guidelines:	Antibiotic utilization
<ul style="list-style-type: none">• Accurate diagnosis• Empiric antibiotic choice• Adjustment of antibiotic with microbiologic data• Timing of antibiotic (surgical prophylaxis)• Duration of therapy•	Patient outcomes
	Cost (antibiotic saved)
Utilization of orderset (or order sheets, etc.)	Length of stay
	Adverse drug effects

Stewardship target:
Management of common infections

New evidence-based guidelines released

Diagnosis and Treatment of Adults with Community-acquired Pneumonia

An Official Clinical Practice Guideline of the American Thoracic Society and
Infectious Diseases Society of America

② Joshua P. Metlay*, Grant W. Waterer*, Ann C. Long, Antonio Anzueto, Jan Brozek, Kristina Crothers, Laura A. Cooley, Nathan C. Dean, Michael J. Fine, Scott A. Flanders, Marie R. Griffin, Mark L. Metersky, Daniel M. Musher, Marcos I. Restrepo, and Cynthia G. Whitney; on behalf of the American Thoracic Society and Infectious Diseases Society of America

THIS OFFICIAL CLINICAL PRACTICE GUIDELINE WAS APPROVED BY THE AMERICAN THORACIC SOCIETY MAY 2019 AND THE INFECTIOUS DISEASES SOCIETY OF AMERICA AUGUST 2019

- Empiric antibiotic therapy
- Defined specific risk factors for infection with MDROs --- NO MORE HCAP
- Duration = 5 days

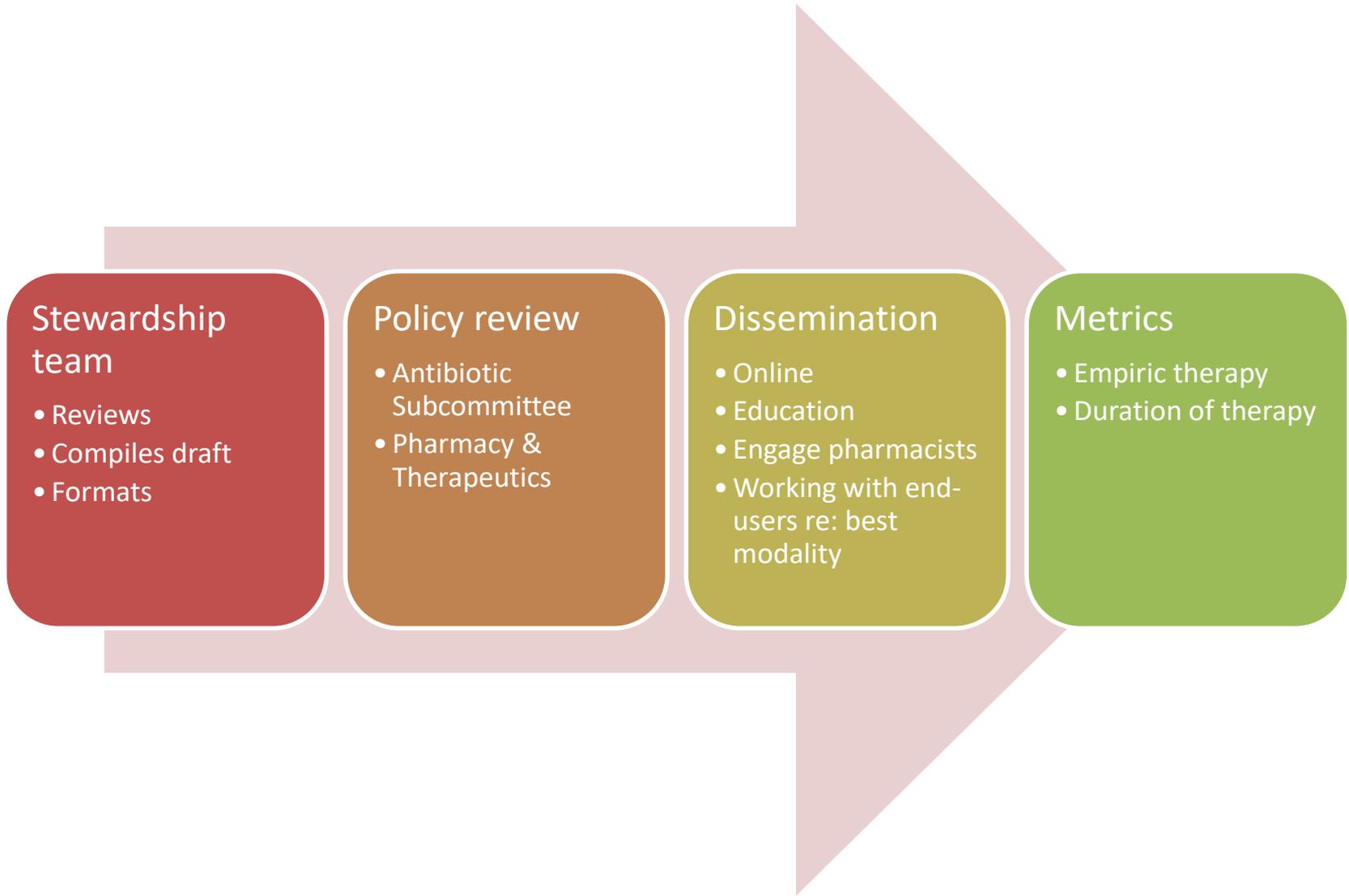
Excess duration of therapy is associated with patient harm

- 43 Michigan hospitals January 2017 – 2018
- 6,481 patients hospitalized with pneumonia
- Excess days = (expected treatment duration – actual duration)
- 67.8% patients (n= 4391) received excess Rx
- 93% of these excess days were after discharge
- No difference in survival, readmissions, repeat ED visits - but **5% per day increase risk of AEs**

Target:
antibiotic upon
discharge from
hospital



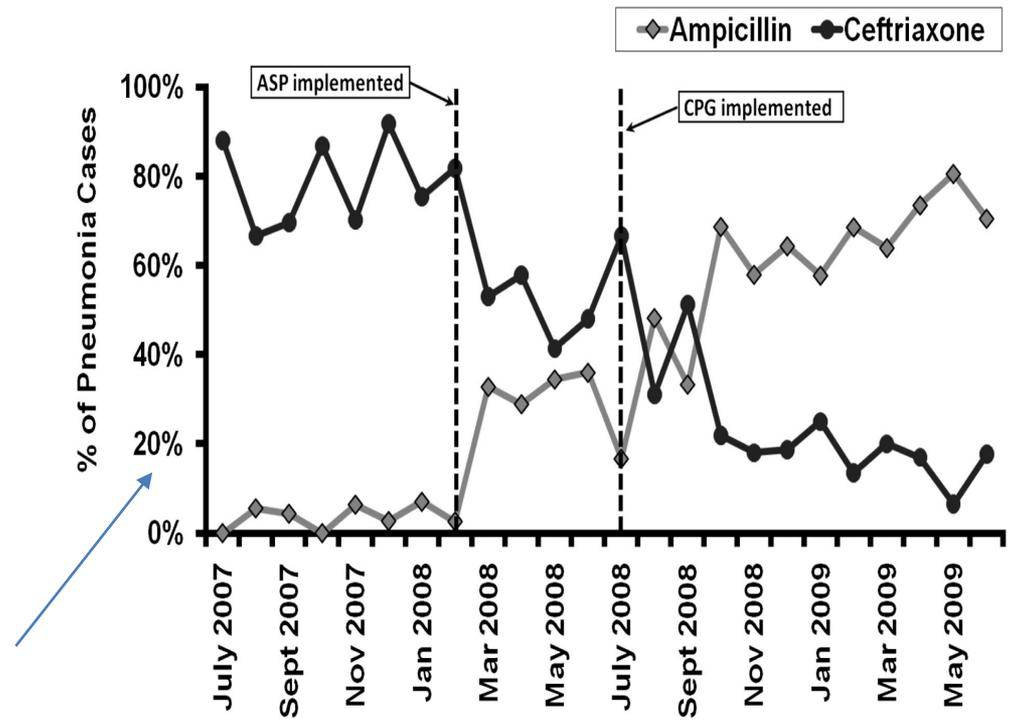
One approach



Example 1

- Before/after study including >1000 children with CAP
- Narrower spectrum antibiotic upon admission & discharge
- No change in treatment failure

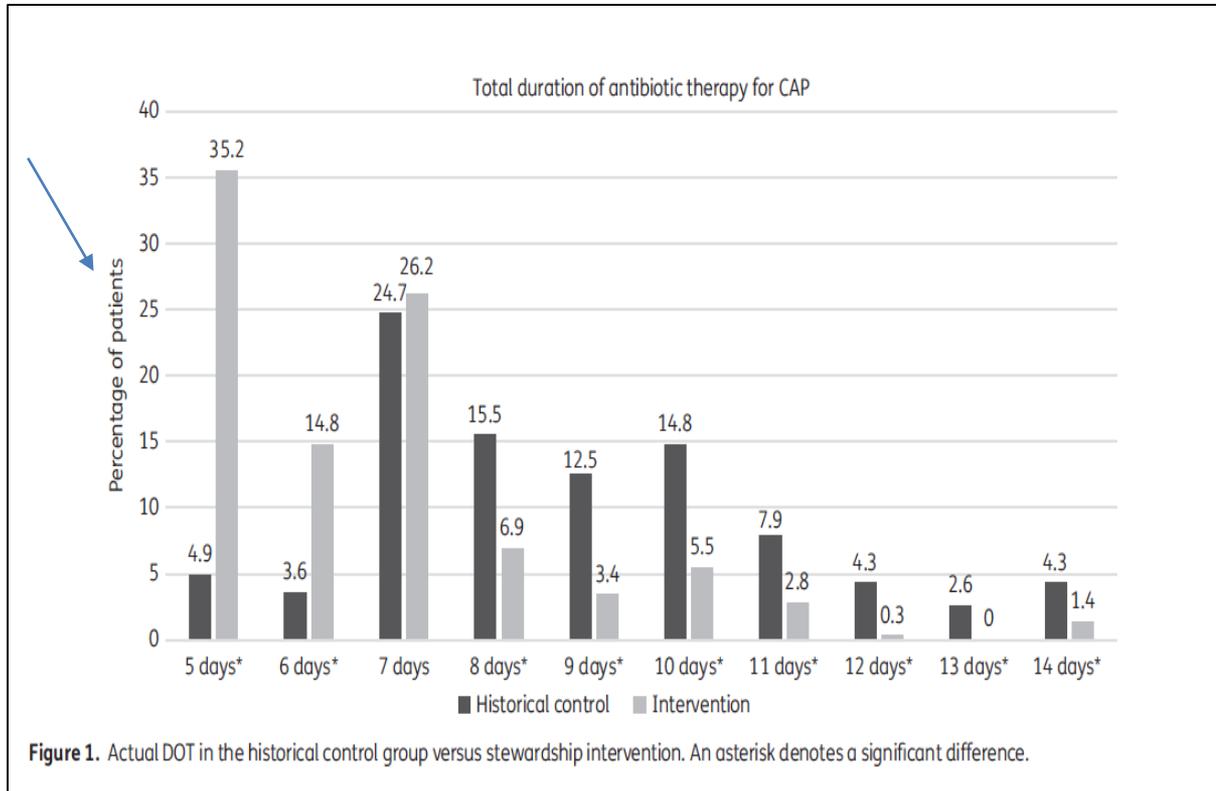
Metric:
Empiric
antibiotic
therapy



Metric:
Duration of
therapy

Example 2

- Multifaceted approach to improve abx for community-acquired pneumonia:
 - Guidelines
 - Education
 - Audit and feedback



Process metrics

Example 3

Dissemination strategy:

- weekly clinical meetings (real cases of hospitalized patients were used)
- the distribution of educational material (pocket booklets)
- posters

Table 1 - Characteristics of the patients hospitalized for community-acquired pneumonia between July of 2007 and October of 2008 at the Marília School of Medicine *Hospital das Clínicas de Marília*, located in the city of Marília, Brazil^a

Characteristic	Pre-implementation period	Post-implementation period	Total	p
Diagnosed patients, n	48	20	68	
Male	28 (58.3)	14 (70.0)	42 (61.8)	0.36
Female	20 (41.7)	6 (30.0)	26 (38.2)	
Age, in years ^b			68.5 ± 19.2	
Recording of SpO ₂ on the medical chart	9 (19)	6 (30)	15 (19)	0.42
CURB-65 score ≥ 3	27 (56.3)	12 (60.0)	39 (57.4)	0.77
Severe sepsis	19 (39.6)	13 (65.0)	32 (47.1)	0.056
Prescribed antimicrobial agents				
Ceftriaxone	26 (54.2)	17 (85.0)	43 (63.2)	0.01
Ciprofloxacin	9 (18.8)	0 (0.0)	9 (13.2)	0.009
Clindamycin	15 (31.3)	3 (15.0)	18 (26.5)	0.16
Azithromycin	4 (8.3)	15 (75.0)	19 (27.9)	0.0001
β-lactam + macrolide	3 (6.3)	15 (75.0)	18 (26.5)	0.0001
Hospital stay, in days ^b	17.8 ± 22	16.1 ± 13.4	17.2 ± 19.8	0.32
Death	17 (35.4)	3 (15.0)	20 (29.4)	0.09

CURB-65: (mortality prediction score) mental Confusion, Urea, Respiratory rate, Blood pressure, and age > 65 years. ^aValues expressed as n (%), except where otherwise indicated. ^bValues expressed as mean ± SD.

Clinical Practice Guideline for the Management of Asymptomatic Bacteriuria: 2019 Update by the Infectious Diseases Society of America^a

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- Recommend screening or treating for asymptomatic bacteriuria in **very few targeted** populations
 - Pregnant women
 - Prior to urologic surgery where mucosal injury is anticipated
- Recommend against screening (and treating) asymptomatic bacteriuria in almost all other groups – including elderly with altered mental status or after a fall unless there are **symptoms** of a urinary tract infection

Prevalence of ASB

Population	Prevalence, %
Children	
Boys	<1
Girls	1 – 2
Healthy women	
Premenopausal	1 – 5
Pregnant	2 – 10
Postmenopausal	3 – 9
Diabetes mellitus	
Women	11 – 16
Men	1 – 11
Elderly in community	
Women	11 – 16
Men	4 – 19

Population	Prevalence, %
Elderly in LTCF	
Women	25 – 50
Men	15 – 50
Spinal cord injury	
Intermittent catheter	23 – 69
Kidney transplant	
First month after	23 – 24
1 month – 1 year after	10 – 17
>1 year after	2 – 9
Indwelling catheter use	
Short-term	3 – 5%/day
Long-term	100

2019 Infectious Diseases Society of America Guideline Update

Screening for and treatment of ASB indicated for:

- Pregnant women
- Invasive urological procedures

Do NOT screen for or treat ASB:

- Elderly
- Diabetics
- Indwelling urinary catheters
- Spinal cord injury
- Long term care facility residents
- Elective, non-urologic surgeries including prosthetic joints*
- Neutropenia*
- Renal or other solid organ transplant*
- Children*

*** Not addressed in
2005 IDSA guideline**

2019 IDSA Guideline Recommendations for Specific Clinical Scenarios

“In older patients with functional and/or cognitive impairment with bacteriuria and delirium without local genitourinary symptoms or other systemic signs of infection, we recommend assessment for other causes and careful observation rather than antimicrobial treatment”

(strong recommendation, low-quality evidence)

Two key approaches to reduce antibiotic treatment of ASB

1) Prevent identification of ASB to begin with

- Eliminate inappropriate or unnecessary urinalyses and urine cultures

2) Prevent antibiotic treatment when ASB identified

- Increase recognition/diagnosis of ASB
- Understand antibiotics not indicated
- Appropriately withhold antibiotics

Summary

- ❑ Multifaceted intervention → diagnostic/ treatment algorithms, audit and feedback, education
- ❑ Targeted ordering of urine cultures *and* decision to treat positive cultures
- ❑ Reduced total volume of urine cultures and rate of antibiotic treatment of ASB
- ❑ More effective on long-term care wards than acute medical wards
- ❑ Ongoing study in non-catheterized patients

Potential areas of focus



Inappropriate treatment of asymptomatic bacteriuria is associated in negative outcomes.

- In one study of >2700 hospitalized patients, abx for ASB was associated with longer hospitalizations.

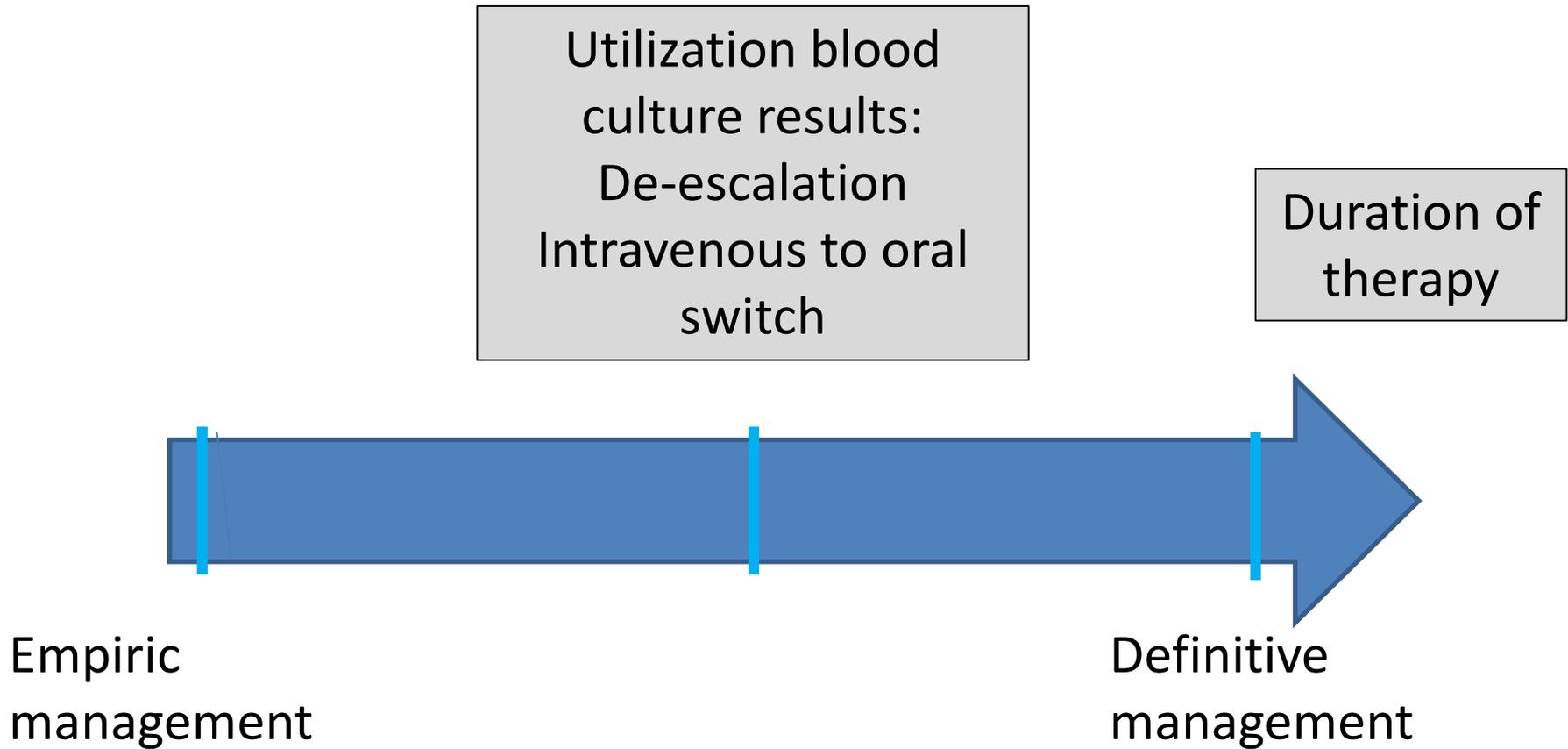
Potential areas of focus



Interventions:

- Education
- Policy
- Engage stakeholders
- Standardized orders
-

Bacteremia: De-escalation



Annex IV: Sample AMS review form

ANTIMICROBIAL STEWARDSHIP PROGRAMMES

IN HEALTH-CARE FACILITIES IN LOW- AND

MIDDLE-INCOME COUNTRIES

A WHO PRACTICAL TOOLKIT

Patient information		
Date:	Department:	Ward:
Patient name:	Age:	Sex: Male <input type="checkbox"/> or Female <input type="checkbox"/>

Antibiotic prescriptions				
Antibiotics prescribed	Dose	Route	Interval	Start date

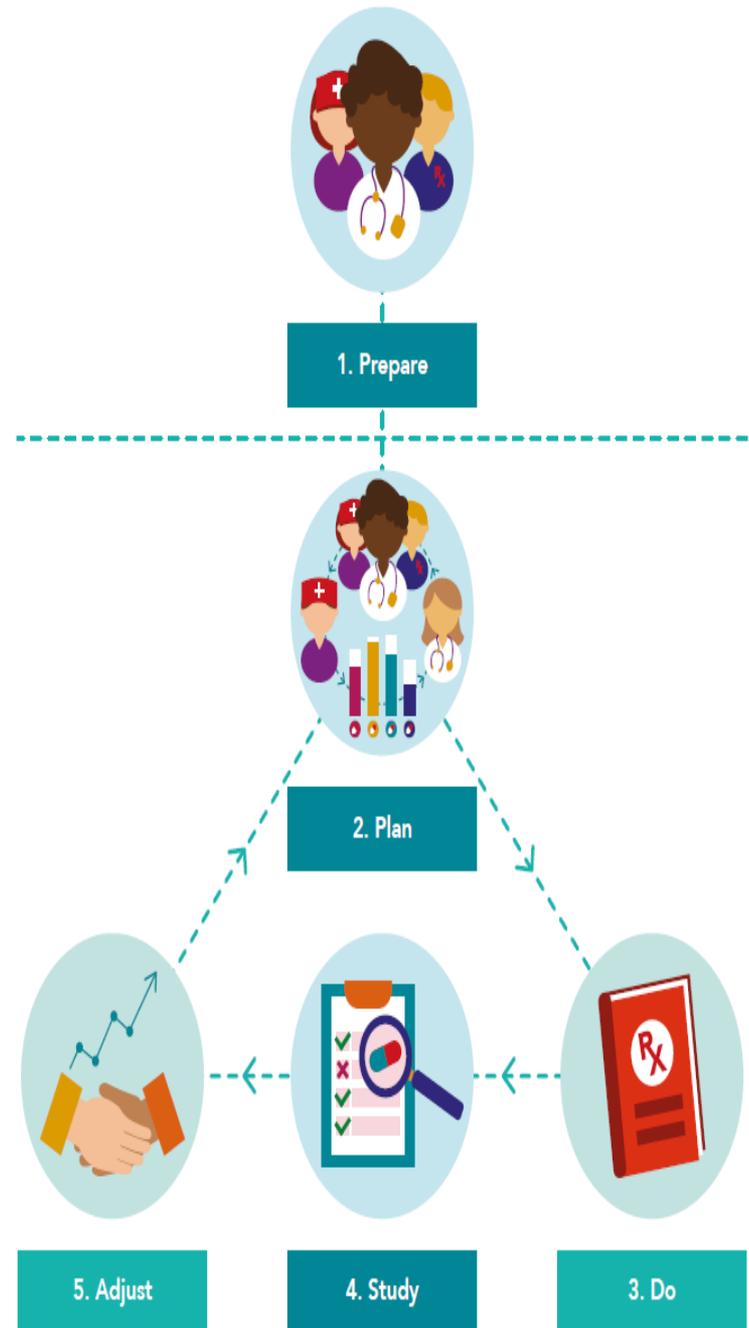
48-hour review of antibiotic treatment				
Is antibiotic treatment reviewed? Yes <input type="checkbox"/> No <input type="checkbox"/>		If yes, what action?		
Escalate <input type="checkbox"/>	Continue <input type="checkbox"/>	De-escalate <input type="checkbox"/>	Stop <input type="checkbox"/>	IV-oral switch <input type="checkbox"/>
Why is antibiotic treatment being continued?				
Continuing clinical signs of infection <input type="checkbox"/>		Confirmed infection <input type="checkbox"/>	Other (comment):	
Microbiology specimens collected? <input type="checkbox"/>	Microbiology results received? <input type="checkbox"/>	Microbiology results acted upon? <input type="checkbox"/>		
Date:	Date:	Comment:		

Targets for stewardship: common infections

1 - appropriate utilization of
urine cultures

2 - ensure appropriate
duration of therapy for
community acquired
pneumonia

3 - promote **re-evaluation of
blood culture results** and
modify antibiotic course



The Approach to the Problem Prescriber

- Carefully plan your approach:
 - Pick your battles
 - Timing is important
 - Avoid heat of the moment confrontations (generate light not heat)
- Do your homework
 - Gather as much data as possible
 - DUE: Service and physician specific for several drugs
 - Discuss with Clinical PharmDs and discretely other MDs
 - Discuss with CMO/Chief of staff
 - Understand the MD's Practice and Patient Population

Critical Success Factors Identified

- Collegial and educational relationship
- Daily review of antimicrobial orders by a consistent accountable team
- Support of hospital/medical leadership
- Development of criteria and guidelines for anti-infective use
- Formulary restriction
- Education of prescribers to insure compliance
 - **But the most important one is investing in the human factor**

The Guidelines

To be Finalized (it toke 8 months)

- The most common syndromes:
- 1. sepsis
- 2. UTI
- 3. Pneumonia (CAP / HAP)
- 4. soft tissue infection
- 5. diabetic foot
- 6. URTI
- 7. Antibiotic surgical prophylaxis
- 8. Endocarditis

8. What are the top three common infectious clinical syndromes at your facility that are either known or estimated?

11 responses

- pneumonia , UTI, Sepsis
- surgical infection, endocarditis, ventilator associated pneumonia
- respiratory tract infections, UTI, GIT
- UTI, pneumonia, diabetic foot infection
- UTI, ventilator associated pneumonia, SSI
- dry socket, pulpits, periodontitis

https://docs.google.com/forms/d/1BISnSiMCDpJwIHGhoctajupLu68D_ymORr6UZEWNK/viewanalytics

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ASSESSMENT OF CURRENT PRACTICES

6/21/18, 8:52 AM

- ESBL, MRSA
- UTI, URTI, Vaginitis
- pneumonia, UTI , wound infection
- UTI, URTI, Gastroenrritis
- pneumonia. UTI, wound infection

Infection	Most likely organisms	1st line empiric treatment	In penicillin allergy	Comments
Cellulitis (non-diabetic)	<i>S. aureus</i> <i>Streptococci</i>	<p>cloxacillin 1-2g q6h iv If erysipelas Or streptococcal infection suspected add Benzylpenicillin 1.2-2.4g q6h Iv Or Cefazolin 1 gm IV q 8 hr</p> <p>Oral switch: cloxacillin 1g qds po If erysipelas or streptococcal infection suspected ADD Amoxicillin 500mg q8h po</p>	<p>Clindamycin 300 mg PO/IV Q8H Or Doxycycline 100 mg po bid Or trimethoprim-sulfamethaxazole double strength tablets po q 12 hrs</p> <p>If severe cellulitis/ risk of MRSA consider adding (Vancomycin 15mg/kg q12h iv)</p>	Duration 7-14 days Consider oral switch following clinical improvement.
Line Infection	<i>S. aureus</i> , <i>Coagulase negative staphylococci and other organisms</i>	Vancomycin ± Cefepime 1-2 g IV Q8H (use higher dose if pseudomonas suspected)	<p>Take blood cultures prior to commencing antibiotics (Vancomycin 15mg/kg q12h iv) If gram negative organisms are suspected add Ciprofloxacin 400mg q12h iv / 500mg q12h po Or broad antibiotics if suspected multidrug resistant organisms</p>	
Necrotizing fasciitis	Multiple organisms including Group A <i>Streptococci</i>	<p>cloxacillin 2g q6h iv + Benzyl penicillin 2.4g q6h iv + Ciprofloxacin 400mg q12h iv + Metronidazole 500mg q8h iv</p>	<p>Clindamycin 600mg q6h iv + [Ciprofloxacin 400mg q12h iv ± Gentamicin] + Vancomycin 15mg/kg q12h iv</p>	Seek urgent surgical advice
Gas gangrene	<i>Clostridium perfringens</i> and other gas producing organisms	<p>Benzyl penicillin 2.4g q6h iv + Ciprofloxacin 400mg q12h iv + Metronidazole 500mg q8h iv</p>	<p>Clindamycin 600mg q6h iv + Ciprofloxacin 400mg q12h iv</p>	Seek urgent surgical advice For mixed infection (aerobic/anaerobic) use broad spectrum antibiotics (imipenem /meropenem/Tazocin)
Infected burns	Send swabs for cultures and sensitivities to direct therapy	<p>Apply silver sulphadiazine 1% cream to the affected areas.</p> <p>Co-amoxiclav 1.2g q8h iv / 625mg q8h po OR Tazocin/meropenem/doripenem/imipenem + vancomycin or Daptomycin</p>		
Surgical wound	Following clean surgery	Co-amoxiclav 1.2g q8h iv / 625mg q8h po	trimethoprim-sulfamethaxazole double strength one tablet po q 12 hrs	Based on culture and sensitivity results and location of surgical site:
Soft Tissue		Co-amoxiclav 1.2g q8h iv / 625mg q8h po		

The education

First part
submitted to
NHRA for CME
hours

- **First part** : Basic concepts in AMR antimicrobial resistance and general guidelines :
 - Part a for physicians : please see attached
 - Part b for nurses please see attached
- **The second** training type will be hands on the program Tools ; will be provided later
- Second part (hand on training)
- 1. MDRO (WHOnet/ GLASS)
- 2. Antibiotic consumption
- 3. HAI surveillance
- **The third part:**
 - Method of implementation and reporting
 - Competencies needed
 - Policies
 - Accreditations

Antibiotic Stewardship Program

The screenshot displays a Microsoft Excel spreadsheet titled "preferenceno_synoptologika_arginino_ABC_Cal_3102 - User Product Activation Failed". The spreadsheet is organized into two main sections. The top section lists various antibiotics with columns for Name of product, Grams per test dose, No. doses per package, Name of antibacterial, ATC code route, DDD (WHO 2008) U, DDD (WHO 2008) N, and No. packages. The bottom section is a detailed list of antibiotic combinations and their DDD values.

Name of product	Grams per test dose	No. doses per package	Name of antibacterial	ATC code route	DDD (WHO 2008) U	DDD (WHO 2008) N	No. packages
Cloxacillin 500 mg	0.5	1	Cloxacillin (Oral)	J01CF02 O	2 g	0.0	0.0
gr. amoxicilin + p. benzoylen			Cloxacillin (Parenteral)	J01CF03 P	2 g	0.3	0.0
			Methicillin	J01CF03 P	4 g	0.0	0.0
			Oxacillin (Oral)	J01CF04 O	2 g	0.0	0.0
			Oxacillin (Parenteral)	J01CF04 P	2 g	0.0	0.0
			Flucloracillin (Oral)	J01CF05 O	2 g	0.0	0.0
			Flucloracillin (Parenteral)	J01CF05 P	2 g	0.0	0.0
			Subactam	J01DD01 P	1 g	0.0	0.0
			Taximobactam	J01DD02			
AMPCILLIN-SULBACTAM 1.5 g	1	1	lin and enzyme inhibitor (Parenteral)	J01CR01 P	2 g	0.5	0.0
AMPCILLIN-SULBACTAM 1.5 g	1	1	lin and enzyme inhibitor (Oral)	J01CR01 O	2 g	1.0	0.0
			lin and enzyme inhibitor (Oral)	J01CR02 O	1 g	0.0	0.0
			lin and enzyme inhibitor (Oral)	J01CR03 O	1 g	0.0	0.0
			lin and enzyme inhibitor (Oral)	J01CR04 O	1 g	0.0	0.0
			Amoxicillin and enzyme inhibitor (Oral)	J01CR05 O	1 g	0.0	0.0
AMOXICILIN-CLAVULANIC ACID 1.2g	1	1	Amoxicillin and enzyme inhibitor (Parenteral)	J01CR02 P	3 g	0.3	0.0
			Amoxicillin and enzyme inhibitor (Oral)	J01CR02 O	1.5 g	0.0	0.0
			Amoxicillin and enzyme inh. (Oral)	J01CR04 O	1.5 g	0.0	0.0
			lin and enzyme inhibitor	J01CR01 P	15 g	0.2	0.0
			lin and enzyme inhibitor	J01CR03 P	15 g	0.3	0.0
			lin and enzyme inhibitor	J01CR05 P	14 g	0.1	0.0
			lin and enzyme inhibitor	J01CR05 P	14 g	0.3	0.0
			Amoxicillin + cloxacillin (250/250) (Oral)	J01CR02 O	2 g	0.0	0.0
			Amoxicillin + cloxacillin (250/250) (Parenteral)	J01CR02 P	2 g	0.0	0.0
			Amoxicillin + Nidococillin (250/250) (Oral)	J01CR03 O	2 g	0.0	0.0
			Amoxicillin + Nidococillin (250/250) (Parenteral)	J01CR03 P	2 g	0.0	0.0
			Amoxicillin + oxacillin (125/125) (Oral)	J01CR04 O	2 g	0.0	0.0

ATC Code	DDD (WHO 2008) U	DDD (WHO 2008) N	No. packages
J01BA01	0.0	0.0	0.000
J01BA02	0.0	0.0	0.000
J01BA03	0.0	0.0	0.000
J01BA04	0.0	0.0	0.000
J01BA05	0.0	0.0	0.000
J01BA06	0.0	0.0	0.000
J01BA07	0.0	0.0	0.000
J01BA08	0.0	0.0	0.000
J01BA09	0.0	0.0	0.000
J01BA10	0.0	0.0	0.000
J01BA11	0.0	0.0	0.000
J01BA12	0.0	0.0	0.000
J01BA13	0.0	0.0	0.000
J01BA14	0.0	0.0	0.000
J01BA15	0.0	0.0	0.000
J01BA16	0.0	0.0	0.000
J01BA17	0.0	0.0	0.000
J01BA18	0.0	0.0	0.000
J01BA19	0.0	0.0	0.000
J01BA20	0.0	0.0	0.000
J01BA21	0.0	0.0	0.000
J01BA22	0.0	0.0	0.000
J01BA23	0.0	0.0	0.000
J01BA24	0.0	0.0	0.000
J01BA25	0.0	0.0	0.000
J01BA26	0.0	0.0	0.000
J01BA27	0.0	0.0	0.000
J01BA28	0.0	0.0	0.000
J01BA29	0.0	0.0	0.000
J01BA30	0.0	0.0	0.000
J01BA31	0.0	0.0	0.000
J01BA32	0.0	0.0	0.000
J01BA33	0.0	0.0	0.000
J01BA34	0.0	0.0	0.000
J01BA35	0.0	0.0	0.000
J01BA36	0.0	0.0	0.000
J01BA37	0.0	0.0	0.000
J01BA38	0.0	0.0	0.000
J01BA39	0.0	0.0	0.000
J01BA40	0.0	0.0	0.000
J01BA41	0.0	0.0	0.000
J01BA42	0.0	0.0	0.000
J01BA43	0.0	0.0	0.000
J01BA44	0.0	0.0	0.000
J01BA45	0.0	0.0	0.000
J01BA46	0.0	0.0	0.000
J01BA47	0.0	0.0	0.000
J01BA48	0.0	0.0	0.000
J01BA49	0.0	0.0	0.000
J01BA50	0.0	0.0	0.000
J01BA51	0.0	0.0	0.000
J01BA52	0.0	0.0	0.000
J01BA53	0.0	0.0	0.000
J01BA54	0.0	0.0	0.000
J01BA55	0.0	0.0	0.000
J01BA56	0.0	0.0	0.000
J01BA57	0.0	0.0	0.000
J01BA58	0.0	0.0	0.000
J01BA59	0.0	0.0	0.000
J01BA60	0.0	0.0	0.000
J01BA61	0.0	0.0	0.000
J01BA62	0.0	0.0	0.000
J01BA63	0.0	0.0	0.000
J01BA64	0.0	0.0	0.000
J01BA65	0.0	0.0	0.000
J01BA66	0.0	0.0	0.000
J01BA67	0.0	0.0	0.000
J01BA68	0.0	0.0	0.000
J01BA69	0.0	0.0	0.000
J01BA70	0.0	0.0	0.000
J01BA71	0.0	0.0	0.000
J01BA72	0.0	0.0	0.000
J01BA73	0.0	0.0	0.000
J01BA74	0.0	0.0	0.000
J01BA75	0.0	0.0	0.000
J01BA76	0.0	0.0	0.000
J01BA77	0.0	0.0	0.000
J01BA78	0.0	0.0	0.000
J01BA79	0.0	0.0	0.000
J01BA80	0.0	0.0	0.000
J01BA81	0.0	0.0	0.000
J01BA82	0.0	0.0	0.000
J01BA83	0.0	0.0	0.000
J01BA84	0.0	0.0	0.000
J01BA85	0.0	0.0	0.000
J01BA86	0.0	0.0	0.000
J01BA87	0.0	0.0	0.000
J01BA88	0.0	0.0	0.000
J01BA89	0.0	0.0	0.000
J01BA90	0.0	0.0	0.000
J01BA91	0.0	0.0	0.000
J01BA92	0.0	0.0	0.000
J01BA93	0.0	0.0	0.000
J01BA94	0.0	0.0	0.000
J01BA95	0.0	0.0	0.000
J01BA96	0.0	0.0	0.000
J01BA97	0.0	0.0	0.000
J01BA98	0.0	0.0	0.000
J01BA99	0.0	0.0	0.000
J01BA00	0.0	0.0	0.000

Quality indicators for the strategic plan (for 5 years)

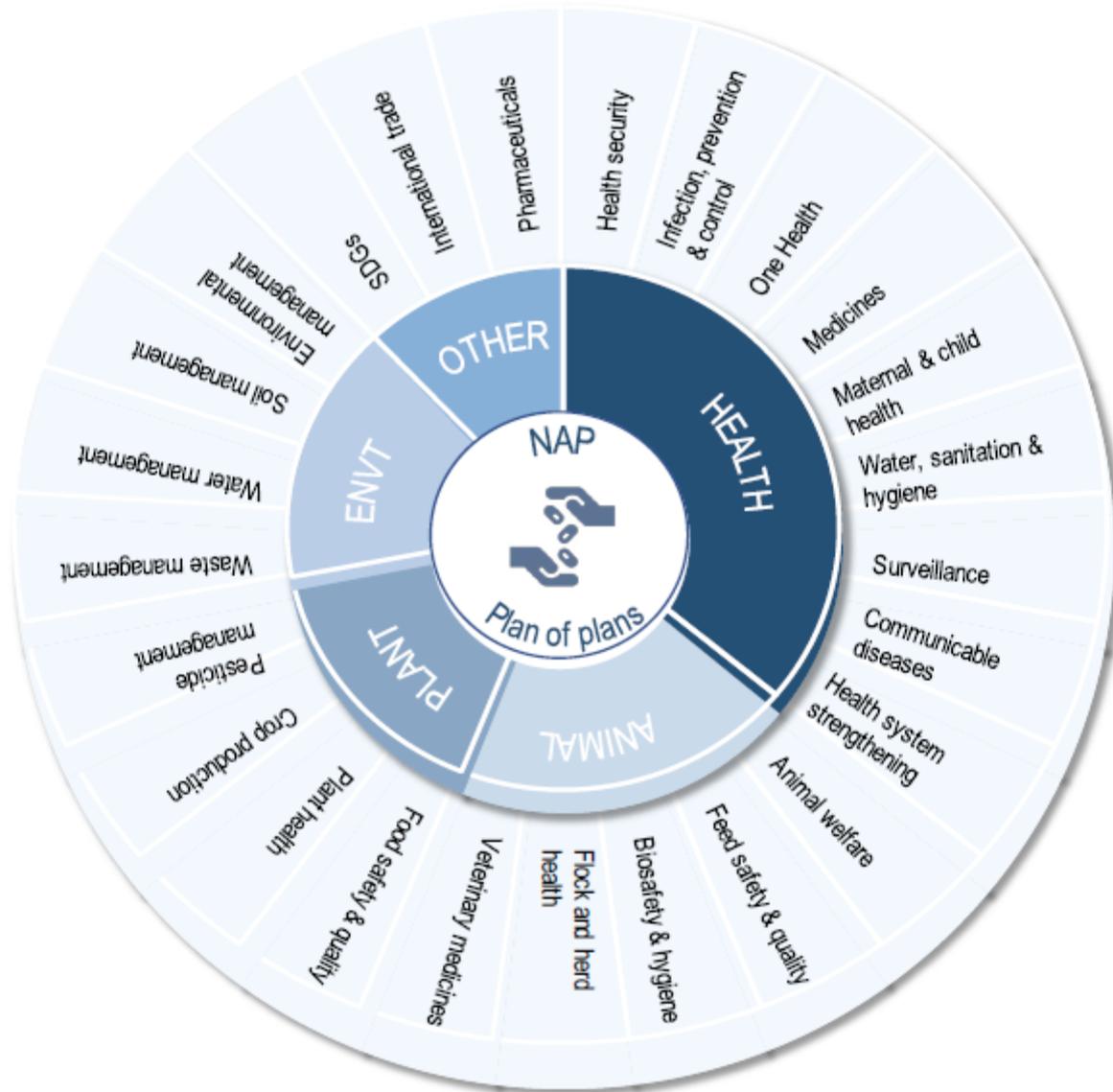
- **Outcome measures**

- Rate of c diff
- mortality rate due to infections
- Length of stay in all hospitals
- The rate and types of antibiotic use per the suggested syndromes
- Compliance with surgical antibiotic prophylaxis
- Number of trained staff /sessions./ workshops
- Number of isolate suggested and their trend and patterns of resistance
- Calculating ddd or dot
- Cost in animals for antibiotic
- Cost in humans
- Rate for compliance with guidelines

- **Process measures**

- Duration of treatment
- Iv to po

Figure 7: With links to many different national plans and strategies, NAPs are often a 'plan of plans'



Lessons I've Learned...

- Take your time and do your homework
- Culture matters and dictates everything else
- Obtain dedicated FTE from Hospital Administration
(Consider home in Quality)
- Reporting structure matters ...
 - Pharmacists and MDs can be pulled away if managers don't buy-in to stewardship
- Obtain leadership skills (local and national workshops)
- Stewardship burnout is real...Pursue scholarly work and outside networking

ELEMENTS OF AN EFFECTIVE ANTIMICROBIAL STEWARDSHIP PROGRAM

Team success

“The ultimate difference between a company and its competition is, in fact, the ability to execute.”

- Larry Bossidy

One size does not fit all



Thank you

Put your leadership and communication skills
together to sell your results to the people in suits

What I've learned through the years