

First ABS studies in my hospital: How to do it, what to learn from it ?

Kornelia Chrapkova

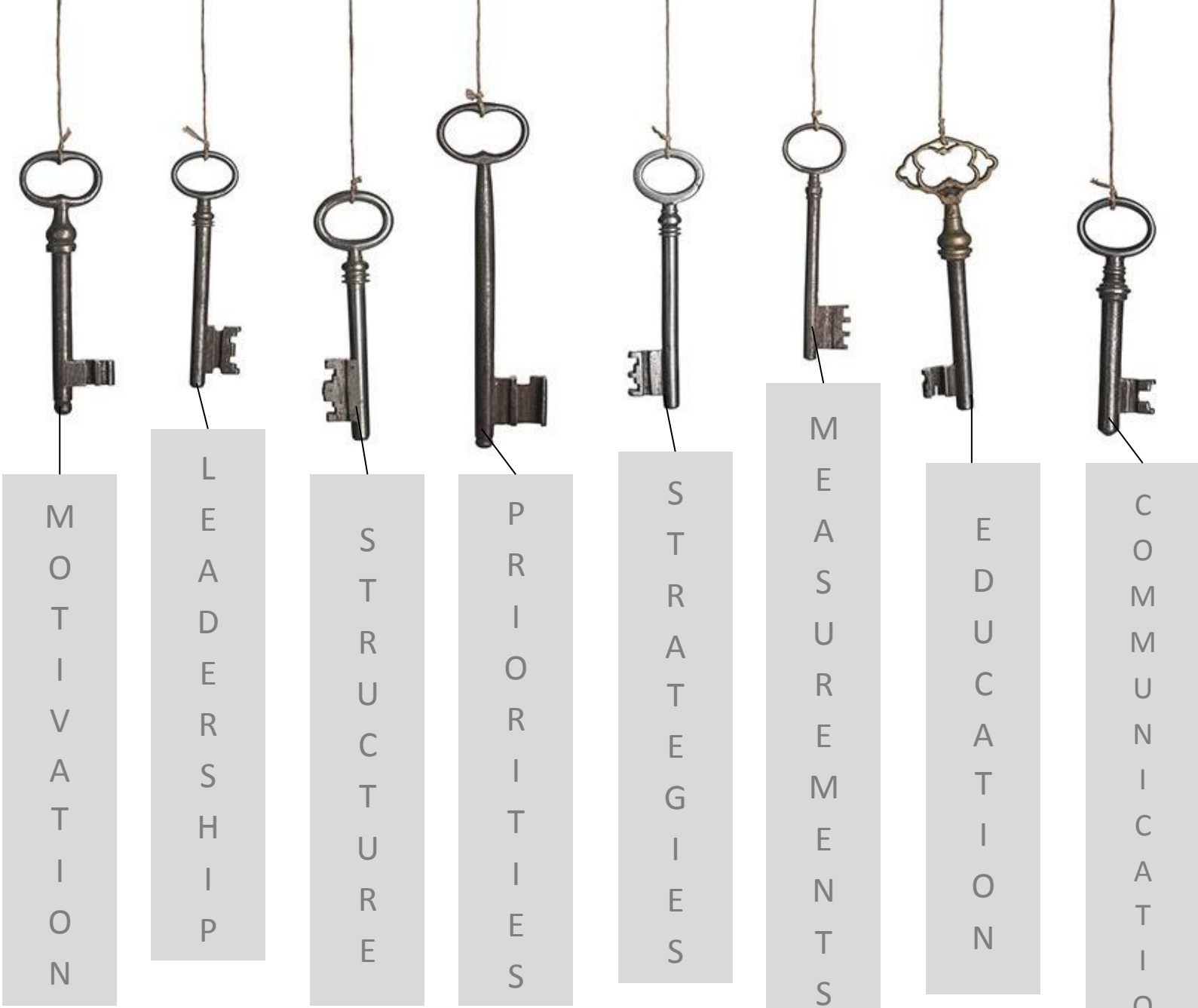
Institute for Cilinical and Experimental Medicine
Prague, Czech Republic

- I have nothing to declare

“The right antibiotic for the right patient, at the right time, with the right dose, the right route and the right duration causing the least harm to the patient and future patients“

www.cdc.gov/getsmart/healthcare/inpatient-stewardship





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Priorities and measuring progress and success



Centers for Disease Control and Prevention
 CDC 24/7: Saving Lives, Protecting People™

Antibiotic Stewardship Driver Diagram



INSTITUTE FOR
 HEALTHCARE
 IMPROVEMENT

Timely and appropriate antibiotic utilization in the acute care setting

Decreased incidence of antibiotic-related adverse drug events (ADEs)

Decreased prevalence of antibiotic resistant healthcare-associated pathogens

Decreased incidence of healthcare-associated *C. difficile* infection

Decreased pharmacy cost for antibiotics

Primary Drivers

Timely and appropriate initiation of antibiotics

Appropriate administration and de-escalation

Data monitoring, transparency, and stewardship infrastructure

Availability of expertise at the point of care

Secondary Drivers

- Promptly identify patients who require antibiotics
- Obtain cultures prior to starting antibiotics
- Do not give antibiotics with overlapping activity or combinations not supported by evidence or guidelines
- Determine and verify antibiotic allergies and tailor therapy accordingly
- Consider local antibiotic susceptibility patterns in selecting therapy
- Start treatment promptly
- Specify expected duration of therapy based on evidence and national and hospital guidelines

- Make antibiotics patient is receiving and start dates visible at point of care
- Give antibiotics at the right dose and interval
- Stop or de-escalate therapy promptly based on the culture and sensitivity results
- Reconcile and adjust antibiotics at all transitions and changes in patient's condition
- Monitor for toxicity reliably and adjust agent and dose promptly

- Monitor, feedback, and make visible data regarding antibiotic utilization, antibiotic resistance, ADEs, *C. difficile*, cost, and adherence to the organization's recommended culturing and prescribing practices

- Develop and make available expertise in antibiotic use
- Ensure expertise is available at the point of care

Leadership and Culture



Identification of effective interventions for your setting



Identification of effective interventions for your setting

- LOCAL NEEDS
 - What can be implemented depends on local needs and issues, geography, available skills and expertise, other resources
- SUPPORT
 - It's important to select the interventions that are more supported by clinical staff
- STEP BY STEP
 - It's not recommended that any facility attempt to implement all of the interventions at once

Identification of effective interventions for your setting

- FRONT-END strategies – preauthorization and restriction
 - Antimicrobial Prescribing Policy
 - Clinical guidelines or Care Pathways
 - Formulary restrictions/approval system – expert



Identification of effective interventions for your setting

- BACK-END strategies
 - Antimicrobial review methods
 - Review of indication for antibiotic and compliance with policy
 - Review of appropriateness of antibiotic choice, dose, route and planned duration,
 - Review of drug allergy
 - Potential for conversion from IV to oral
 - Requirement for TDM
 - Prospective audit and feedback



Identification of key measurements for improvement

- Structural indicators
- Process measures
- Outcome measures
- Balancing measures



The key is also:

Establishing what to measure, the frequency of measurement and how data will be communicated and acted upon

Starting with ABS in my hospital: Institute for Clinical and Experimental Medicine



Institute for Clinical and Experimental Medicine

- 315 beds(111 intensive care unit beds)
- 4 specialized centres:
 - Cardiology centre
 - Transplant centre :
 - Nephrology, Hepatogastroenterology, Transplant Surgery,
 - Diabetology centre
 - Centre for experimental medicine
- 5 clinical pharmacists
 - 1 fulltime, 4 part time, covering 5 wards

Antimicrobial Stewardship

OUR STARTING POINT

- Preauthorization and restriction
 - Electronically – microbiologist to authorise
 - Microbiologist – consulting during working hours
 - Formulary of restricted antibiotics

Antimicrobial Stewardship

Starting with Training – making experts

- 2011-2012: mandatory training in Antimicrobial Stewardship organised by Ministry of Health
 - consultant of ICU, microbiologist, clinicians, chief pharmacist, clinical pharmacist
- Weakness:
 - lack of motivation
 - resistance to change
 - lack of trust, lack of acknowledgement - clinical pharmacist –new role in the hospital

**CAN WE START
THE**

ANTIMICROBIAL

STEWARDSHIP

**OVER
AGAIN?**

**I WASN'T
READY...**



Antimicrobial Stewardship Leadership, Structure and Organization

- Microbiologist, consultant of ICU, clinical pharmacist, 1 clinician per every ward
- **Team for Prevention and Control of infection**
- **Drug Committee Support**





Identification of the need

Definition of priorities

Key measurements

Antimicrobial Stewardship Prescribing Quality Assessment

Assessment of an adherence to a local
and international guideline for
SURGICAL ANTIMICROBIAL
PROPHYLAXIS



Assessment of an adherence to a local and international guideline for SURGICAL ANTIMICROBIAL PROPHYLAXIS (AP)

Surgical Site Infection Prevention Policies and Adherence in California Hospitals, 2010

Laurie J. Conway, RN, MS, CIC¹, Monika Pogorzelska, PhD, MPH¹, Elaine L. Larson, PhD, RN, FAAN, CIC^{1,2}, and Patricia W. Stone, PhD, RN, FAAN¹

¹Columbia University School of Nursing, New York, New York

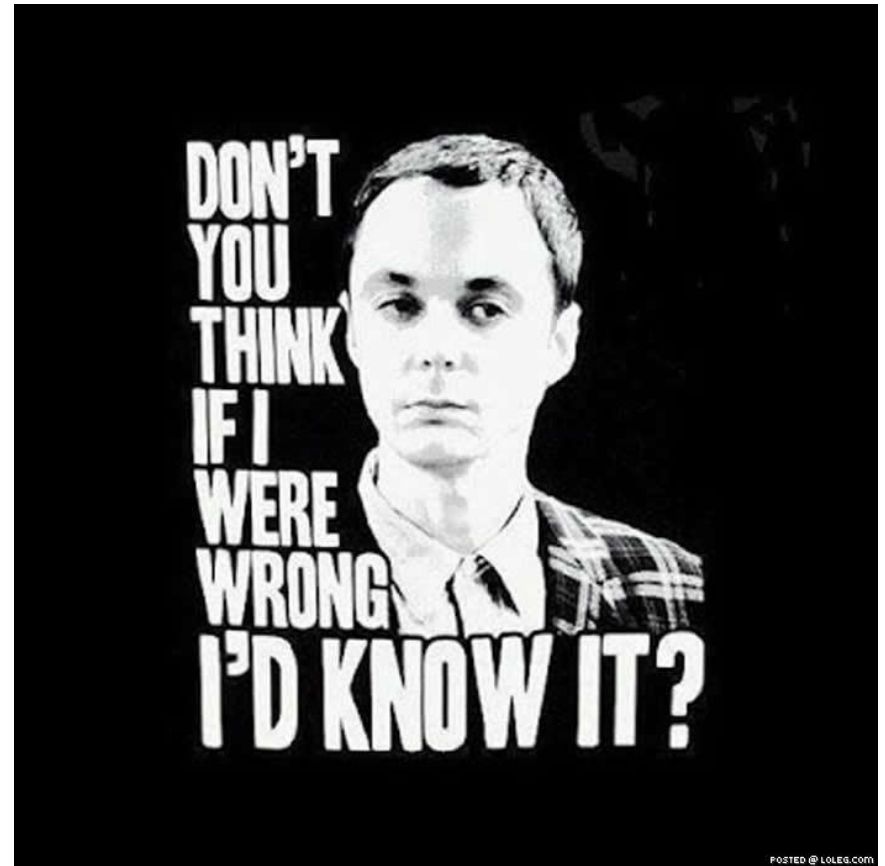
²Columbia University Mailman School of Public Health, New York, New York

Surgical site infections (SSIs) are common, costly, and preventable; 55% may be prevented with current evidence-based strategies.¹ SSIs occur at a rate of more than 290,000 infections per year and cost approximately \$25,500 per infection, and US hospitals could therefore save more than \$4 billion and prevent thousands of deaths annually by implementing SSI prevention strategies.² We sought to describe the presence of and adherence to SSI prevention policies in California hospitals. Specifically, we examined the adoption of



Why making an AUDIT?

- Key approach in changing prescriber's behaviour
 - Individual provider's performance is monitored
 - And confidentially compared/assessed
 - The feedback is likely to be accepted without controversy when is confidential
- Passive educational efforts are good to inform not to change



Audit at Transplant Surgery

- Prospective audit Dec 2014/Jan 2015
- Assessment of 50 abdominal and vascular surgeries
- Clinical pharmacist, microbiologist and clinicians
- Adherence to a local and international recommendation (BRATZLER, D.W., et al. *Clinical practice guidelines for antimicrobial prophylaxis in surgery*. Am J Health-Syst Pharm. 2013)assessed
- 8 process measures - quality assessment
- Outcome measures weren't monitored because of the complex aetiology of SSI (many contributing factors)

Process measures		Adherence to IG	Adherence to LG
Indication	AP	100 %	Not able to assess
	Therapy	69%	Not able to assess
Adequate choice of antibiotic		68%	Not able to assess
Correct dose of antibiotic		24%	Not able to assess
Preoperative timing of antibiotic		30 %	Not able to assess
Per operative (repeated) administration of antibiotic in longer surgical procedures		0%	Not able to assess
Per operative (repeated) administration of antibiotic in surgeries with blood loss $\geq 1.5l$		0%	Not able to assess
Duration of AP		100 %	Not able to assess

There was NO surgical procedure which would be 100% adherent to all process indicators

Potential barriers for adherence



- Disagreement of local and international guidelines
- Non-availability of local guidelines for every clinician
- Vague local guidelines
- Lack of education
- Lack of multidisciplinary approach (authors were anaesthesiologists only)
- Interdisciplinary disagreement (responsibility for AP, surgeon vs. anaesthesiologist), logistic problems

What Can I do with the situation?

How can I improve quality of AP prescribing?

Appropriate surgical antibiotic prophylaxis:

- ↓ SSI,
- ↓ consumption of antibiotics,
- ↓ prevalence of antibiotic resistant health-care associated pathogens

Primary drivers

- Appropriate indication,
- Duration ,
- Timing,
- Monitoring, Transparency,
- Local available guidelines,
- Expertise,
- Education,

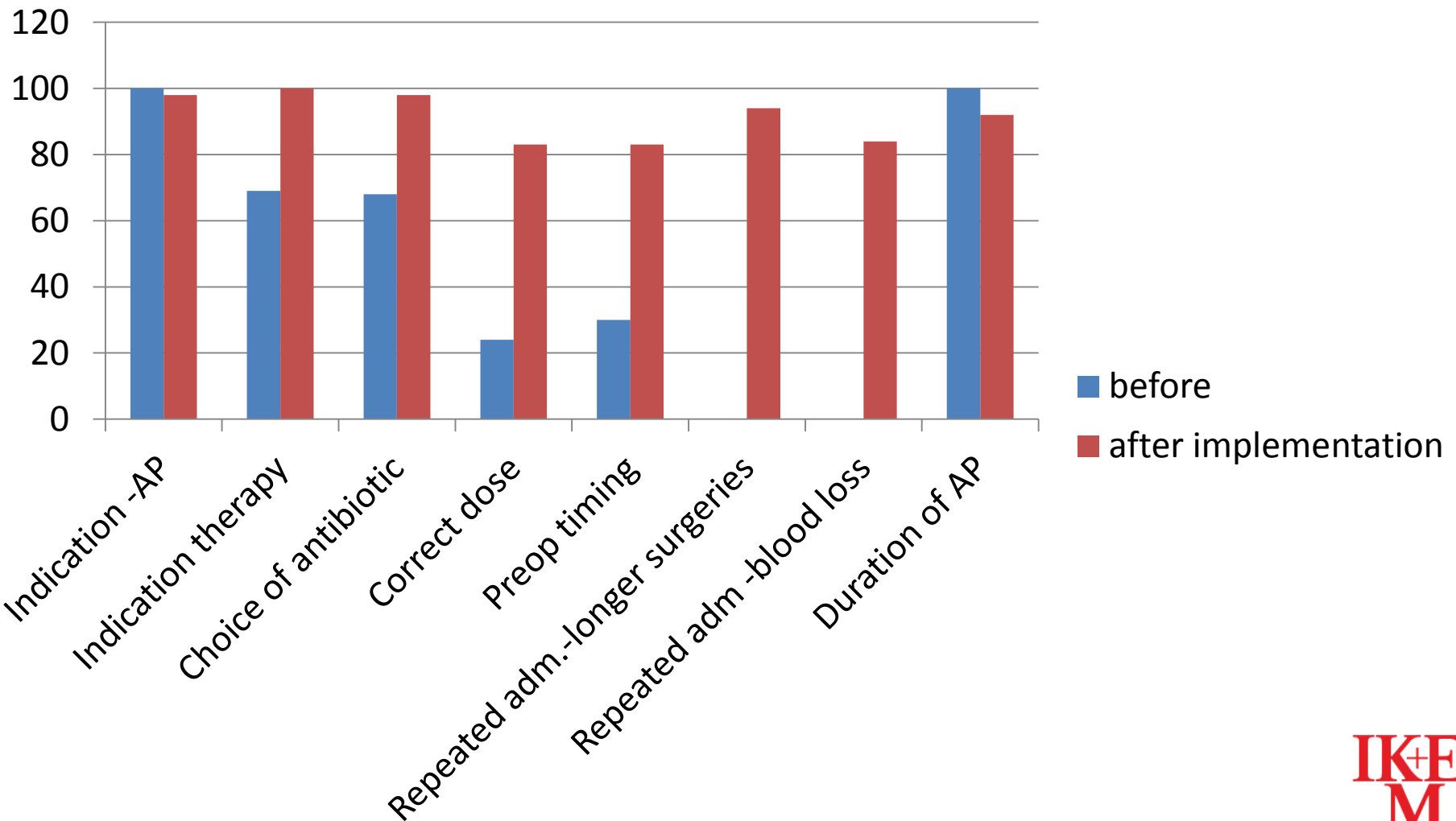
Secondary drivers

- Identification of patients/surgeries who require AP,
- Choice of correct antibiotic,
- Correct dose (children, obese, underweight, renal/liver failure)
- Options in allergy,
- Correct time of administration,
- Duration- when to extend prophylaxis over 24 hrs, start treatment,
- Monitor feedback and make visible data regarding antibiotic utilization,
- Develop expertise (microbiologist, pharmacist, clinicians),
- Communicate,
- Ensure expertise is available at the point of care,
- Use guidelines up to date, review every 2 yrs,
- Educate

Results of the audit 2 years after the implementation

Process measures		Adherence to LG
Indication	AP	98 %
	Therapy	100 %
Adequate choice of antibiotic		98 %
Correct dose of antibiotic		83 %
Preoperative timing of antibiotic		83 %
Per operative (repeated) administration of antibiotic in longer surgical procedures		94 %
Per operative (repeated) administration of antibiotic in surgeries with blood loss $\geq 1.5l$		84 %
Duration of AP		92 %

Assessment of adherence to guidelines Before and 2 yrs after implementation



Antimicrobial Stewardship Prescribing Quality Assessment **LOCAL GUIDELINES**



Local Guidelines- a starting point

- No willingness to share/display local guidelines –arguments:
 - Following “own” guidelines
 - Prescribing according to “own and best” clinical practice
 - “Guidelines are very restrictive”



A ROLE for the AUTHORITY to change the situation

Creating local guidelines

- Development of a local guideline by a local stakeholder group provides “sense of ownership”
- Identifying a need of necessary clinical guidelines
 - GAP analysis according to your local situation - urgency
 - Observation - clinical pharmacy practice, interventions, communication with clinicians, suggestions

The best indicator is when clinicians demand a guidance.

List of local guidelines

- Respecting national and international evidence based guidelines
- Stakeholders – clinicians, microbiologist, pharmacists
- Review and update every 2 years
 - **Clostridium difficile infection**
 - Observation, gaps in prescribing, infection control, lack of knowledge, lack of guidelines
 - **Splenectomy (vaccination, use of emergency antibiotics)**
 - Observation, no vaccination before/after splenectomy, no follow up, recommendation, lack of guidelines
 - **Treatment of infective endocarditis**
 - Observation - Lack of guidelines,
 - **UTI treatment outpatients/ inpatients**
 - Clinicians requirement
 - **COLOMYCIN- new old antibiotic**
 - Creating guidelines about correct dosing (until availability of new EMA recommendation) – risk of underdosing → failure of therapy

Antimicrobial Stewardship Prescribing Quality Assessment

Therapeutic drug monitoring

TDM



*"I stopped taking the medicine because I prefer
the original disease to the side effects."*

TDM -starting point



- Coordinated by clinicians only:
 - No previous training
 - Adjusting doses - “traffic light” system
 - Lack of knowledge about PK/PD characteristic, principle of TDM
 - Measuring concentrations as “needed”:
 - wrong use of laboratory resources



overdosing, failure of therapy, adverse events

What can I do with the situation ?

How can I change the old practice?

Appropriate antibiotic
treatment with NTI
antibiotic

↑ safety,
↑ therapeutic effect,
↓ cost,
↓ hospital stay,
↓ antibiotic resistance

Primary drivers

Indication,
Dosing,
Monitoring,
Local guidelines,
Expertise,
Education

Secondary drivers

Choice of a correct antibiotic,
Correct dose (renal failure, MIC,
disease)

Pharmacokinetic modeling,
Improve the use of laboratory
resources,
Interpret laboratory data,
concentrations,
Duration of the treatment
Options in allergy,
Collaboration of microbiologist,
clinicians, pharmacists,
Communicate,
Create local guidelines,
Create monitoring tool ,
Monitor feedback and make
visible recommendation,
Educate

TDM- today' s practice

- Automatic pharmacist consultation on wards with established clinical pharmacy service
- On request- where clinical pharmacy service is not provided routinely
- Every clinical pharmacist on ward is responsible for:
 - Identification of drugs that need TDM
 - Checking the indication, duration of the antibiotics
 - Guiding about measuring and timing of measurements of concentrations
 - Communication with microbiologist, biochemist, clinician
 - Interpreting the measured concentrations and predicting dosing (using MWPHARM)
 - Giving feedback to the prescriber (electronically and orally)
 - Education of the clinical staff (nurses , clinicians)

LABORATORNÍ PŘÍRUČKA

PRACOVNÍŠTĚ LABORATORNÍCH METOD

přednosta prof. MUDr. Antonín Jabor, CSc.



[Všeobecné informace](#)



[Laboratorní vyšetření PLM](#)



[Kontakty](#)



[Referenční meze](#)



[Žádanky](#)



[Terapeutické monitorování léčiv](#)



[Externí laboratoře](#)



[Laboratoře IKEM mimo PLM](#)

Verze 17, 15.08.2017



[Žádanka TDM antibiotika](#)

[Základní informace k TDM](#)

[Ukázka interpretace](#)



Terapeutické monitorování léků (TDM)

Kontakty:

Pracoviště laboratorních metod:

MUDr. Janka Franeková Ph.D., kl. 5225

Oddělení klinické farmacie a lékové informační centrum

Mgr. K. Chrapková PG Dip., kl. 5274, (8243)

PharmDr. Iva Prokopová Ph.D., kl. 5274 (8244)

Mgr. Eliška Dvořáčková kl. 5274 (8244)

Antibiotika teoretické podklady

[Aminoglykosidy](#)

[Vankomycin](#)

Imunosupresiva teoretické podklady

[Tacrolimus](#)

[Everolimus](#)

[Cyklosporin](#)

IKEM

Selection of patients with NTI antibiotics in hospital

CHRAPKOVÁ Kornélia (koch) x

ZLATOKOP **Klinická farmacie IKEM - monitoring**

Obecná konzilia
Klinická farmacie IKEM - konzilia
Klinická farmacie IKEM - monitoring
Farmakoterapeutická doporučení

Pacienti Dokumenty Laboratoře Mikrobiologie Vyšetření Formuláře Klinik

Vyhledávání Ambulance Konzilia Hospitalizace Operace TX program Extern

Dnes Sestavy

řehled Monitoring ... evidence

ing

KTCH 13.9.2017 - dd.mm.rrrr TDM amikacin gentamicin vankomycin

Doporučení

13.9.2017 Dnešní (13.9.) naměřená koncentrace vankomycinu (po podání první 12.9.dávky) je 4.6 umol/l. Doporučuji podávat 750 mg á 24 hod. Cílové rozmezí (6.9-10 umol/l). Prosím o kontrolní odběr koncentrace VAN před podáním ATB s ranními odběry v pátek (15.9.)
dávka odpovídající DP | stejný | DVOŘÁČKOVÁ Eliška (dvoe)

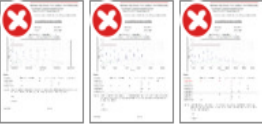
15.9.2017 Dnešní (15.9.) naměřená koncentrace vankomycinu je 9.8 umol/l. Koncentrace je v terapeutickém optimu (6.9-10 umol/l). Doporučuji podávat 750 mg á 24 hod v 18 hod, 60 min infuzí. Cílové rozmezí (6.9-10 umol/l). Prosím o kontrolní odběr koncentrace VAN před podáním ATB s ranními odběry v pondělí (18.9.).
dávka odpovídající DP | stejný | DVOŘÁČKOVÁ Eliška (dvoe)

18.9.2017 Dnešní (18.9.) naměřená koncentrace vankomycinu je 12.6 umol/l. Koncentrace je mírně nad optimem (6.9-10 umol/l). Doporučuji úpravu dávky na 250 mg á 24 hod v 8 hod, 30 min infuzí. Cílové rozmezí (6.9-10 umol/l). Prosím o kontrolní odběr koncentrace VAN před podáním ATB s ranními odběry ve středu (20.9.).
nižší dávka než v DP | stejný | DVOŘÁČKOVÁ Eliška (dvoe)

interní poznámky o sledování

Ukončení ? důvod ukončení

Příložené soubory


13.9.2017 15.9.2017 18.9.2017

11
25.8.
6.9.

KAR

Accessible feedback/recommendation for a prescriber

VIDEŇSKÁ 1958 / 9
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INSTITUT KLINICKÉ A EXPERIMENTÁLNÍ MEDICÍNY
PRACOVISŤE LABORATORNÍCH METOD
Prednosta: Prof. MUDr. Antonín Jabor CSc.

Kontakt: MUDr. Janka Franeková, Ph.D., 5225, 737 205 963, Mgr. Kornélia Chrapková, PG Dip, 5274, 8243

Interpretace terapeutického monitorování léků

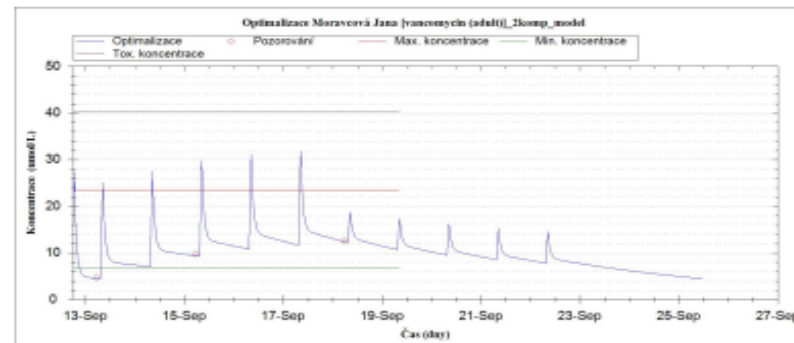
vancomycin (adult)

Výška: 172 cm

Osdělení: KUIP

Váha: 80 kg

Ošetřující lékař:



Historie

Datum & Čas	Dávka [mg]	Počet dávek	Třet [h]	Třet [h]	Konc [mg/l]	Konc [umol/l]	S ₀ Kreatinin [umol/l]
18/9/2017 8:00:47	250	5	24	1	0		
18/9/2017 8:00:05					18,81	12,8	
17/9/2017 7:01:39					0		187
15/9/2017 7:24:36					0		238
15/9/2017 6:00:38					14,83	9,8	
13/9/2017 8:00:23	750	5	24	1	0		
13/9/2017 5:55:07					6,87	4,6	
12/9/2017 18:00:29	1000	1		1	0		

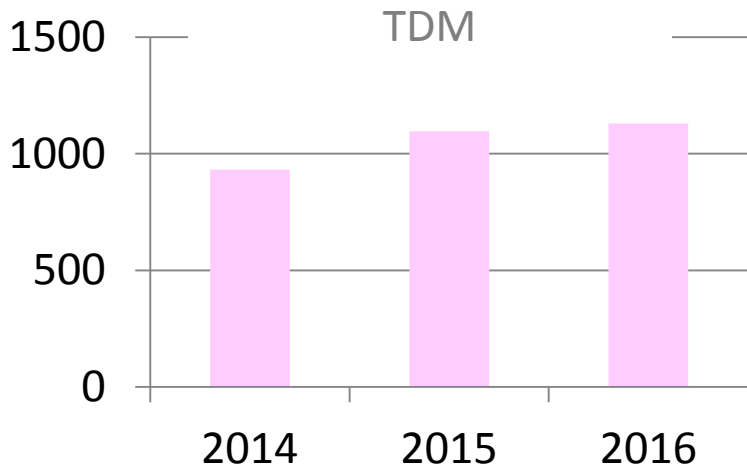
Doporučení: Dnešní (18.9.) naměřená koncentrace vancomycinu je 12,8 umol/l. Koncentrace je mírně nad optimem (8,9-10 umol/l). Doporučuji úpravu dávky na 250 mg & 24 hod v 8 hod, 30 min infuzí. Cílové rozmazi (8,9-10 umol/l). Prosím o kontrolní odběr koncentrace VAN před podáním ATB s ranními odběry ve středu (20.9.).

Děkuji

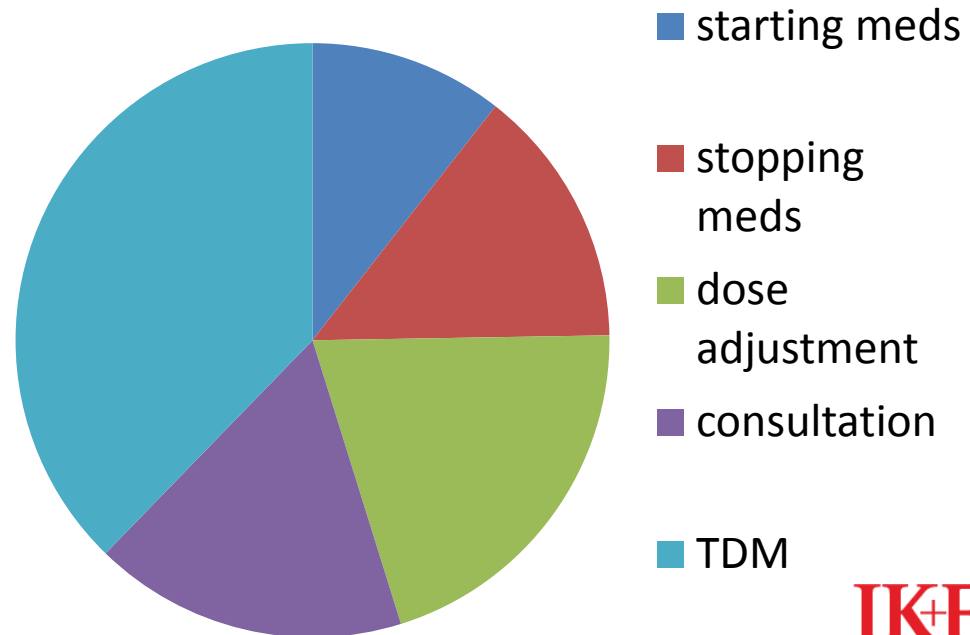
Dvořáčková E



TDM done by pharmacists



Pharmacists interventions 2016





TDM provided by clinical pharmacists was also a very important key for implementing clinical pharmacy service on wards.

Take home messages

- Assemble a multi-professional antimicrobial stewardship
- Establish a clear aim that is shared by all the stakeholders
- Start with core evidence-based stewardship interventions depending on local needs
- Plan measurement to demonstrate their impact
- Try not to implement all interventions at the same time
- Educate

Thank You!
😊