

Antibiotic policy group in the **Ghent University Hospital: experiences and results**

Séminaire de pathologie infectieuse
Cliniques Universitaires Saint-Luc, Bruxelles

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Content

- ➲ **Introduction**
- ➲ **Activities Antibiotic Policy Group UZ Gent**
 - ➲ Formulary and guidelines
 - ➲ Restrictions/support antibiotic prescribing
 - ➲ Surveillance
 - ➲ Antibiotic consumption and feedback
 - ➲ Resistance profiles
 - ➲ Audit
 - ➲ Drug Use Evaluation
 - ➲ Time of initialisation of antibiotics
 - ➲ Kidney failure and antibiotic dose
 - ➲ Therapeutic drug monitoring
 - ➲ Parenteral to oral conversion
 - ➲ Future projects
- ➲ **References for the clinical pharmacist**
- ➲ **Conclusion**

Introduction: who are we?



- ➲ **1.062 hospital beds**
- ➲ **34.000 admissions**
- ➲ **300.000 beddays**
- ➲ **4.799 staff members**
 - ➲ 660 physicians
 - ➲ 1.500 nurses
 - ➲ 20 pharmacists
- ➲ **Mean LOS: < 9 days**

Introduction: Antibiotic Policy Group

➊ Multidisciplinary composition (19)

- ➔ Infectiologists (4) (prof. dr. D. Vogelaers – president)
- ➔ Hospital hygiene physician (1)
- ➔ Microbiologist (1)
- ➔ Virologist (1)
- ➔ Intensive care physicians (4)
- ➔ Haematologist (1)
- ➔ Orthopedic physician (1)
- ➔ Pharmacists (5) (secretary)
- ➔ Scientific collaborators (2)

➋ Monthly plenary meeting

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Activities Antibiotic Policy Group: general information

- ➲ **Infectious Diseases Society of America and the Society for Healthcare Epidemiology of America Guidelines for Developing an Institutional Program to Enhance Antimicrobial Stewardship**
 - ➲ Clinical Infectious Diseases 2007;44:159–177

Table 1. Infectious Diseases Society of America–United States Public Health Service grading system for ranking recommendations in clinical guidelines.

Category, grade	Definition
Strength of recommendation	
A	Good evidence to support a recommendation for use
B	Moderate evidence to support a recommendation for use
C	Poor evidence to support a recommendation for use
Quality of evidence	
I	Evidence from ≥1 properly randomized, controlled trial
II	Evidence from ≥1 well-designed clinical trial, without randomization; from cohort or case-controlled analytic studies (preferably from >1 center); from multiple time-series; or from dramatic results from uncontrolled experiments
III	Evidence from opinions of respected authorities, based on clinical experience, descriptive studies, or reports of expert committees

NOTE. Adapted from [1].

Activities Antibiotic Policy Group: formulary and guidelines

- ➲ Local formulary (book and intranet):
 - ➲ Product information (dose, indication)
 - ➲ Guidelines: empirical, documented and prophylactic treatment (AI)
 - ➲ Recent guidelines
 - ➲ Empirical treatment for abdominal infections
 - ➲ HIV occupational post exposure prophylaxis
 - ➲ Bacterial meningitis and meningocccen sepsis
 - ➲ Extended and continuous infusions of antibiotics
 - ➲ Administration and monitoring of glycopeptides and aminoglycosides
 - ➲ Monthly revision of one chapter
 - ➲ The Sanford Guide for Antimicrobial therapy
 - ➲ Technical information for drug administration

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Activities Antibiotic Policy Group: restrictions/support antibiotic prescribing

- ➲ **Elimination of antibiotic stock on the wards**
- ➲ **Implementation of antibiotic/antimycotic order form (BII)**
 - ➲ Requirements for physicians
 - ➲ General
 - ➲ Motivation of therapy (empirical, documented and prophylactic treatment)
 - ➲ Start date
 - ➲ Clinical focus
 - ➲ Why no enteral administration (for high bioavailable products)?
 - ➲ Detailed motivation for linezolid
 - ➲ Information for physicians
 - ➲ Price per unit
 - ➲ IV/PO possibility
 - ➲ Attestation necessary
 - ➲ Limited dosages delivered by pharmacy



VPE

Patientgegevens (in te vullen of klever)

- Te leveren met volgende ronde
 Dringend leveren tegen uur

VOORSCHRIFT GERESERVEERDE ANTIBIOTICA EN ANTIMYCOTICA

(BUITEN RESERVEKAST)

EEN VOLLEDIG INGEVULD VOORSCHRIFT IS GELDIG
VOOR 4 DAGEN THERAPIE

Buiten de openingsuren van de apotheek zijn de producten
beschikbaar in de centrale spoedkast: -1K12C, -1K2 en 1K6
(kosten rekening te houden)

- teruggave uit centrale spoedkast genomen

MOTIVATIE THERAPIE

- EMPIRISCH (kiem onbekend)
 GERICHT (kiem vermelden):
 PROFYLACTISCH

STARTDATUM THERAPIE:

DIAGNOSE/KLINISCH BEELD (infectie).....

Geneesmiddel	Prijs per stuk (euro)	Advies	Dosis /24uur	Gevraagd aantal	Autor	Geleverd aantal	Apr.	Terug aantal
Ambisome 50 MG IV	161,33	Attest voorzakelijk				46322		
Abelcet 100 MG/20ML IV	124,86	Attest voorzakelijk				136986		
Avelox 400 MG PO	3,45					1542892		
Avelox 400 MG IV (2)	32,95	Eualteer N/P/O switch				1829637		
Cancidas 50 MG IV	483,88	Attest voorzakelijk				1596045		
Cancidas 70 MG IV	615,48	Attest voorzakelijk				1596144		
Ciproxine 250 MG co PO	0,69					186476		
Ciproxine 500 MG co PO	1,07					18707		
Ciproxine 200 MG IV (2)	15,49	Eualteer N/P/O switch				186773		
Ciproxine 400 MG IV (2)	27,88	Eualteer N/P/O switch				1292672		
Colistineb 2 miljoen E	6,24					2347794		
Diflucan 50 MG co PO	1,74					270610		
Diflucan 150 MG co PO	5,52					270808		
Diflucan 200 MG co PO	5,65					93238		
Diflucan 1400 MG 200 MG/5ML PO sir	41,23					1241994		
Diflucan 350 MG 50 MG/5ML PO sir	12,20					1241796		
Diflucan 200 MG IV (2)	17,41	Eualteer N/P/O switch				270907		

		100%
1	2	3

Diflucan 1400 MG/5ML PO sir	41,23			1241994	
Diflucan 350 MG 50 MG/5ML PO sir	12,20			1241796	
Diflucan 200 MG IV (2)	17,41	Buiteler M/PO switch		270907	
Glazidim 500 MG flacon	6,81			784809	
Glazidim 1 G flacon	13,20			392750	
Glazidim 2 G flacon	26,11			392651	
Maxiprim 1 G IV	13,85			370776	
Maxiprim 2 G IV	27,40			109668	
Meronem 500 MG IV	15,71			164306	
Meronem 1000 MG IV	28,28			164603	
Targocid 200 MG IV	32,42			911701	
Targocid 400 MG IV	61,75			911996	
Tavanic 250 MG co PO	2,16			1482318	
Tavanic 500 MG co PO	3,47			1327910	
Tavanic 500 MG IV (2)	30,35	Buiteler M/PO switch		1492909	
Tazocin 4 G IV	24,01			57210	
Vancocin 500 MG IV	11,20			1000484	
Vancocin 1000 MG IV	20,16			2047787	
Vfend 50 MG co PO	12,05	Attest voodzakelijk		1837258	
Vfend 200 MG co PO	42,13	Attest voodzakelijk		1717393	
Vfend 200 MG IV (2)	156,56	Attest voodzakelijk Buiteler M/PO switch		1717591	
Vfend PO susp 40 MG/ML 70 ML	674,80	Attest voodzakelijk		2207640	
Zyvoxid 600 MG co PO (1)	65,82	Indicate voodzakelijk		1610693	
Zyvoxid 600 MG IV (1)(2)	65,82	Buiteler M/PO switch Indicate voodzakelijk		1608319	
Zyvoxid 100 MG/5ML susp. 150 ML (1)	329,12	Indicate voodzakelijk		1672655	

(1) Indicatie Zyvoxid

- Infectie met MRSA, MR-CNS, ARE, VRE

Resistent of intermediair gevoelig aan glycopeptiden

Of

- Gevoelig aan glycopeptide maar
 - glycopeptide intolerant OF
 - IV toediening niet meer nodig/mogelijk

Zyvoxid opgestart in overleg met infectioloog. Ja / neen

Zyvoxid voor verderzetting thuis moet op ambulant voorschrijf met attest.

(2) Enterale

toediening (oraal of via maagsonde)
NIET mogelijk wegens:

- Braken
- Diarree
- Postoperatieve context
- Darmparalyse

Transport

- Buizenpost
- Koerier
- Ronde
- Datum:

Klaargezet door

Gecontroleerd door:

Datum, stempel**+ handtekening arts**

(conform KB 19/10/1978 art 5)

Activities Antibiotic Policy Group: restrictions/support antibiotic prescribing

④ Support by infectiologists/microbiologists

- ④ Weekly antibiotic meetings (SICU, MICU, orthopedic ward, pediatric ward ..) (AIII)
- ④ Consultation on demand
- ④ Spontaneous consultations based on (AI)
 - ④ Positive hemocultures
 - ④ Antibiotic prescriptions from pharmacy
 - ④ Link database pharmacy and microbiology
- ④ Education (BII)



Vorige



Ga naar

LabView - Microbiologie & Antibiotica

Laatste update pagina: 20:46

Patienten Li [REDACTED]

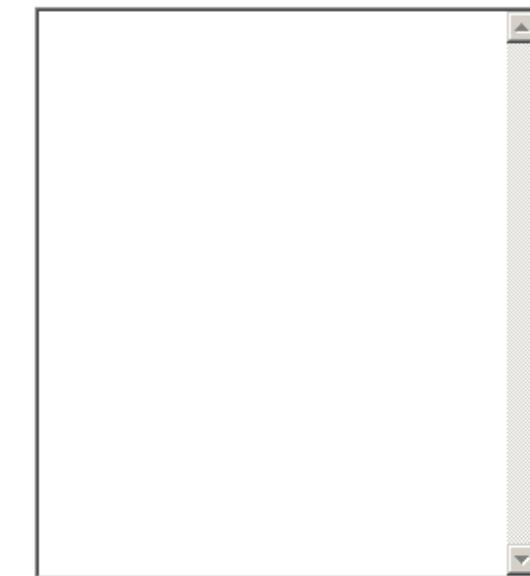
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Datum	Nr	Type staal	Kweek / Antibiotherapie	Aantal
4/12/2006	(061204-2381)	B.A.L.-vocht	1:: Pseudomonas aeruginosa ++	Antibiogram
4/12/2006	(061204-2100)	Keeluitstrijk	1:: Pseudomonas aeruginosa +-	Antibiogram
4/12/2006	(061204-2098)	Aspiraat bij geintubeerde	1:: Pseudomonas aeruginosa ++	Antibiogram

11/12/2006	GLAZIDIM 2 G BOLUS	6
11/12/2006	TAZOCIN FL 4 G/500 MG INJ	4
7/12/2006	TAZOCIN FL 4 G/500 MG INJ	16
6/12/2006	GLAZIDIM 2 G BOLUS	3
5/12/2006	AUGMENTIN 1 G IV FLAC	1
5/12/2006	GLAZIDIM 2 G BOLUS	4
30/11/2006	CIPROXINE COMP 500MG	2

9 stalen.



Geel: Volgt, Groen: geen kweek, Rood: Kweek

Vorige Pagina

Hoofdmenu

Heraanloggen

Help

Historische resultaten

Indien u problemen vaststelt, gebruik indien nodig Zielab en verwittig Dr. Tom Fiers (4565)

Activities Antibiotic Policy Group: restrictions/support antibiotic prescribing

④ Support by pharmacists

- ④ Prospective follow up of IV/PO in the central pharmacy (AI)
- ④ Clinical pharmacists
 - ④ Pediatric oncology
 - ④ Pediatric intensive care unit
 - ④ Geriatric unit
 - ④ Medical intensive care unit
 - ④ Abdominal surgery
 - ④ Emergency department (funded project “Clinical pharmacy - Federal government)
- ④ Education for nurses (BII)

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Activities Antibiotic Policy Group: surveillance and feedback of consumption (BIII)

- ➲ **ATC/DDD classification** (Anatomical Therapeutic Chemical classification /Defined Daily Dose)
 - ➲ Nominator
 - ➲ DDD
 - ➲ Costs (euro)
 - ➲ Denominator
 - ➲ Patient-days
 - ➲ Admissions
- ➲ **Level of analyses**
 - ➲ Global hospital level
 - ➲ Ward level
- ➲ **Monthly review by Antibiotic Policy Group**
- ➲ **Assigned physician by medical discipline for drug consumption**

Activities Antibiotic Policy Group: surveillance and feedback of consumption

Table 1. Evolution antibiotic consumption (correction for admissions and beddays)

year	amount (euro) per admission	% diff	amount per 100 beddays	% diff	DDD per admission	% diff	DDD per 100 beddays	% diff
2004	99,24	#VERW!	1.088,61	#VERW!	5,84	#VERW!	64,06	#VERW!
2005	104,54	5,34	1.146,92	5,36	5,84	0,01	64,08	0,03
2006	97,43	-6,81	1.112,13	-3,03	5,63	-3,58	64,29	0,33
2007	89,22	-8,42	1.040,07	-6,48	5,57	-1,15	64,90	0,94

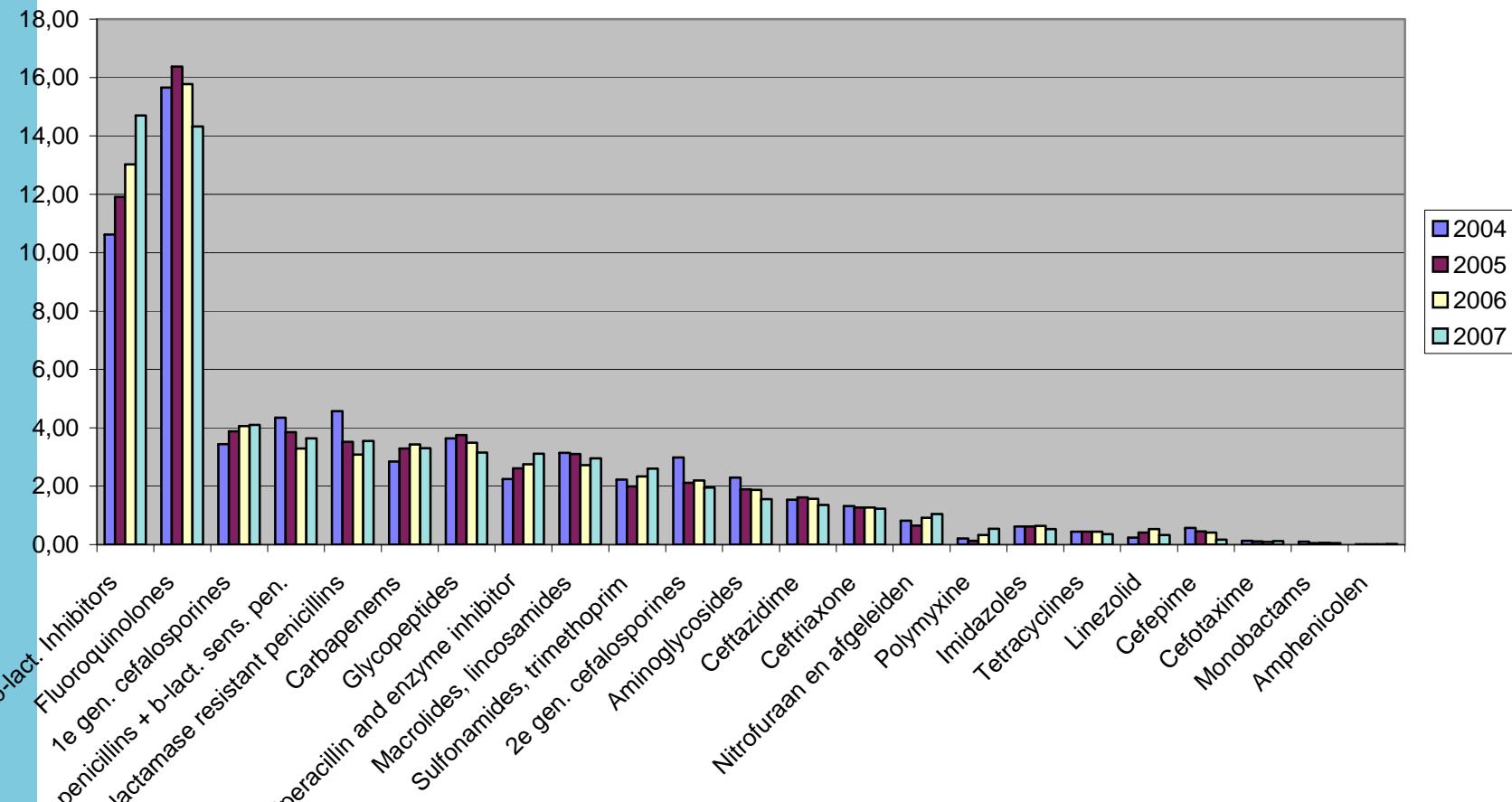
Table 2. Evolution antibiotic consumption in DDD per 100 beddays for products with high bioavailability

year	DDD IV	% diff	DDD PO	% diff	DDD PO+IV	% diff	%IV/IV+Po	% diff
2004	7,37	#VERW!	10,18	#VERW!	18	#VERW!	42	#VERW!
2005	7,16	-2,82	11,15	9,54	18	4,35	39	-6,87
2006	5,42	-24,38	12,09	8,49	18	-4,37	31	-20,92
2007	3,72	-31,24	12,35	2,10	16	-8,21	23	-25,09

Activities Antibiotic Policy Group:

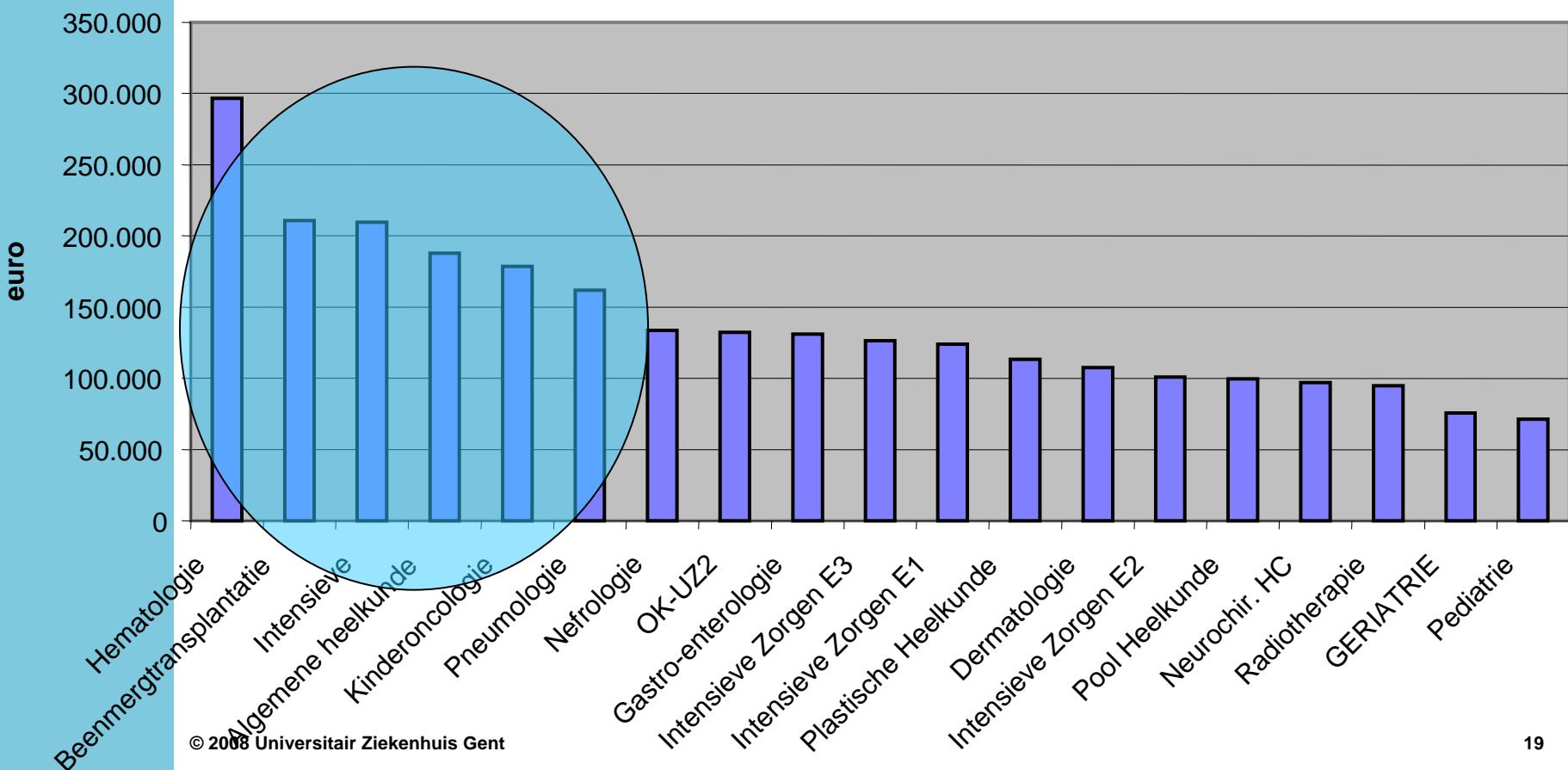
surveillance and feedback of consumption

Evolution antibiotic consumption



Activities Antibiotic Policy Group: surveillance and feedback of consumption

Top 20 : Wards with highest antibiotic consumption (in euro)





Activities Antibiotic Policy Group: surveillance of resistance profiles

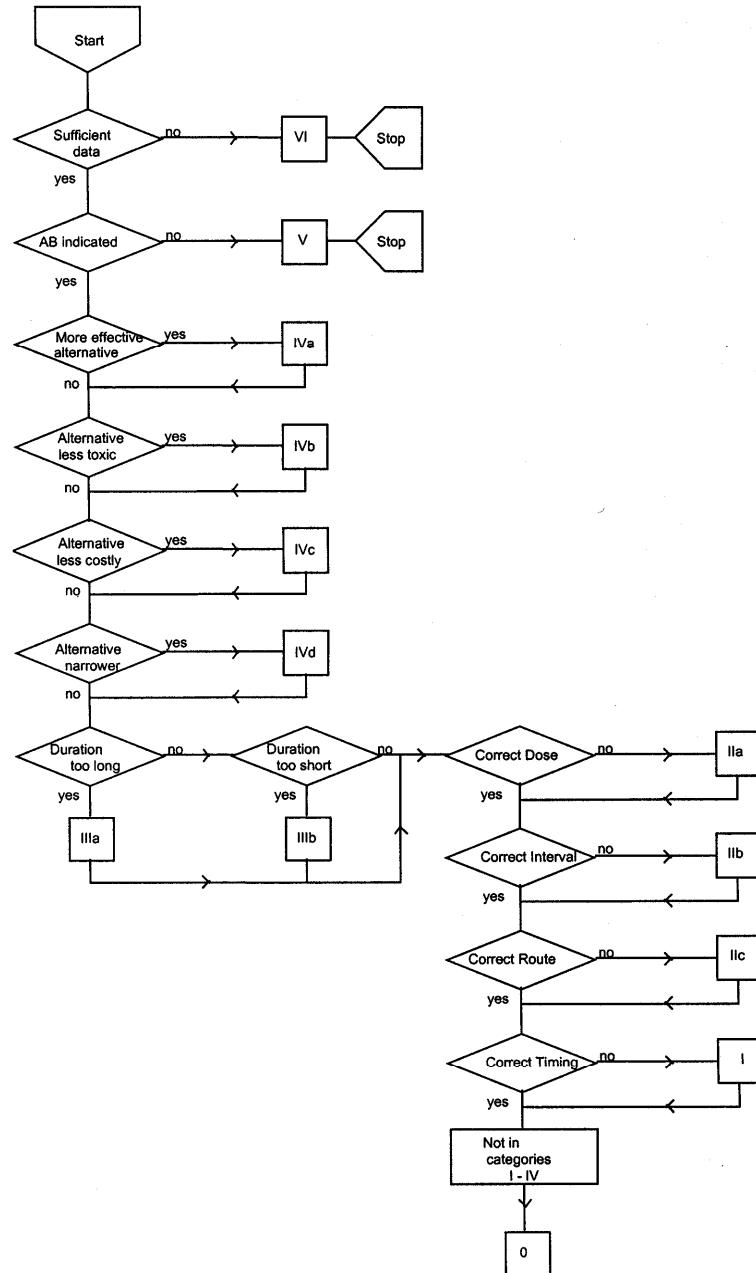
- ⌚ Hospital hygiene committee
- ⌚ Resistance patterns are available in the formulary

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Activities Antibiotic Policy Group: Audit – Drug Use Evaluation (AIII)

- ⌚ Experience with vancomycin, teicoplanin, fluoroquinolones, piperacilline/tazobactam
- ⌚ Main outcome measures
 - ⌚ Degree of concordance with predefined intra-muros published guidelines
 - ⌚ initial empiric therapy
 - ⌚ subsequent therapy beyond 72 hours
 - ⌚ Evaluation by infectiologist (Gyssens et al. J Antimicr Chemother 1992;30:724-727)
 - ⌚ indication, duration, dosage



Activities Antibiotic Policy Group: Audit – Drug Use Evaluation

- ➲ Retrospective observational study in 2001
- ➲ Surgical intensive care unit in Ghent University Hospital
- ➲ Outcome:
 - ➲ Degree of concordance of initial empiric and subsequent directed therapy beyond 72 hours with piperacilline/tazobactam with predefined intra-muros published guidelines.

Activities Antibiotic Policy Group: Audit – Drug Use Evaluation

	Indication antibiotic therapy	Number of courses	Inappropriate initial therapy	More effective alternative	Less broad alternative	Inappropriate continuation (>72hours)	More effective alternative	Less broad alternative	Excessive length
	PIPERACILLINTAZOBACTAM COURSES								
	Abdominal sepsis	9	3	1	2	4	2	2	-
	Late onset nosocomial infection	14	3	1	2	8	1	5	2
	Others	4	4	2	2	2	1	1	-
	Total courses	27	10	4	6	14	4	8	2
	Percentage (%)	100	37	15	22	52	15	30	7

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Activities Antibiotic Policy Group: Audit: time of initialisation of antibiotics

- ➲ **Concomitant observational study (AIII)**
- ➲ **Patient inclusion criteria**
 - ➲ a proven or suspected infection
 - ➲ community acquired pneumonia (CAP), meningitis, pyelonephritis, erysipelas, intra-abdominal infections and exacerbation of chronic obstructive pulmonary disease (COPD)
 - ➲ transfer to a ward after admission to the ED
- ➲ **Measured indicators**
 - ➲ time interval between admission to ED and first antibiotic administration
 - ➲ antibiotic selection in accordance with local guidelines

Activities Antibiotic Policy Group: Audit: time of initialisation of antibiotics

Patients characteristics

Number of patients	65
Type of infection	
CAP (%)	35 (54)
<i>intra-abdominal infections</i> (%)	9 (14)
exacerbation of COPD (%)	8 (12)
<i>pyelonephritis</i> (%)	3 (5)
<i>erysipelas</i> (%)	3 (5)
<i>meningitis</i> (%)	1 (2)
others (%)	6 (9)
Mean age, years	61
Gender ratio; M:F	32:33

Outcomes

Time interval (hours) between admission to ED and first antibiotic administration (S.D)	3,9 (3,7)
<i>first dose in the E.D. (N=46) (S.D)</i>	2,4 (1.3)*
<i>first dose on the ward (N=19) (S.D)</i>	7,5 (4,9)*
Percentage treatments in accordance with local guidelines %	86%

* P < 0.01

Conclusion

- Administration of the first antibiotic dose on the ward is associated with longer time intervals.

Improvements

- First dose on ED
- Critical antibiotics available on ED
- Accelerate delivery of antibiotics by pharmacy

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Activities Antibiotic Policy Group: Audit: Kidney failure and antibiotic dose (AIII)

- ⦿ **Retrospective observational study**
- ⦿ **Inclusion criteria**
 - ⦿ > 18 years
 - ⦿ Admitted on internal medicine, abdominal surgery or nephrology (April 2006 – March 2007)
 - ⦿ eGFR < 60 mL/min/1,73m² (Modification of Diet in Renal Disease (MDRD) formula)
 - ⦿ Antibiotic-treatment
- ⦿ **Exclusion criteria**
 - ⦿ Dialysis patiënts
- ⦿ **Measurement kidney function**
 - ⦿ eGFR met MDRD-formula (serum creatinine, age, sex and race)
 - ⦿ creatinine clearance with Cockcroft & Gault (if weight available)
- ⦿ **Evaluation antibiotic dose**
 - ⦿ By panel (infectiologist, intensivist, pharmacist) based on creatinine clearance

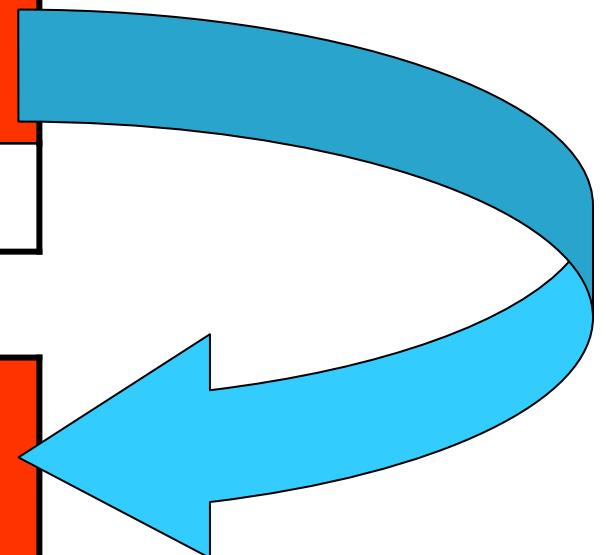
Activities Antibiotic Policy Group: Audit: Kidney failure and antibiotic dose

Patient characteristics

Number of patients	99
Nephrology	27
Internal medicine	21
Abdominal surgery	51
Mean age (SD) (years)	65 (15)

Activities Antibiotic Policy Group: Audit: Kidney failure and antibiotic dose

Number administered doses	Number correct doses (%)	Number incorrect doses (%)
1.364	1.068 (78,3%)	296 (21,7%)



Number administered doses with overdose (%)	Number correct doses with underdose (%)	Number doses not possible to evaluate
183 (61,8%)	74 (25,0%)	39 (13,2%)

Activities Antibiotic Policy Group: Audit: Kidney failure and antibiotic dose

➊ Improvements

- ➏ Warning in the on-line electronic laboratory database
 - ➏ Patiens between 18 and 70 years and MDRD < 60 ML/MIN/1.73m²
 - ➏ “Possible increased kidney function. Take this in account by selecting and dosing drugs”.
- ➏ Letter to al physicians
- ➏ Task for clinical pharmacist

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Activities Antibiotic Policy Group: Audit: Therapeutic drug monitoring (AIII)

- ➲ **Retrospective observational study**
- ➲ **Inclusion criteria**
 - ➲ Treatment with teicoplanine or vancomycin
 - ➲ Admission on infectious diseases, abdominal surgery and nephrology
- ➲ **Exclusion criteria**
 - ➲ =< 1 day treatment
- ➲ **Outcome measurement**
 - ➲ Degree of concordance with intra-muros guidelines
- ➲ **Evaluation antibiotic dose and level**
 - ➲ Panel of 2 senior pharmacists

Activities Antibiotic Policy Group: Audit: Audit: Therapeutic drug monitoring

Patient characteristics - results

Number of patients	50
Mean age (SD) (years)	56,8 (13,4)
Number vancomycin treatments (continuous infusion)	39
<i>Mean treatment duration (days)</i>	14,6
<i>Number doses evaluated</i>	583
Number teicoplanin treatments	11
<i>Mean treatment duration (days)</i>	10,8
<i>Number doses evaluated</i>	108

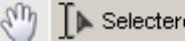
Activities Antibiotic Policy Group: Audit: Therapeutic drug monitoring - vancomycin

Correct loading dose	17/28	61%
Correct maintenance dose day 1	25/30	83%
Correct maintenance dose day 2	25/30	83%
Number correct doses	268/583	46%
Correct concentration infusion	35/39	90%

Improvements

- Update guidelines for administration and monitoring of glycopeptides (flow chart)
- Pop-up in the on-line electronic laboratory database linked to results drug monitoring

opslaan



Selecteren



122%



Adobe Reader 7.0

**Zie richtlijn 1A
(INTERMITTENT INFUUS)**

CONTINU INFUUS

START DAG 1: 0 uur

1. LADINGSDOSIS

15mg/kg lichaamsgewicht oplossen in glucose 5% of NaCl 0.9 %

- 1) Dosis geven over 2 uur (Red Man syndrome)
- 2) Maximale concentratie infuus 0.5 % (1% bij vochtrestrictie)
 - Vb 500 MG vancomycine in 50 ML NaCL 0,9% of glucose 5%
 - Vb 1000 MG vancomycine in 100 ML NaCL 0,9% of glucose 5%
 - Vb 1500 MG vancomycine in 250 ML NaCL 0,9% of glucose 5%
- 3) Check andere medicatie (zie 2.11.) (eventueel infuus maximaal 2 x 2 uur/ dag onderbreken)

DAG 1: 2 uur later

2. Onmiddellijk NA beëindigen LADINGSDOSIS

GEEF ONDERHOUDSDOSIS:

30 mg/kg over 24 uur

- 1) Maximale concentratie infuus 0.5 % (1% bij vochtrestrictie)
 - Vb 1000 MG vancomycine in 100 ML NaCL 0,9% of gluc 5%
 - Vb. 2000 MG vancomycine in 250 ML NaCL 0,9% of gluc 5%
 - Vb 3000 MG vancomycine in 500 ML NaCL 0,9% of gluc 5%

- 2) Check andere medicatie (zie 2.11.) (eventueel infuus maximaal 2 x 2 uur/ dag onderbreken)

NA 24 uur

Bloedafname voor spiegelbepaling niet via infuus waarlangs medicatie is toegediend.

Bloedafname voor spiegelbepaling

DAG 2

3. GEEF ONDERHOUDSDOSIS:

30 mg/kg over 24 uur

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Activities Antibiotic Policy Group: Audit: Parenteral to oral conversion (AIII)

**“Big diseases need big treatments
and big treatments come in
syringes”**



Activities Antibiotic Policy Group: Audit: Parenteral to oral conversion

- ➡ **Definitions** (Nathwani et al. JAC 1997;39:441-446)
- ➡ **Streamlining therapy:**
 - ➡ narrowing of the target specific pathogen, encompassing the overall strategy of altering the antimicrobial regimen in response to culture and susceptibility results (Quintilani)
- ➡ **Sequential therapy:**
 - ➡ Conversion from IV to oral formulation of the same medication (maintaining equivalent potency)
- ➡ **Step-down therapy:**
 - ➡ conversion from IV to oral agent of the same or different class of agent, with reduction in potency
- ➡ **Switch therapy:**
 - ➡ conversion from IV therapy with one drug to an oral formulation of a different medication without losing potency

Activities Antibiotic Policy Group: Audit: Parenteral to oral conversion

➲ Advantage for patiënts

- ➲ faster mobilisation
- ➲ shorter stay in hospital
- ➲ reduced risk of adverse effects
 - ➲ pulmonary embolism
 - ➲ phlebitis (no IV line)
 - ➲ catheter associated infection

➲ Advantage for hospital/community budget

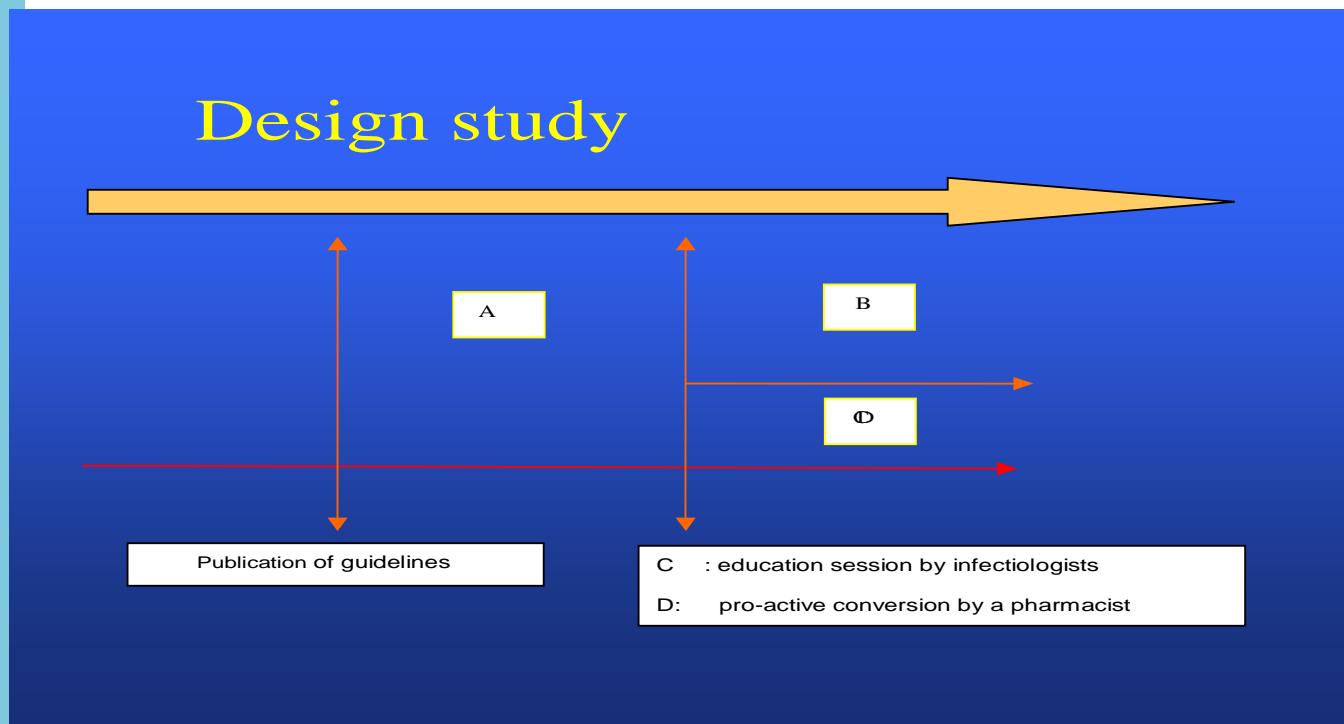
- ➲ reduction in drug acquisition cost
- ➲ reduction in drug wastage
- ➲ reduction laboratory costs, monitoring serum concentration and renal function
- ➲ reduced preparation and administration time

Activities Antibiotic Policy Group: Audit: Parenteral to oral conversion

- ➲ Guidelines for IV/PO switch differ according to different literature data.
 - ➲ body temperature < 38°C during 24 hours
 - ➲ decreasing or normal leukocyte count
 - ➲ no unexplained tachycardia
 - ➲ intact functional gastro-intestinal tractus absence of malabsorption (no vomiting, no diarrhoea)
 - ➲ a functional gastric feeding tube
 - ➲ no planned operation within 24 hours
 - ➲ no severe sepsis
- ➲ For which products?
 - ➲ Levofloxacin, ciprofloxacin, moxifloxacin, clindamycine, metronidazol, fluconazol, voriconazol linezolid
- ➲ Pitfalls
 - ➲ Drug-drug interactions
 - ➲ Drug-food interactions
 - ➲ Crushing of tablets

Activities Antibiotic Policy Group: Audit: Parenteral to oral conversion

- controlled before and after prospective study



Characteristic	Period B	Period C	Period D	P
	N= 36	N= 21	N=24	
Type of infection				
Urinary tract infection	11	5	12	0,117
Skin and soft tissue infection	4	7	2	0,048
gastro-intestinal infections	7	0	6	0,061
Pneumonia	3	1	1	0,857
Prothesis-infection	0	5	0	0,001
Osteomyelitis	0	2	0	0,065
Others	11	1	3	0,037
Pathology index (a) (source: Medicare) (S.D)	1,7 (1,4)	1,5 (0,9)	1,6 (1,4)	0,966
Mean age, years (S.D.)	62 (15)	59 (16)	58 (15)	0,574
Gender ratio, m/v	22/14	9/12	14/10	0,398
Mean time start therapy until fulfilment of criteria (days) (S.D)	5,2 (5,2)	5,8 (6,2)	5,0 (4,7)	0,934

Activities Antibiotic Policy Group: Audit: Parenteral to oral conversion

Patient outcome

	GROUP A	GROEP B	GROEP C	P
	N= 36	N= 21	N=24	
Mean LOS (S.D)	29,8 (27,4)	23,9 (19,8)	24,7 (21,9)	0,619
Mean duration FQ IV treatment in days (S.D)-	8,6 (6,6)	9,3 (7,9)	4,7 (4,5)	0,029
Mean extra IV treatment in days (S.D.)	4,1 (5,8)	3,5 (4,9)	1,0 (1,3)	0,006
Mean cost caused by extra IV treatment in euro (SD) (a)	188,7 (292,9)	103,6 (131,5)	44,8 (60,2)	0,037

Activities Antibiotic Policy Group: Audit: Parenteral to oral conversion

Year 2005



**Slikken
is soms beter
dan prikken!**

WAART

- Antibiotica moeten in het algemeen niet meer intraveneus gegeven worden.
- Gedreven door een aantal mogelijk orale alternatieven.

WAARNEEMEN

- Invoering van antibioticale middelen met een goede bioavailability (amideen en wortelkanaalroute bereikt de cel of bacterie rechtstreeks via de bloed sel) na toediening (Tazocin®, ciprofloxacin (Ciproflox®), amikacine (Amikin®), ciprofloxacin (Ciproflox®), ciprofloxacin (Ciproflox®), norfloxacin (Noroxin®), levofloxacin (Levofloxacin®), gatifloxacine (Gatifloxin®)).
- Patiënt kan meer of vroegere voorkeuring veranderen via mogelijke.

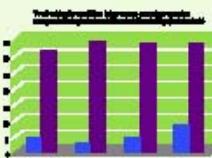
2. Onderzoek moet andere gezondheidsoverlast of reënval. Dit kan worden veroorzaakt door voorkeursgedrag te wijzigen. (d: UZ Gent, Interne Apotheek en Zorggebied Slikken is soms beter dan prikken).

WAAROM?

- Conformiteit waarbij een intraveneuze behandeling die helemaal vergaard werd met een orale dosering naar deze therapie moet kunnen doorgaan.
- Werkzaam:
 - langer niet meer nodig (meer beschikbaarheid, verlaagd risico op complicaties en risico's)
 - behandeling
 - patiënt
 - medische en zorgkosten

HIGH RISK

Prof. dr. D.Wagenveldt, MSc
Apo. E.Bogaerts, MSc



Tijd	Percentage
1e week	10%
2e week	20%
3e week	30%
4e week	40%
5e week	50%
6e week	60%
7e week	70%
8e week	80%
9e week	90%
10e week	100%



Activities Antibiotic Policy Group: Audit: Parenteral to oral conversion

Year 2007

Further promotion:

- By clinical pharmacists
- Frequently audits
- Follow up usage indicator
 - % IV/IV+PO
 - in 2007: 23%



SLIKKEN BLIJFT SOMS BETER DAN PRIKKEN



Proficiat!

Dankzij uw medewerking bij het frequenter starten of sneller switchen van producten met een hoge biologische beschikbaarheid werd in 2006 bijna 65.000 euro bespaard en de patiënt even doeltreffend behandeld.

Tavanic 500 MG PO (=3,74 €) is 10 x goedkoper i.v.m. IV (=30,35 €).

DE ANTIBIOTICABELEIDSGROEP REKENT VERDER OP U!

Producten met een hoge biologische beschikbaarheid zijn: levofloxacin (Tavanic®), ciprofloxacin (Ciproxine®), moxifloxacin (Avelox®), linezolid (Zyvoxid®), clindamycine (Dalacin®), metronidazol (Flagyl®), fluconazol (Diflucan®) en voriconazol (Vfend®).

Consulteer ook Intranet
<http://serapis/docz/apotheek/UZGeneesmiddelenbulletin24Prikken.pdf>



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Future projects

- ➲ Reducing duration of antibiotic treatment
 - ➲ Procalcitonin: better diagnostic marker?
- ➲ Extended and continuous infusion of antibiotics (All)
 - ➲ Implementation in daily practice
 - ➲ Pitfalls
 - ➲ Incompatibilities
 - ➲ Loadingdose
- ➲ ABS project: validation of indicators
 - ➲ *Staphylococcus aureus* bacteremia
 - ➲ Prophylaxis in surgery
 - ➲ IV/PO switch
- ➲ Decision support system into CPOE (computer physician order entry) (BI)

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References for the clinical pharmacist

Electronic databases	- Up to date (http://www.utdol.com/online/about/contact_us.html) - Micromedex (www.micromedex.com/products/clinicalxpert) - Clinical Pharmacology (http://www.goldstandard.com)
Handbooks	- Handbook of injectable drugs. Trissel L. (ISBN 1-58528-016-X) - Interpretation laboratory data. M. Lee ASHP (ISBN 1-58528-059-3) - Clinical pharmacy and therapeutics. Walker R., Edwards C (0-443-07137-3) - Antibiotic policies. Gould I. (ISBN 0-306-48500-1) - The Sanford Guide to antimicrobial therapy
Websites	- The Infectious Diseases Society of America (IDSA). http://www.idsociety.org/ - European Society of Clinical Microbiology and Infectious Diseases. http://www.escmid.org/ - The Cochrane Collaboration. Cochrane Reviews “Infectious Diseases”. http://www.cochrane.org/reviews/en/topics/72.html - European Society of Clinical Pharmacy. http://www.escpweb.org/

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Conclusion

- ➲ Multidisciplinary team approach is necessary
- ➲ Combination of different interventions can reduce inappropriate use of antibiotics
- ➲ Quality indicators in development or already available

Thank you for your attention