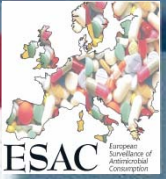


# Rationale



- Survey of antibiotic consumption is the first step towards optimising its use
- Benchmarking is an important initial step to identify needs of improvement

# ESAC-2 Hospital Care Subproject: 2005-2007

Ansari F, Johnston A, Molana H, Davey P, Dundee, UK  
Erntell M, Sweden  
Neotide, Finland  
Ferech M, Goossens H, Antwerp, Belgium  
The ESAC II hospital care study group



- ***Objectives:***

1. Standardise a measure for longitudinal analysis of antibiotic use by hospitals
  - Numerator: Defined Daily Doses
  - Denominator: comparison of bed days *versus* admissions
2. Standardise a method for point prevalence survey (cross sectional analysis) of antibiotic use
3. To develop methods that can be applied in all participating countries in ESAC-3.

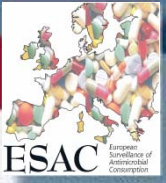
- ***Project components***

1. Questionnaire (hospital characteristics)
2. Longitudinal Survey (pharmacy stock data, monthly, 6 yrs, J01plus, bed days and admissions)
3. Point Prevalence Survey (STRAMA tool adopted for ESAC-2 HC)

- Standardised data from one hospital in each participating country
- A practical method that can be rolled out
  - To other hospitals in each participating country
  - To other countries in ESAC-3
- A platform for statistical analysis:
  - Trends within hospitals
  - Comparison between hospitals or countries

# Composition Hospital Subproject

5

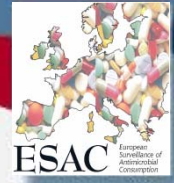


- **Members: 23 countries (including England, Northern Ireland, Scotland, Wales)**

Austria (Sigrid Metz), Belgium (Hilde Jansens), Czech Republic (Jiri Vlcek) Croatia (Arjana Tambić Andrašević), Denmark (Birgit Molstad), England (Conor Jamieson), Estonia (Piret Mitt), Finland (Outi Lyytikainen), France (Isabelle Patry, Xavier Bertrand), Germany (Michaela Steib-Bauert), Greece (Anastasia Antoniadou), Latvia (Elina Pujate), Lithuania (Ilma Bertulyte), Malta (Peter Zarb), Netherlands (Margreet Filius, Claire van Nispen tot Pennerden), Northern Ireland (Sheila Maltby), Norway (Cecile Syrrist), Poland (Pawel Grzesiowski), Slovenia (Milan Cizman), Sweden (Mats Erntell), Scotland (Faranak Ansari), Turkey (Denis Gür) and Wales (Maggie Heginbotham)

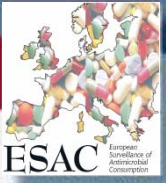
# ESAC-2 Longitudinal Study 2005-7

## *Methods*



- Assignments of ATC/DDD and calculations of DBD (DDD/100 bed days) & DAD (DDD/100 Admissions) centrally in Dundee
- Estimation of discharge (DIS) and LOS (Length of stay) from Occupied Bed Days (OBD) and Admissions (AD)
- Time series analysis for DDD, DBD, DAD, LOS, DIS and effect of clinical activity variables on DDD
- Percentage of annual change for each variable.

# Calculation of Clinical Activity



## Bed days

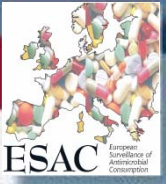
- **Occupied bed days (OBD)** calculated from daily census of the number of occupied beds at a specified time (e.g. midnight).
- **Hospital care days** calculated from daily census of the number of patients that have occupied each bed in a 24 hour period (can be >1 patient per day).
- **Patient days** calculated based on length of stay, e.g. date of discharge-date of admission -1.

## Admissions

- Does the number of admissions include day cases? If so can day cases be excluded? If not, does the hospital have data about % day cases by year?
- How are birth admissions included? Do they count as one admission or two? Has a consistent method been used throughout the study period?



# Time Series Analysis



- For time series analysis we built the following dynamic regression model:

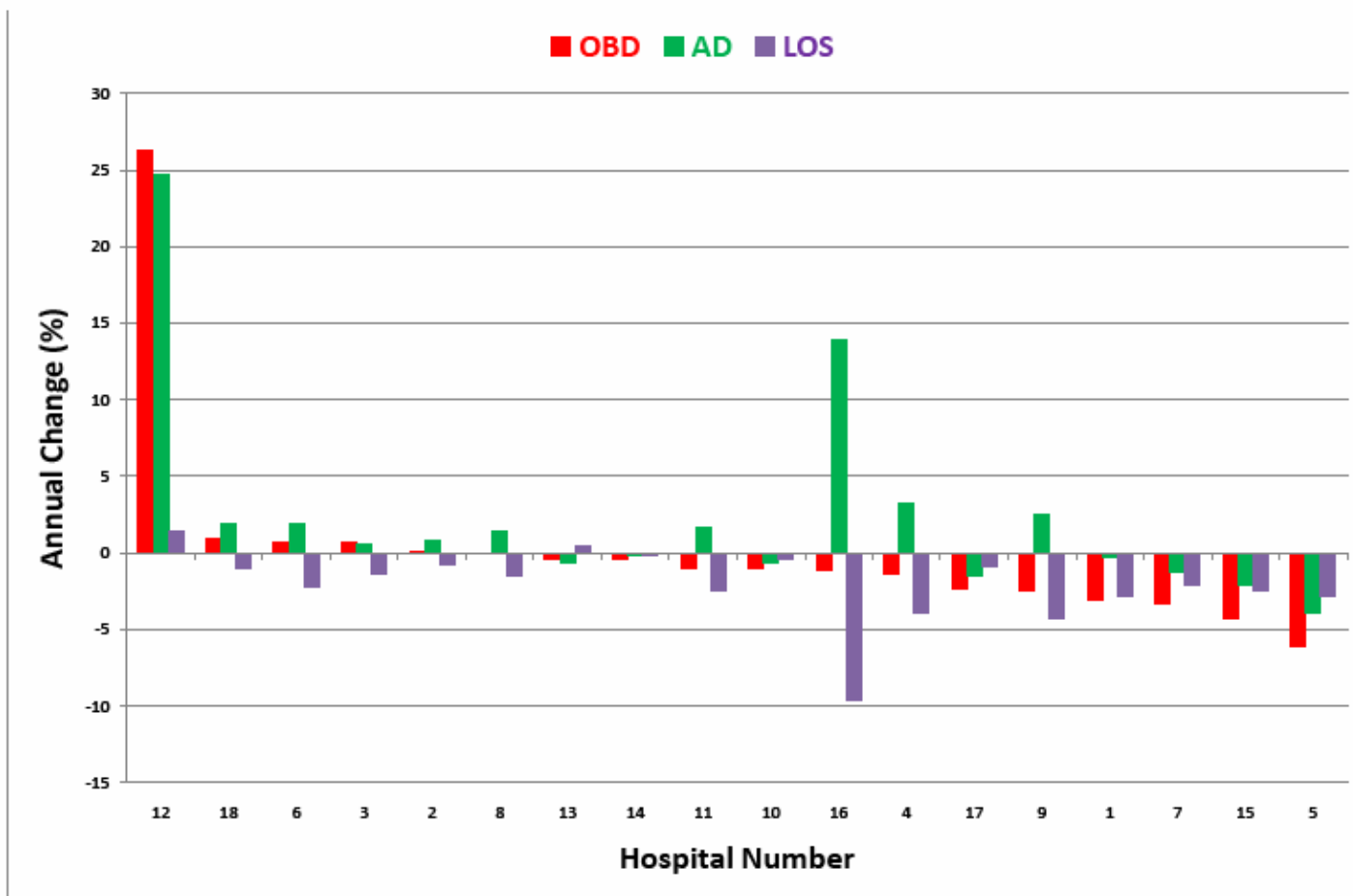
$$Y_t = \sum_{s=0}^S (a_s Y_{t-1-s} + \beta_s A_{t-s} + \gamma_s B_{t-s} + \delta_s R_{t-s} + \phi_s L_{t-s}) + \sum_{j=0}^J \mu_j t^j + \sum_{i=2}^{12} \theta_i DM_{i,t} + \sum_{k=1}^K \lambda_k DI_{k,t} + \varepsilon_t$$

- In order to scale the changes and find a way to compare the trends between hospitals we calculated the percentage of change/ year according the following formula:

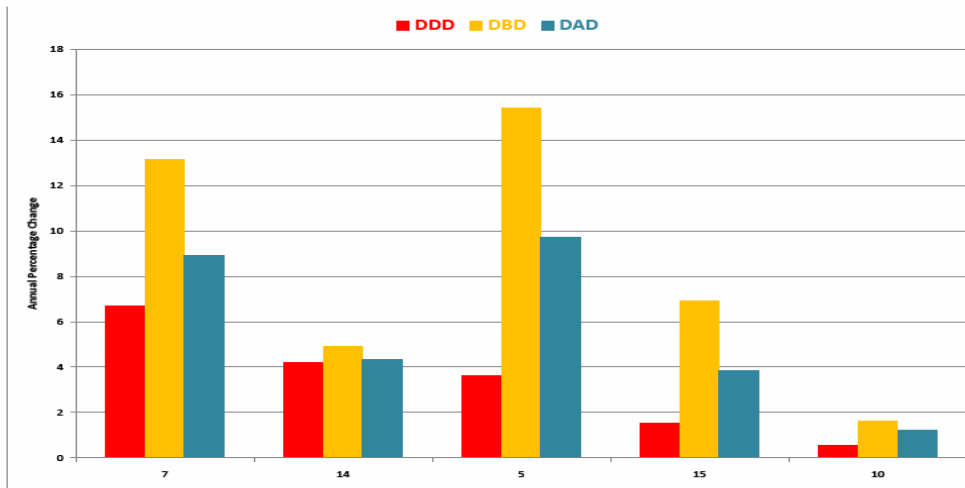
$$\% \Delta \text{ Variable} = [(\text{Slope} * 12) / \text{Intercept}] * 100$$



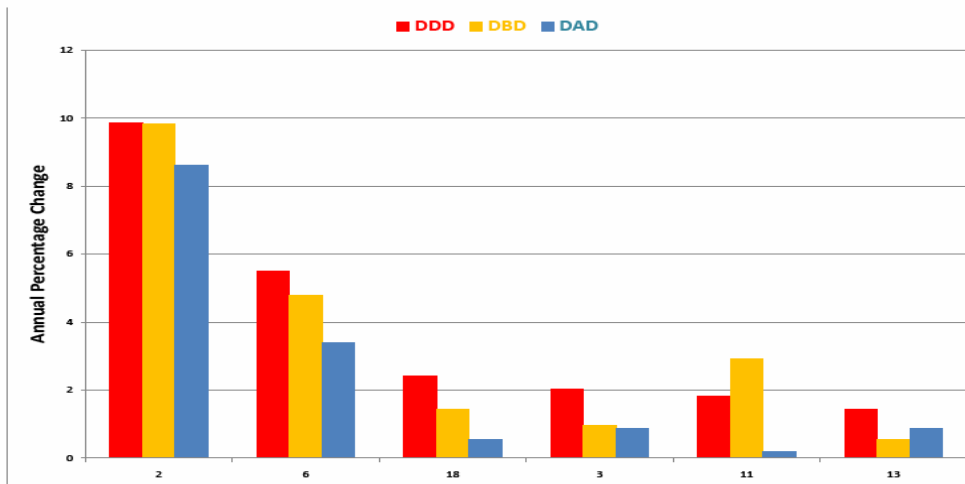
# Annual Changes (%) in OBD, AD and LOS; ranked by change in AD



# Annual Changes (%) in Antibiotic Use Measured as DDD, DBD, DAD; ranked by change in DDD

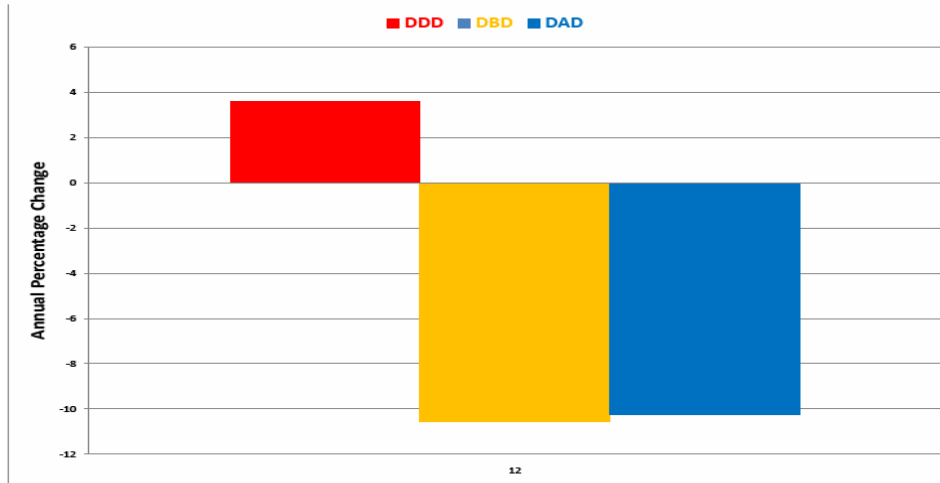


DDD increasing  
**DESPITE**  
Clinical activity  
decreasing



DDD increasing  
**PARTIALLY  
EXPLAINED BY**  
Clinical activity  
increasing

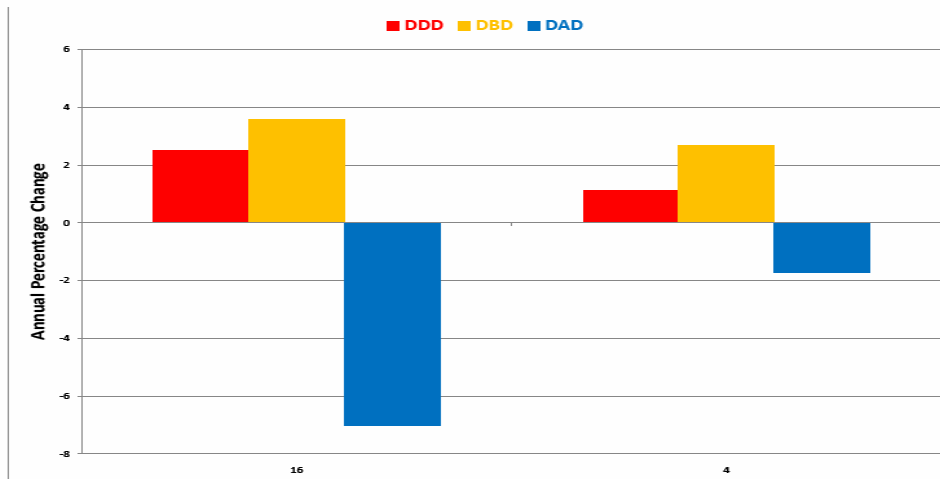
# Annual Changes (%) in Antibiotic Use Measured as DDD, DBD, DAD; ranked by change in DDD



DDD increasing

**ENTIRELY  
EXPLAINED BY**

Clinical activity  
increasing

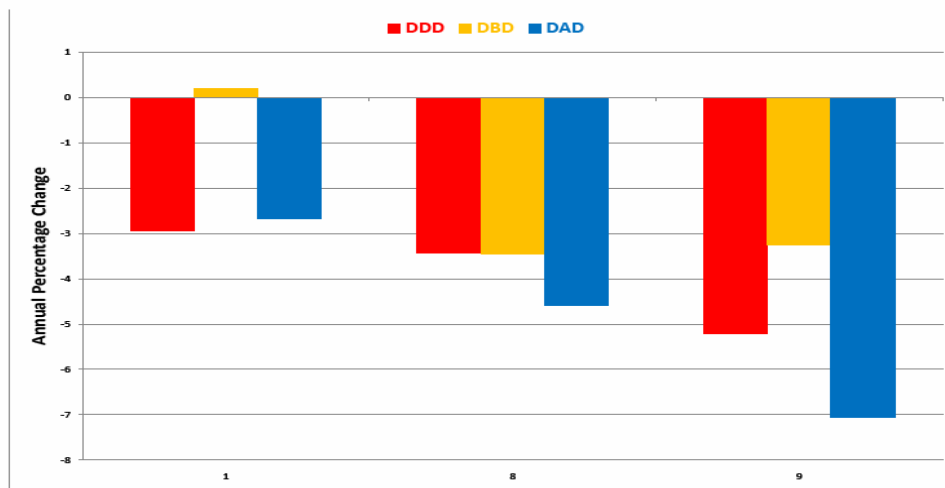


DDD (& DBD)  
increasing

**ENTIRELY  
EXPLAINED BY**

Admissions  
increasing &  
LOS decreasing

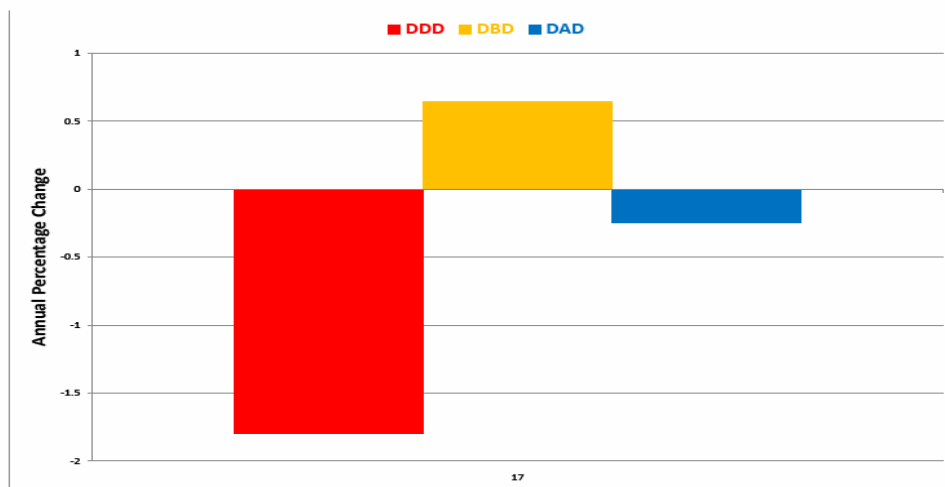
# Annual Changes (%) in Antibiotic Use Measured as DDD, DBD, DAD; ranked by change in DDD



DDD decreasing

**ONLY PARTIALLY  
EXPLAINED BY**

Clinical activity  
decreasing



DDD decreasing

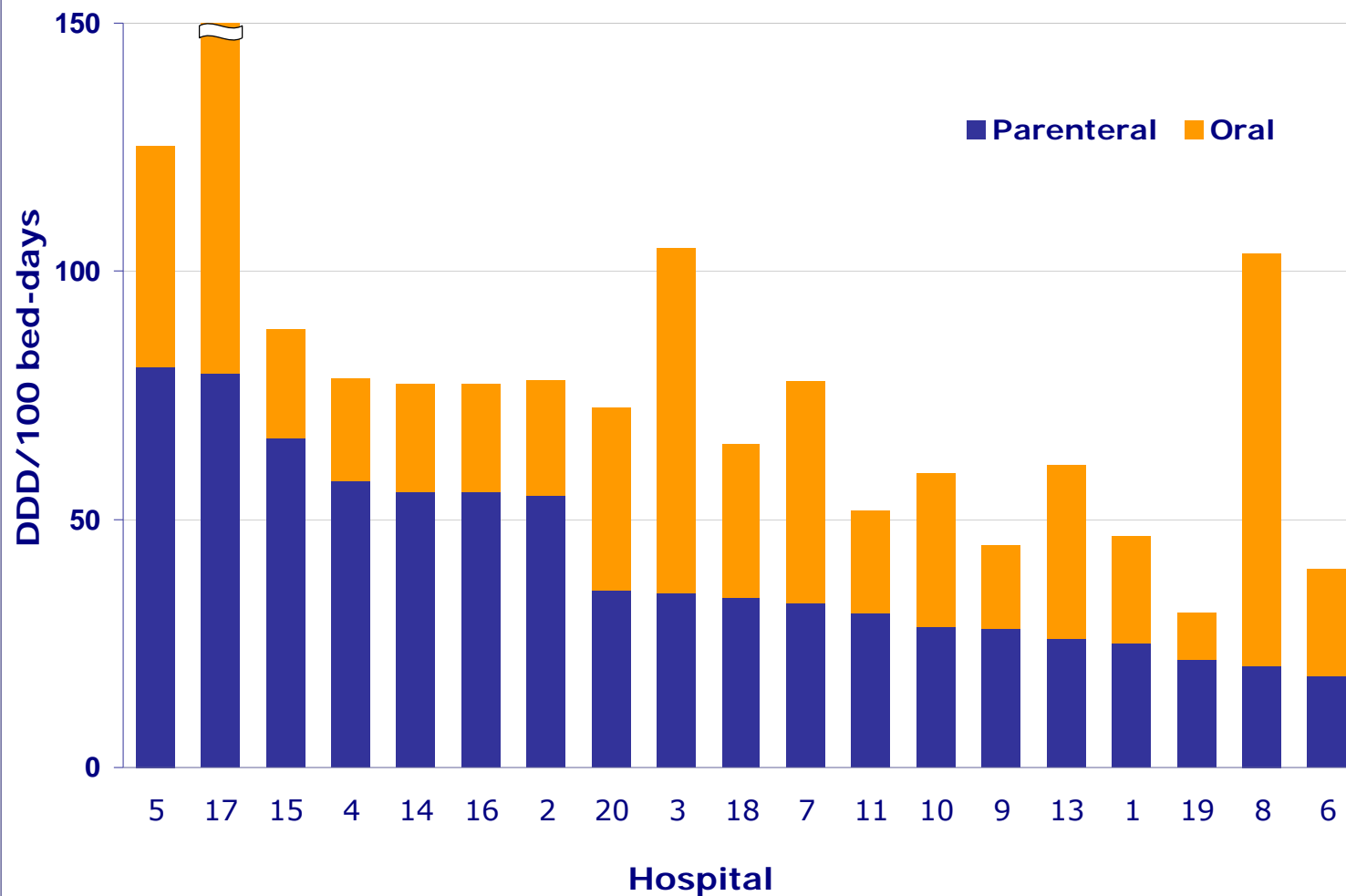
**ENTIRELY  
EXPLAINED BY**

Clinical activity  
decreasing

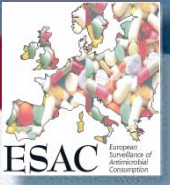
# ESAC-2, HC LS-Results



## Total Use of Antibacterials, 2005

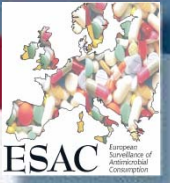


# Conclusions, Longitudinal Survey



- While patient specific data are not available at international level, DDD, despite limitations, is still the cornerstone in DUR studies.
- Present DDD with adjustment for clinical activity:
  - LOS decreased over time in 16 of 18 hospitals
  - Antibiotic use can be influenced by the number of admissions and by the patients' length of stay
  - Adjustment for clinical activity should be done with both admissions and occupied bed days
  - These results have relevance to microbiological surveillance as well as to prescribing surveillance
- Big variation in IV antibiotic use

# ESAC-2, Point Prevalence Study



## Pattern of antibacterial use in relation to diagnosis in 20 European hospitals in 2006

- Swedish STRAMA protocol and web-based software (Skoog, ECCMID 2004)
- Demographic data, dosing and indications
- 19 pre-defined diagnosis groups
- Therapy reasons: community or hospital acquired infection or prophylaxis



# ESAC-2, HC PPS- Methods



Hospital		Department		Patient (10 fig no)			Sex		ICU Admitt.dept Ward		
							F / M		□ □ □		
Name of the drug	Dose per administration* g/mg	No of doses per day**	Route of administration Oral/ Parent /Rectal	Diagnose-group Code	Immuno-suppression*** Yes / No	Foreign material-assoc. Yes / No	Indication for therapy**** A B 1 – 5 C 1 – 3 D	Relevant culture before therapy Yes / No / Unknown	Indication for given therapy in records Yes / No	Assessment of given treatment 1-5*****	
1											
2											
3											
4											
5											

\* Combinations (Bactrim, Eusaprim and combinations against TBC) are registered in numbers of tablets or ml mixture, or ml injection fluid, respectively.

\*\* Dose every 16th h = 1,5 dose/day, every 18th h = 1,3 dose/day, each 36th h = 0,67 dose/day, each 48th h = 0,5 dose/day etc

\*\*\* Immunosuppression, e.i. chemotherapy, hematological malignancy, neutropenia, dialysis, severe hepatic failure, HIV, primary immunodeficiency, steroids for >1 month corresponding to prednisolon ≥15 mg/d.

\*\*\*\* Indication for therapy:

A. Community acquired infection

B. Hospital acquired infection: B1. Post-operative infection (infection at operation site, debut <30 days after surgery, or <1 year after surgery with implants)

B2. Other intervention related infections (e.i. CAD-, CVC- and PVC-related, VAP)

B3. C. difficile enterocolitis (debut in hospital or <30 days after discharge)

B4. Other hospital acquired infections, debut ≥48 h after admittance to hospital, ind infection from other dept., not B1-3.

B5. Infection is present at admittance from another hospital

C. Per-operative prophylaxis; prescribed for: C1. one dose C2. one day C3. >1 day

D. Medical prophylaxis (immunosuppression, HIV, prophylaxis against relapsing infections, e.i. UTI, etc)

Hospital acquired means infection associated with treatments in hospitals. B 2-4 includes infection debut during stay alt. <30 days after discharge.

\*\*\*\*\* Evaluation of given treatment:

1. Adequately directed therapy (drug, dosing, and an etiology verified by culture)

2. Adequate empirical therapy / prophylaxis (drug and dosing)

3. Obviously incorrect dosing, e.i. un-reduced dose in relation to age and renal function (creatinin clearance <50 ml/min and given full dose of drug)

4. Obviously incorrect choice of drug = ineffective therapy / prophylaxis

5. Other faults

Protocol signed by:		Registered date/sign		Patient nr:	
---------------------	--	----------------------	--	-------------	--

# ESAC-2, HC PPS- Methods



Protocol for registration of patient with ON-GOING antimicrobial treatment - Microsoft Internet Explorer

File Edit View Favorites Tools Help

Address <https://www.neotide.fi/pps/cgi-bin/scgi.exe/en/patient/>

Google Search 26 blocked ABC Check Autolink Autofill Options

## Protocol for registration of patient with ON-GOING antimicrobial treatment

Patient number: 4748

Hospital	Department	Year of birth	Sex	ICU Admitt.dept Ward	Study
	cardiology				test

Name of drug	Dose per administration	Diagnose group	Immuno-suppression	Foreign material assoc.	Indication for therapy	Relevant culture before therapy	Indication for relevant therapy in records	Assessment of given treatment
1.								
2.								
3.								
4.								
5.								

Save More rows

Back to the menu

The following treatments are registered: antiviral treatments.

**Dose per administration:** Combination

**Number of doses per day:** Dose every

**Diagnose group:** Indication for antimicrobial. For prophylaxis each antimicrobial. For prophylaxis ch

**Immunosuppression:** e.g. chemotherapy

**Indication for therapy:**

A. Community acquired infection

B. Hospital acquired infection

B1. Postoperative site, debut <30 days after surgery, or <1 year after surgery with implants)

B2. Other intervention related infections (e.g. CAD-, CVC- and PVC-related, VAP)

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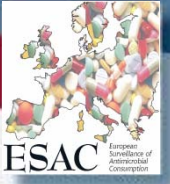
Hospital acquired means infection associated with treatments in hospitals. B 2-4 include infection debut during stay alt. <30 days after discharge.

**Evaluation of given treatment:**

Done

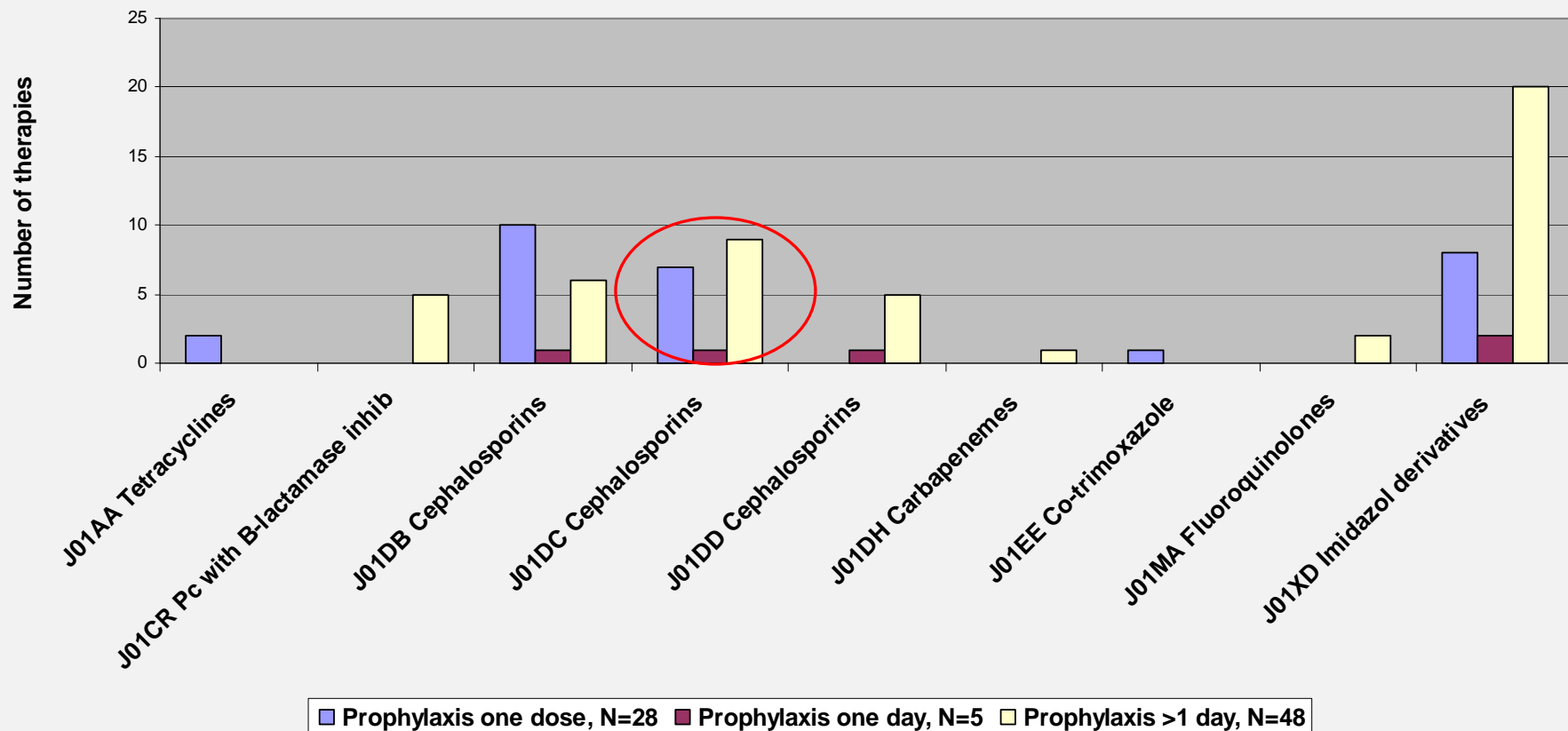
start 5 Microsoft Office O... ESAC II-HC List.xls ESACII HC Protocol V... Microsoft PowerPoint ... 401 Authorization Re... Protocol for registrati... Internet 11:19

# ESAC-2, HC PPS-Results



- 20 hospitals –
- 11,571 patients
- 3,496 patients treated with antibiotics
  - 30.2% (19.0 – 41.4) of all admitted
  - 47.5% women
  - 371 children - 10.6%

## Drugs used in peri-operative prophylaxis therapies for the lower GI-tract

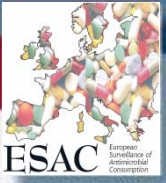


# Conclusions, Point Prevalence Survey



- **Three fold differences in antibiotic use**
- **Pneumonia is the commonest indication for treatment**
- **Peri-operative prophylaxis is too long**
- **The web-based PPS provides a tool for comparison of antibiotic prescribing in European hospitals**

# Key findings



## *A set of unique measures*

- Reveal the differences between hospitals and countries and time trends
- Provide a basement for quality indicators

## *Different denominators of clinical activity*

- Provide meaningful time trends
- Avoid comparing hospitals with differences!

## *LS plus PPS*

- More precise view

*>>> Still to go for quality indicators....*



# ESAC-3 Hospital Care Subproject: 2007-2010

Ansari F, Davey P, Dundee, UK

Tom Bajek, Ljubliana, Slovenia

Drapier N, Muller A, Goossens H, Antwerp, Belgium

The ESAC III hospital care study group



# ESAC-3 Hospital Care Subproject 2007-2010



- To establish a European network for surveillance of antibiotic use.
- To organise a European wide longitudinal (together with EARSS) and point prevalence survey.
- To develop quality indicators of antimicrobial consumption in the hospital care sector (together with ABS).
- To improve antibiotic prescribing in Europe in specific patient groups (e.g. surgical prophylaxis, together with ESGAP)
- To use a set of clinical activity denominators for longitudinal studies to validate time trends in hospital antibiotic use.

# ESAC-3 Hospital Care Subproject 2007-2010



## Components PPS

- Develop webPPS Q1-2008, University of Antwerp
- PPS-1 (at least 2 hospitals/country) Q2-2008
- Roll out PPS methodology Q4-2008
- European wide PPS Q2-2009
- Release of PPS-2 results released at Annual AMR Day (November 18, 2009)

**Invitation to AS MANY  
Belgian hospitals as  
possible to participate in  
PPS Q2-2009**

**Target date: November  
18, 2009 (Annual  
European AMR Day)**