

CARE-ICU - Controlling Antibiotic REsistance in Intensive Care Units

- First report from a web based program for improved infection control in European ICUs

Håkan Hanberger^{1,2}, Dilek Arman³, Hans Gill⁴, Vlastimil Jindrák⁵

Smilja Kalenic⁶, Andrea Kurcz⁷, Monica Licker⁸, Paul Naaber⁹, Elizabeth A Scicluna¹⁰, Václav Vanis⁵, Sten M Walther¹ and Care ICU study group, IPSE

¹Faculty of Health Sciences, Linköpings Universitet, Sweden, ²Swedish Institute for Infectious Disease Control, Solna, Sweden, ³Dept. Clin. Microbiol. and Infect. Dis., Gazi University School of Medicine, Ankara, Turkey, ⁴Department of Biomedical Engineering, Linköpings Universitet, Linköping, Sweden, ⁵Dept. of Clin. Microbiol. Antibiotic Centre, Na Homolce Hospital, Praha, Czech Republic, ⁶Reference Centre for Hospital Infections Clinical Hospital Centre Zagreb, Croatia, ⁷National Center for Epidemiologia, Budapest, Hungary, ⁸Microbiology Department, University of Medicine and Pharmacy, Timisoara, Romania, ⁹Dept Clinical Microbiology, United Laboratories, Tartu University Clinics, Tartu, Estonia, ¹⁰Infection Control Unit, Mater Dei Hospital, Msida, Malta

Objective

To report results from a European ICU surveillance programme with the aim to generate data for bechmarking and audit of microbial resistance, infection control procedures and antibiotic consumption.

Design

Observational study.

Interventions

None.

Settings, patients and participants

Thirty-four ICUs¹ from eight countries participated in the large pilot, which took place during 2005. The median annual number of admissions to ICU was 551 and the median summated length of stay was 2 595 days.

Measurements and results

Antibiotic consumption varied widely from 348 to 4 992 Defined Daily Dosages per 1 000 occupied bed days (DDD₁₀₀₀), with a median of 1 254 DDD₁₀₀₀. A median of 11.6% (range 0-100%) of *S. aureus*² were MRSA and the corresponding figures for ESBL phenotype (resistance to 3rd generation cephalosporins) of *E. coli*² and *K. pneumoniae*² were 3.9% (0-80%) and 14.3% (0-77.8%) respectively. Antibiotics to which more than 90% of isolates of a species were susceptible were defined as treatment alternatives (TA₉₀). More than half of all ICUs had no or only one TA₉₀ for *P. aeruginosa*². Screening for alert microorganisms in patients at risk on admission was commonly omitted, and there was a lack of single rooms for isolation and cohort nursing care of patients colonised or infected with alert organisms.

1 The ICU short names consist of the 2 character Internet top level domain name (Cz=Czech Republic, Ee=Estonia, Hr=Croatia, Hu=Hungary, Mt=Malta, Ro=Romania, Se=Sweden, Tr=Turkey) followed by 2 characters for the type of ICU (Me=Medical, Mx=Mixed, Ne=Neonatal, Ns=Neurosurgical, Ot=Other, Su=Surgical, Th=Cardiothoracic) and a sequence number.
2. Susceptibility was based on data for at least 5 isolates.

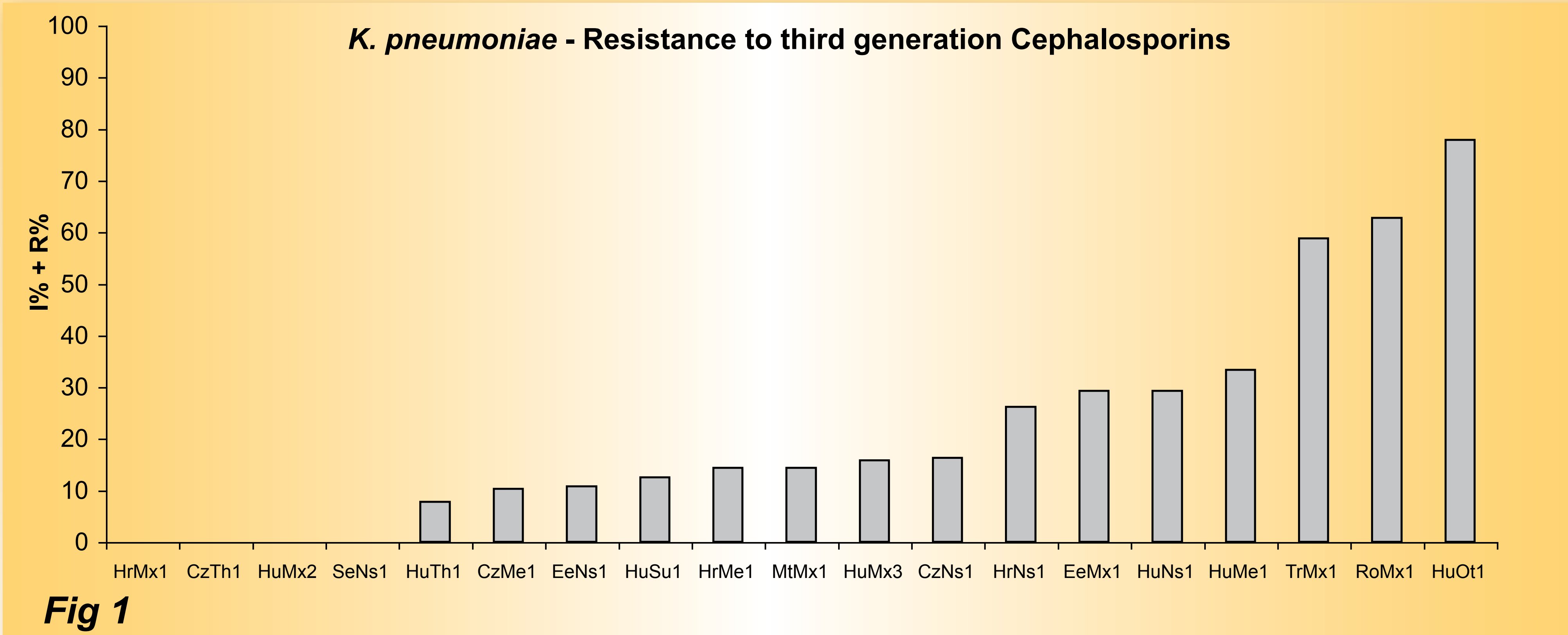


Fig 1

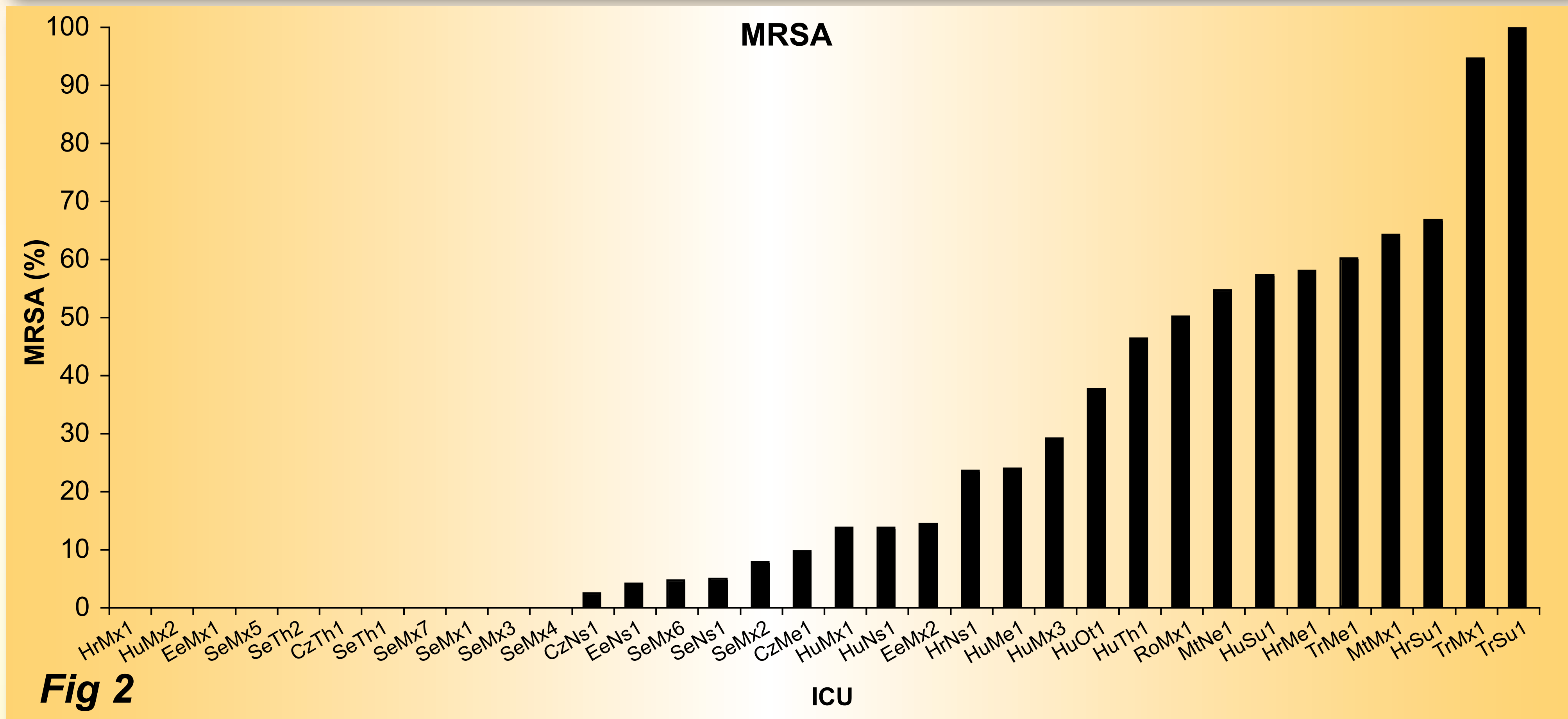


Fig 2

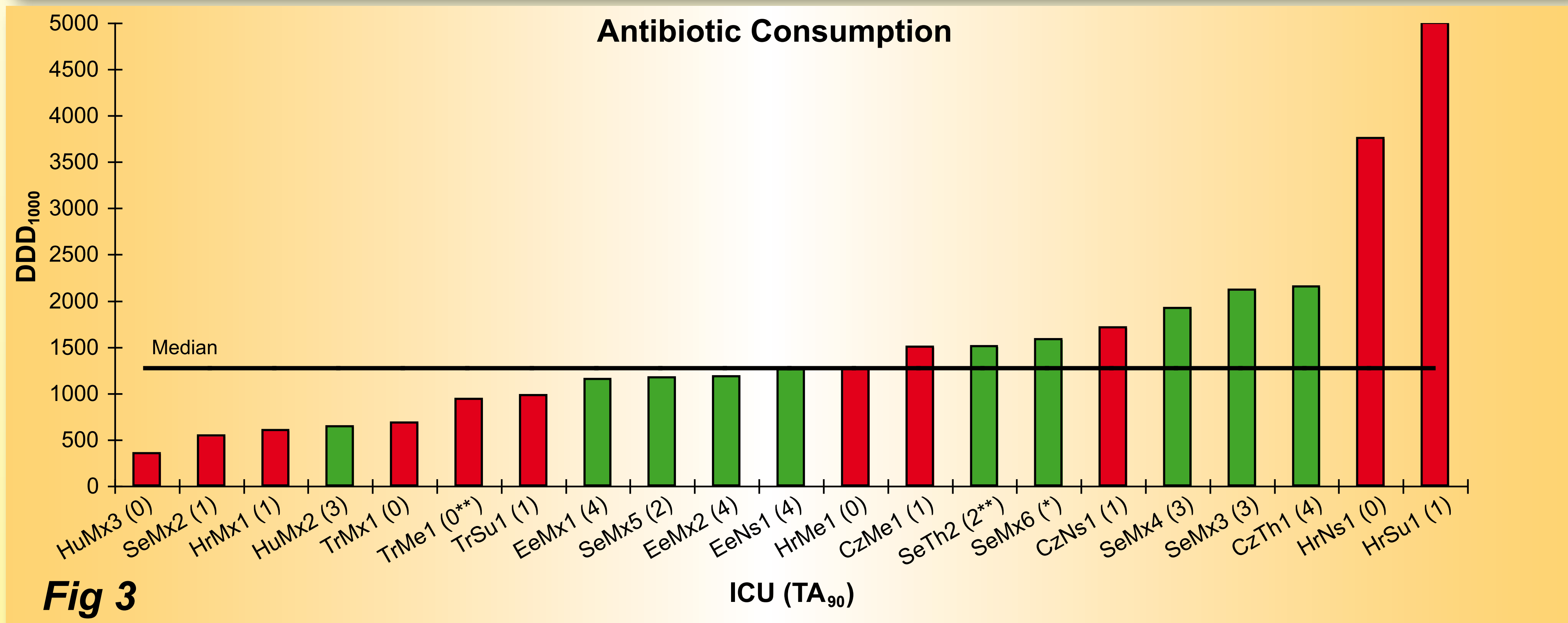


Fig 3

Fig 3. ≤ 1 TA₉₀ including aminoglycoside, ceftazidime, ciprofloxacin and carbapenem against *P. aeruginosa* are denoted with red bars and TA₉₀>1 are denoted with green bars. Susceptibility was based on data for at least 5 isolates except for one ICU where no *P. aeruginosa* isolates were found (*) and two ICUs with < 5 isolates (**).

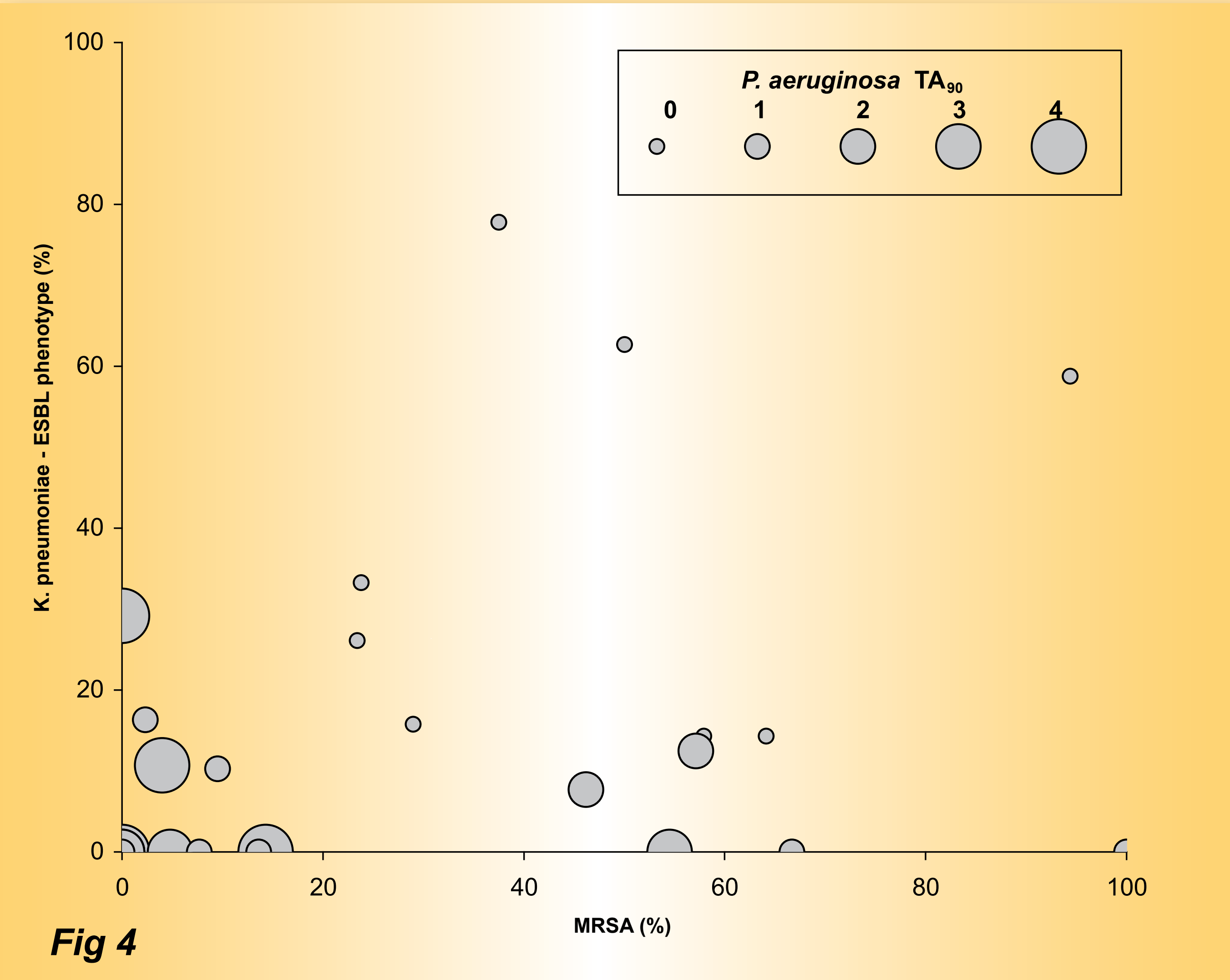


Fig 4

Fig 4. Frequency of MRSA, resistance to third generation cephalosporin in *K. pneumoniae* (ESBL phenotype) and TA₉₀ in *P. aeruginosa* for each ICU. The bubble size indicates the number of TA₉₀ for *P. aeruginosa* for each ICU.

Conclusion

The surveillance programme provided fast and easy access to results for benchmarking and comparative audit of antibiotic policy and infection control in participating ICUs.

This data can help to identify and establish interventions in infection control and antibiotic policy tailored for each individual participating ICU.

Care ICU is a part of the project *Improving Patient Safety in Europe* (IPSE) funded by the European Commission Directorate General for Health and Consumer Protection (DG SANCO).

Contact: hakha@imk.liu.se, +46-705-797102