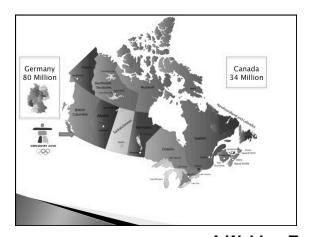


My thanks to Dr Linda Hoang and Dr Michael Gardam

Outline IPAC in Canada Recent Trends in Antibiotic Resistant Organisms in Canada Changing Approaches Emerging Organisms





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IPAC - Canada

IPAC Canada is a national, multidisciplinary association committed to the wellness and safety of Canadians by promoting best practice in infection prevention and control through education, standards, advocacy and consumer awareness.



IPAC Canada's Roles

- initiating and coordinating effective communication and cooperation among all disciplines united by infection control activities
- supporting and/or developing effective and rational infection control practices
- standardizing infection control practices
- promoting research in areas related to infection control
- promoting and facilitating infection control education for both infection control practitioners and other personnel working in hospitals, nursing homes and related institutions.

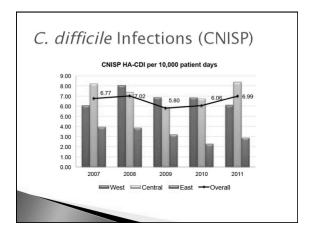
Public Health Agency of Canada

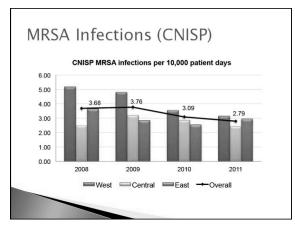
- National Surveillance (CNISP)
 - Surveillance data submitted from 52 large hospitals from across Canada
- Centre for Communicable Disease and Infection Control -
- · National IPAC Guidelines

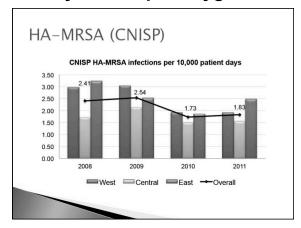
Provincial Bodies

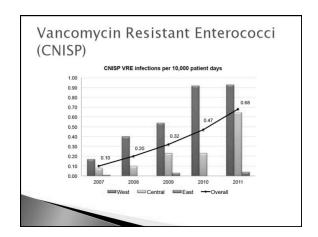
- Public Health Ontario (PIDAC/RICN)
- Manitoba Department of Health
- Alberta Health Services
- BC Provincial Infection Control Network
- Infection Control Nova Scotia
- Infection Control Newfoundland and Labrador
- Each Province produces their own guidance documents!

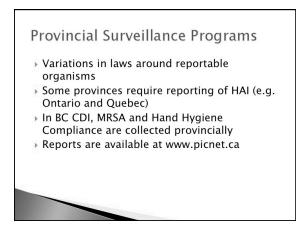
Recent Trends in ARO in Canada

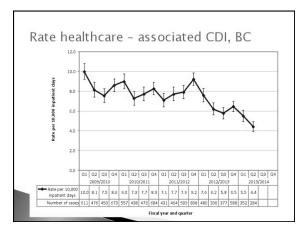


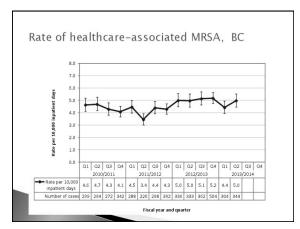




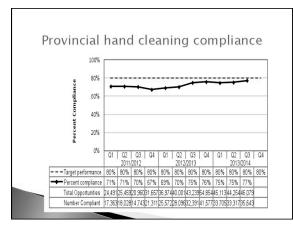








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New Approaches

- > Changes to approach to VRE in Ontario and BC
- Evaluation of changes

Early concerns about VRE

- Emerging organism
- Highly drug resistant
- Poor therapeutic options
- Resistance gene transfer to more pathogenic organisms i.e. MRSA

VRE

- Enterococci in general are intrinsically drug resistant
- Do not produce toxins
- Frequently found with other organisms
- Infections typically occur in patients with considerable pre-existing morbidity
- High mortality rate but contribution of vancomycin resistance is not clear

Typical VRE infections in adults

- Urinary tract
- · infection vs. colonization
- Often catheter-associated
- Bacteremia >> endocarditis
- · Often catheter associated
- Frequently polymicrobial (GI source etc.)
- Deep tissue infections and wounds
- Usually polymicrobial

Very rarely caused by VRE

- Respiratory tract infections
- Meningitis
- > Skin, soft tissue infections

Specific versus non-specific control measures

- Specific control measures
- · Admission screening
- · Contact isolation, periodic screening
- Non-specific control measures
 - Hand hygiene
- · Environmental cleaning
- Practice bundles
- · Antimicrobial stewardship

Does VRE increase length of stay?

- Yes, but why?
- Marker for severe morbidity
- Small percentage of patients develop significant infections
- · Infection control measures themselves
- · increase LOS
- · Isolation
- · Slowing of patient transfers

Decision

- Stop specific VRE control measures
- 2010 Vancouver Island Health screening only patients admitted to designated units.
 - Patients in other patient care areas are not placed on additional precautions unless they have diarrhea
- 2012 4 Hospitals in Ontario stopped screening for VRE, isolating cases or reporting outbreaks
- 2013 2 other health authorities adopted similar policies
- Prospectively collect data to inform future practice.

Evaluation

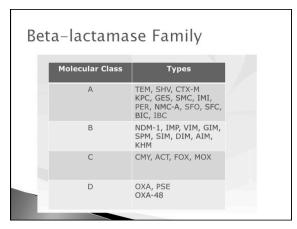
- Results Pending...
- Negative clinical impact has been negligible
- · Substantial decrease in isolation costs
- Increased ability to focus on other organisms

Emerging Organisms

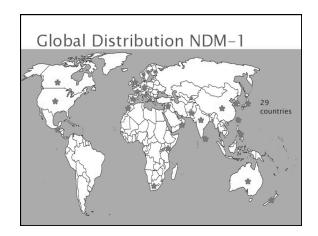
СРО

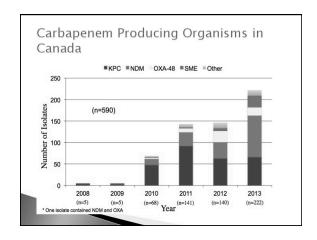
What are Carbapenemase Producing Organisms (CPO)?

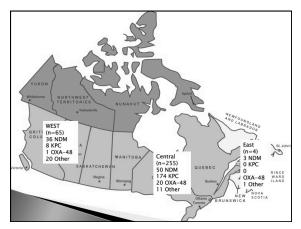
- Carbapenemases are a class of enzymes that inactivate carbapenem antibiotics by hydrolysing them.
- Carbapenem antibiotics, often referred to as "last resort antibiotics":
 - Imipenem
 - Meropenem
 - Ertapenem
- Carbapenemases most commonly in E. coli and Klebsiella spp., (Enterobacterieaceae) but have also been found in other Gram-negative species.

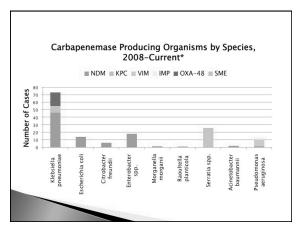


| | Delhi Metallo-beta- amase (NDM-1) |
|--------------------------|--|
| | rts in 2008 of Swedish and UK travelers to Indian ontinent |
| | then, reports of high endemicity in Indian, Pakistan angladesh hospitals |
| cities • 51/ • 2/5 | -1 genes in sewage and water reservoirs in some Indian 171 (30%) waste water seepage 0 (4%) communal drinking water Iples |
| Walsh et al | The Lancet Infectious Diseases, 2011, 11: 355-62 |
| | |

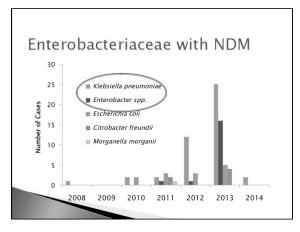


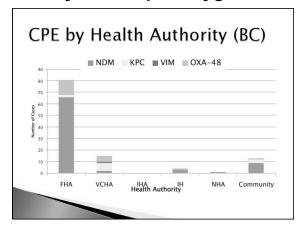




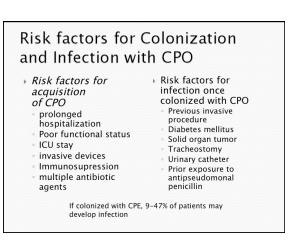


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How are these organisms transmitted? 1. Patient-to-patient 2. Shared Health Care equipment 3. Environmental Contact (environmental reservoirs) 4. Health care workers (Primarily hands)



CPO Measures Implemented Screening/Active surveillance On admission to Unit Weekly point prevalence All contacts of suspect or confirmed cases, at 0, 7 and 21 days Precautions Private room and staff cohorting and dedicated equipment Cohorting of patients and staff "CPO" nursing assignments & dedicated ward Hand hygiene & PPE (goal: 100%) Weekly audits Antimicrobial stewardship

Infection Control Processes

- Screening for all admitted patients
- Question: "Have you been hospitalized or had renal dialysis outside of Canada anytime in the previous 6 months?"
- If yes: patient will have rectal screen for CPO

Next Steps and Challenges

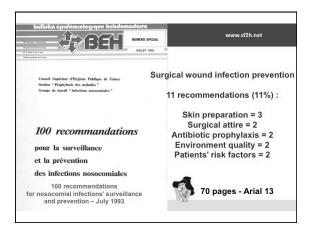
- Better and faster testing
- Develop Real-time PCR method for screening specimens directly
- Maintain aggressive infection control state & CPO alerts between facilities
- Continued Provincial level surveillance with infection control data
 - Collaboration with PICNet
- Further explore genomic characteristics of BC strains and transmission behaviour
 - · Whole Genome Sequencing

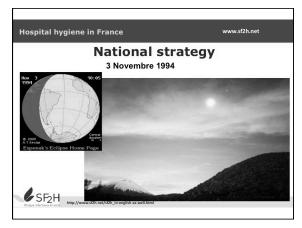
Summary

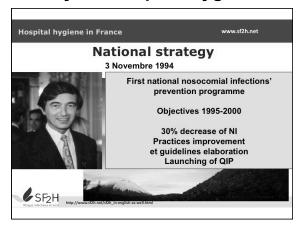
- CPO are an emerging pathogen with global spread, now in Canada
- CPE can spread within institutions
- The most vulnerable patients are the most at risk to become colonized and infected
- Treatment of infections is complex
- Control of spread requires full compliance with precautions and antibiotic stewardship

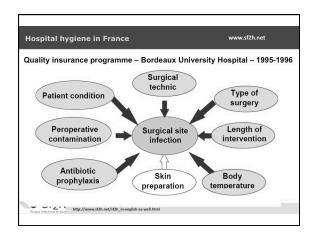


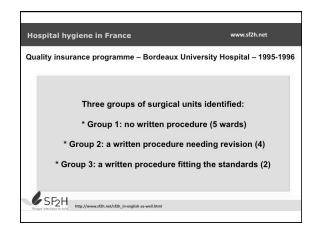


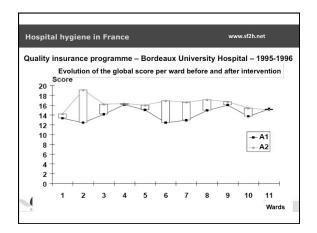


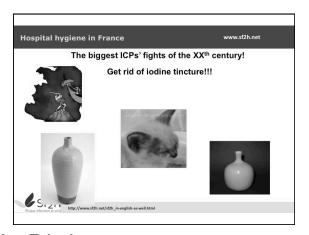




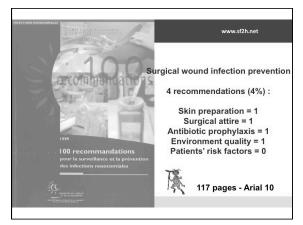


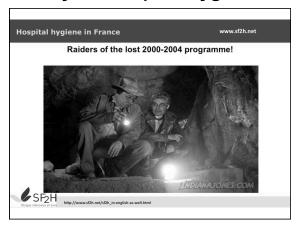


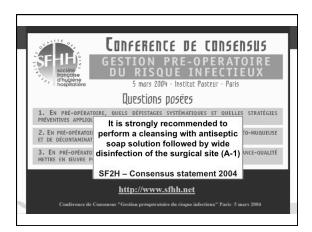


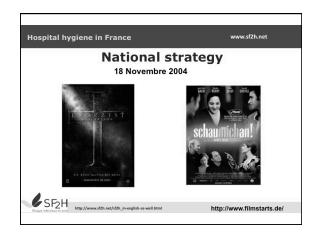


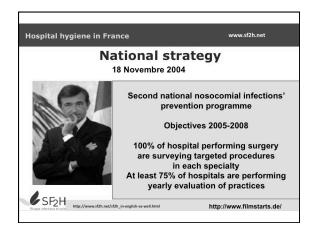
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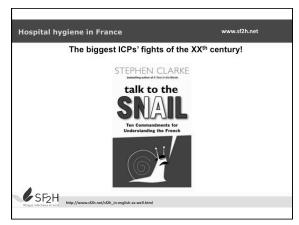


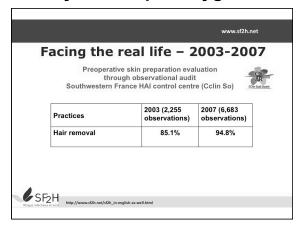


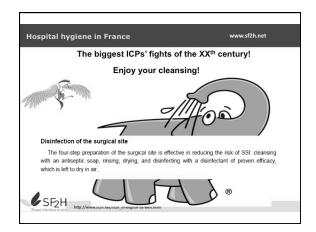


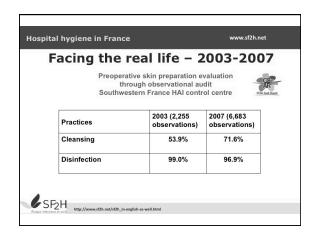


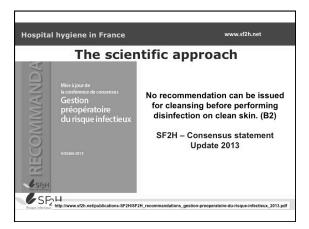
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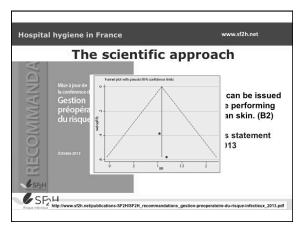




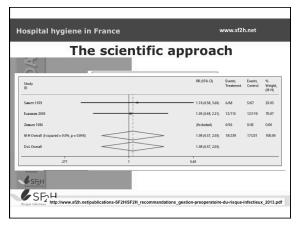


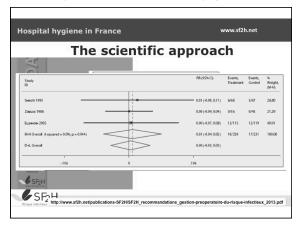


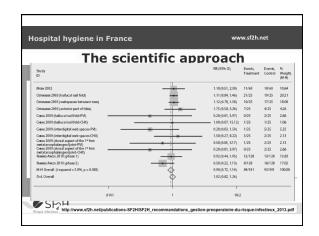


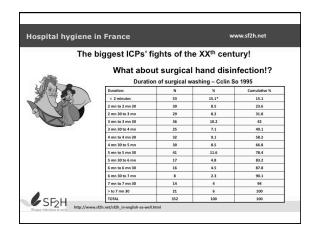


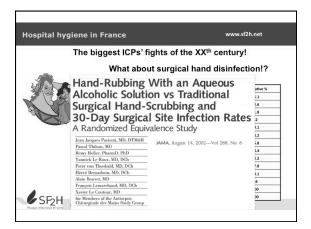
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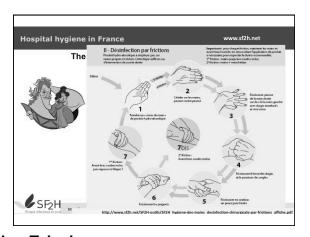




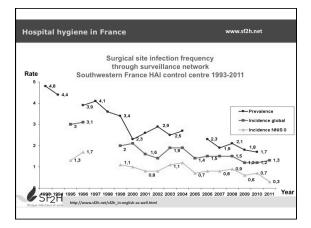


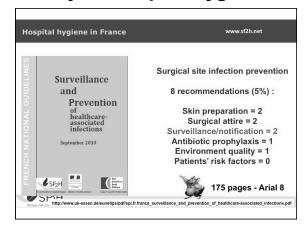


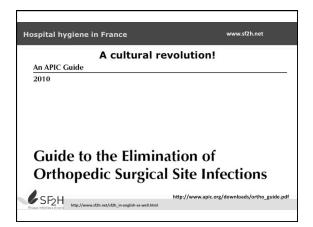




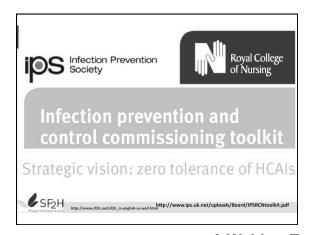
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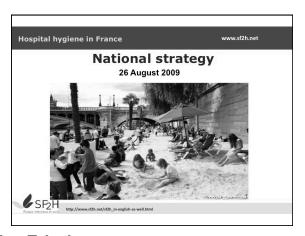




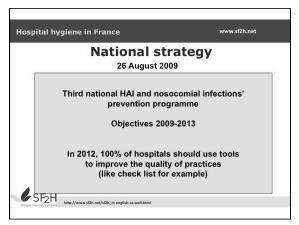


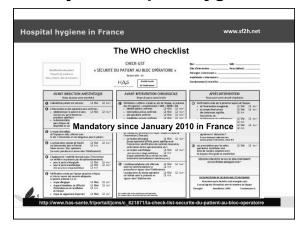


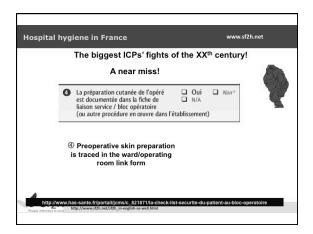


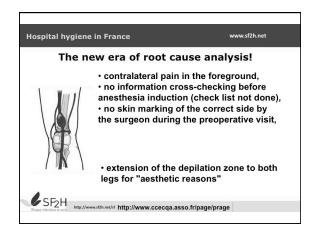


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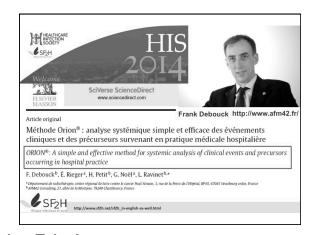




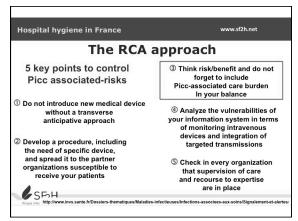




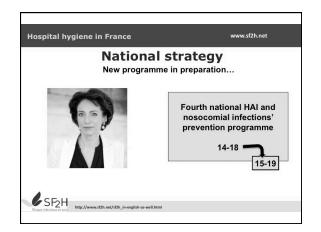


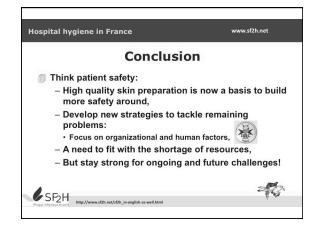


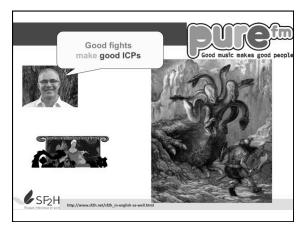
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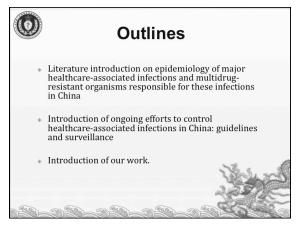


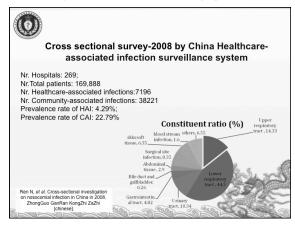


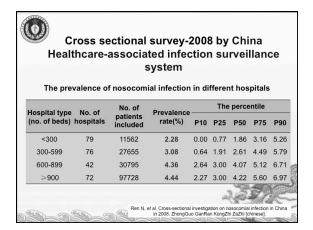


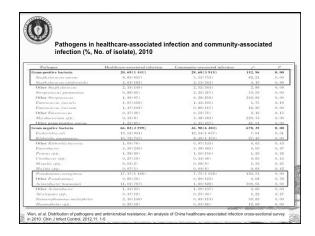


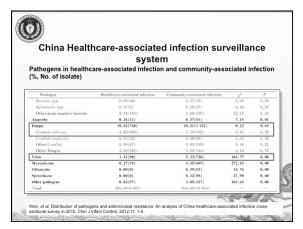
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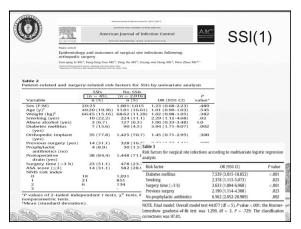




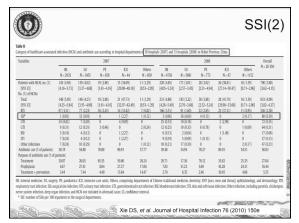


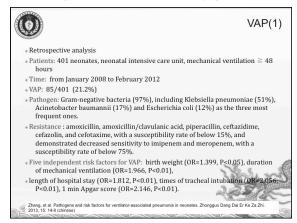


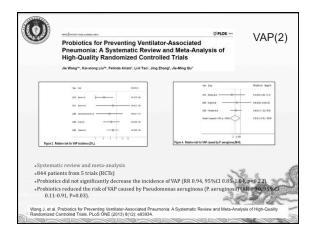


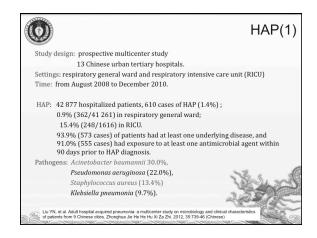


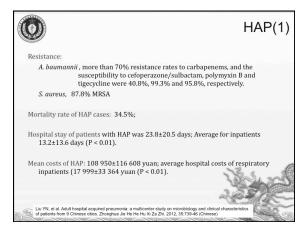
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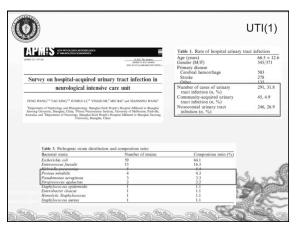




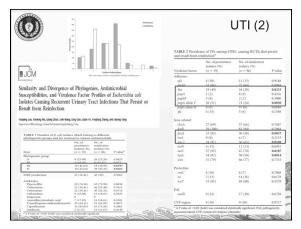


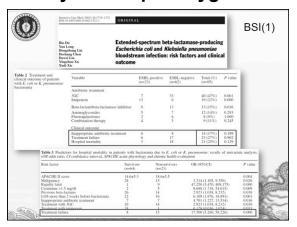


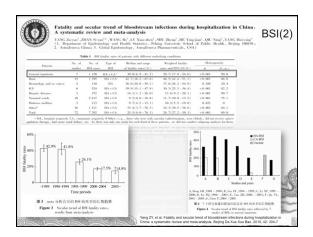


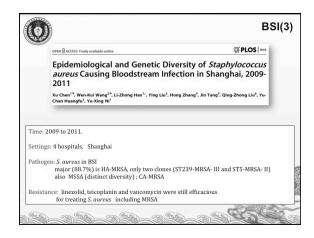


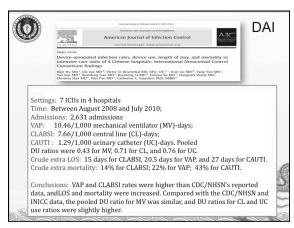
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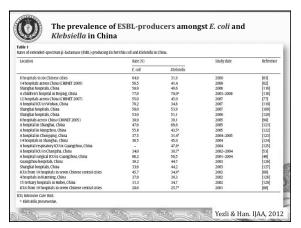




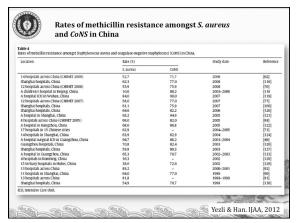


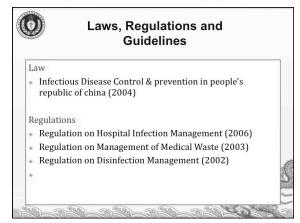




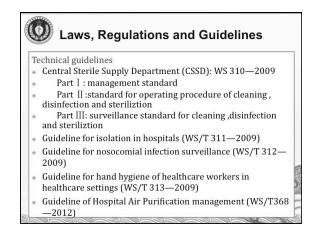


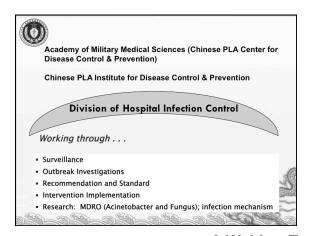
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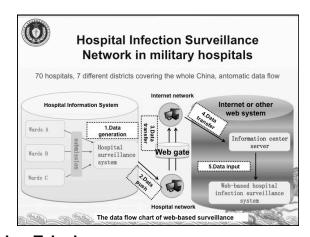




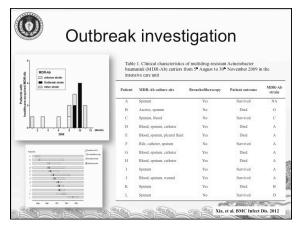


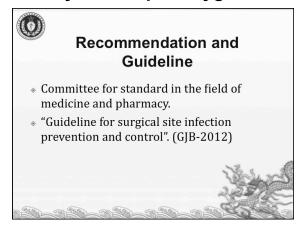


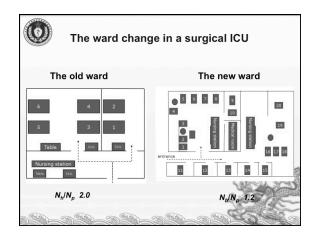


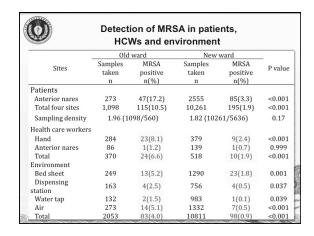


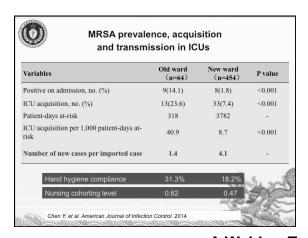
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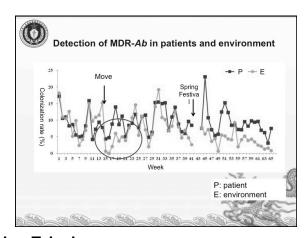




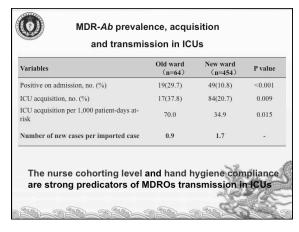


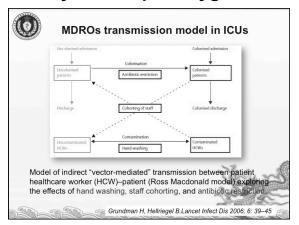


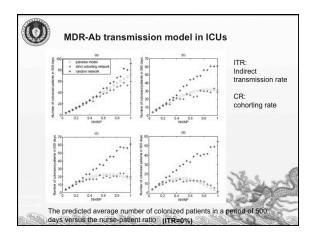


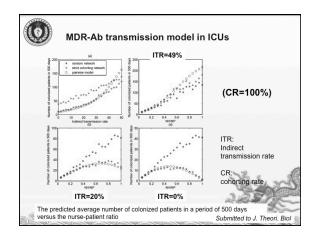


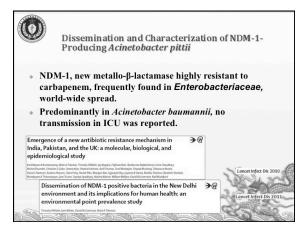
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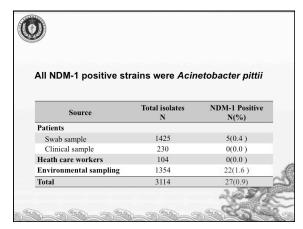




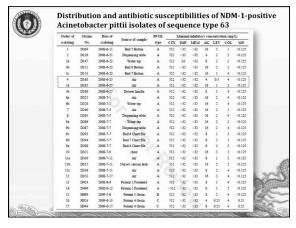


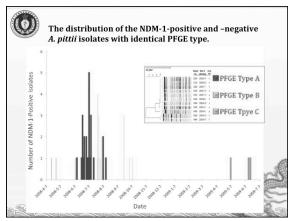


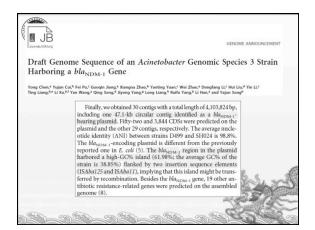


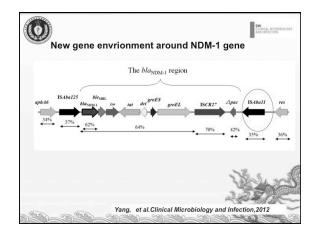


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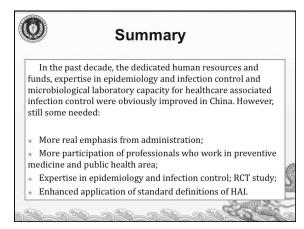








| E Table, P | nterob | acter | iac | eae, in I | ns, Fam Hospital | , B | | | | | | I, Beijir | ıg. |
|----------------------------|-------------------------------|----------------------------------|-----------------------|---|--|------------|----------|----------|----------|----------|-----|-----------|-------|
| Isolate no. | Organism | Source | ST† | bla _{NOM-1} plasmid size, type | Prevalence of RDs | MICs, mg/L | | | | | | | |
| | | | | | | CTX | | TZP | | MEM | | AK | LVX |
| IR5028 | Escherichia coli | Urine | 167 | ≈190 kb, A/C | blatem. blacks.1.‡ blactx*group 1, 2, 9- aac(6')-lb-cr,‡ armA± | >256 | >256 | >256 | >32 | >32 | >32 | >256 | >32.0 |
| IR5047 | Klebsiella pneumoniae | Blood | 1,240 | ≈50 kb, X3 | blackut, qnrS,‡ | >256 | 64 | >256 | 8 | 6 | >32 | 2 | 0.38 |
| IR5337 | Providencia rettgeri | Urine | NA‡ | ≈190 kb, A/C | blaser, blacky, mtC‡ | | >256 | - 200 | >32 | >32 | 4 | >256 | >32.0 |
| R5338 | Enterobacter cloacae | Urine | NA | ≈50 kb, X3 | bla _{SHI} ,‡ bla _{OXA-1} , bla _{CTX-group 2} | >256 | >256 | >256 | >32 | >32 | >32 | 1.5 | 2.0 |
| IR5343 | Raoultella ornithinolytica | Abscess | NA | ≈70 kb, N | qnrS‡ | >256 | 32 | >256 | >32 | >32 | >32 | 2 | 0.75 |
| piperacillis †The allel | n-tazobactam; IMP | , imipenem; fr nulti-locus ST | MEM, me loci for a | ropenem; ETP, ertag given isolate are co | pe; RDs, resistance of enem; AK, amikacin; mbined into an allelic | LVX. lev | ofloxaci | n; NA, r | not appl | licable. | | e; TZP, | |



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