**FIDSSA 2017 Congress Abstracts
Oral Presentations**

**ID: 8435**

**Category:** Infection Control (ICSSA)

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**ABSTRACT TITLE:** IMPLEMENTATION OF A MOBILE APPLICATION FOR HAND HYGIENE MONITORING: SMART, SIMPLE AND INTERACTIVE

**Introduction**

Hand Hygiene (HH) is globally regarded as the single most effective infection prevention intervention used to reduce the spread of micro-organisms in healthcare settings. However, healthcare workers’ compliance to HH is estimated at < 50%. Comprehensive HH programmes are thought to include multi-modal strategies with focus on core elements such as: infrastructure, knowledge, awareness, measurement, feedback and behaviour change. This study aimed to implement a novel mechanism of HH monitoring across a network of 56 private South African hospitals.

**Methods**

A mobile application (app) was developed as a platform for data collection of HH observations according to the World Health Organisation’s five moments campaign. An additional measurement of ‘bare below the elbows (BBE)’ was included. Access to the app was granted to identified observers following completion of a training module and competency assessment. Observations were required to be recorded weekly and a compliance dashboard was designed and distributed weekly to enable regular feedback to front-line staff.

**Results**

A total of 259 283 observations for HH were recorded during the 47-week study period of which 214 869 were compliant (82.9%). Evaluation of HH compliance per moment indicates: before patient contact 81.7% (n = 63 486); before aseptic task 91.8% (n = 17 284); after body fluid exposure risk 93.0% (n = 19 175); after patient contact 84.8% (n = 55 355) and; after contact with patient’s surroundings 77.6% (n = 59 569). Average BBE compliance was 89.6% (n = 131 725). A significant improvement in overall HH compliance was noted comparing data of 2016 to 2017 (p = < 0.05). Compliance of all categories of staff has improved over time 2016 vs 2017. Improvements from doctors (72.3% vs 75.3%), registered nurses (90.4% vs 91.5%) and pharmacy staff (73.4% vs 79.2%) were significant (p < 0.05).

**Conclusion**

HH compliance of healthcare workers in this study was markedly higher than that of previous studies. The implementation of the mobile application made data collection quick, simple and allowed for the availability of real time data to front-line staff to drive improvement.

**ID: 8680**

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ABSTRACT TITLE: ODYSSEAN MALARIA OUTBREAK AT A BUSH LODGE IN MADIKWE GAME RESERVE, NORTH WEST PROVINCE, OCTOBER–NOVEMBER 2015

Introduction
In November 2015, the National Institute for Communicable Diseases (NICD) was notified of a malaria case at a lodge in Madikwe Game Reserve, North West Province – usually a non-transmission area in South Africa. A few other people with fever, headache and flu-like symptoms at the same establishment were reported. An investigation to determine the possible cause/source of the illnesses was undertaken.

Methods
A structured questionnaire was used to gather information on demographic, clinical and exposure history. Interviews were conducted with employees and managers; blood samples were collected. Environmental assessment of the residence and immediate surroundings was conducted. Blood smear microscopy and PCR analysis for the detection of malaria parasites were done.

Result
Four laboratory-confirmed malaria cases were identified. All were female, employed, and resided at the lodge staff residences. Dates of symptom onset were 17 October 2015 (case 1), 21 October (case 2), 22 October (cases 3, 4). Fever was reported in all; headache, dizziness and painful joints in three. Two were admitted to hospital. No recent blood transfusions were reported. None of the cases had travelled to malaria-endemic areas. However, travel history to possible malaria transmission areas was reported in eight of the 33 staff members interviewed. Cases 1 and 2 occupied adjoining rooms, cases 3 and 4 shared a room. Staff residences were within 50–60 m of the parking bay for establishment vehicles, including those returning from malaria-endemic areas. There was no evidence of free-standing water that could enable mosquito breeding. No asymptomatic infections were identified: all laboratory investigations for malaria were negative.

Conclusion
It is most likely that the cases acquired malaria from an imported, infected mosquito – the phenomenon of odyssean malaria. Spraying vehicles with a pyrethroid insecticide with residual effect would reduce the risk of transporting vector mosquitoes but this would be difficult to implement.

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ABSTRACT TITLE: ASSESSING THE IMPACT OF ANTIMICROBIAL STEWARDSHIP WARD ROUNDS USING AN APP-BASED DATA COLLECTION TOOL IN TWO PUBLIC HOSPITALS IN THE EASTERN CAPE, SOUTH AFRICA

Frere Hospital is a 900-bed tertiary/speciality public hospital in the Eastern Cape, South Africa. The burden of disease consists of chronic conditions, acute and chronic infections with a major impact of HIV and TB.

The main challenges in our institution are a lack of resources (staff and finances) but also a lack of awareness of the importance of rational antimicrobial use. As a clinical pharmacist at Frere Hospital, I attend the Antimicrobial Stewardship rounds weekly with the ID Specialist and team which commenced in 2015 and we recently looked at an app as a tool to assess the impact of antimicrobial stewardship ward rounds. Attendance at the conference would be a fantastic opportunity to learn from other hospitals and pharmacists across Southern Africa and share our skills.

As mentioned above, one of the challenges in the institution is finances and we do not have a pharmacy budget for such.

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ABSTRACT TITLE: PROCALCITONIN-GUIDED ANTIBIOTIC THERAPY FOR SUSPECTED AND CONFIRMED SEPSIS OF PATIENTS IN A SURGICAL-TRAUMA INTENSIVE CARE UNIT

Introduction
Procalcitonin (PCT) is a useful sepsis marker to guide duration of antimicrobial therapy. PCT-guided algorithms have demonstrated value in reducing duration of antibiotic therapy in critical care patients. There is a paucity of data supporting the use of PCT-guided
antibiotic algorithms in trauma patients and in patients from developing countries.

**Methods**

A prospective study was conducted in the surgical trauma intensive care unit (ICU) at Charlotte Maxeke Johannesburg Academic Hospital in April 2014 to July 2015 in a two period cross-over design. Patients with suspected or confirmed sepsis were recruited consecutively in two periods of almost equal length. In the first period, 40 patients were recruited as controls and antibiotics were discontinued as per standard of care. In the second period, 40 patients were recruited into the intervention group and antibiotics were discontinued if the PCT decreased by ≥ 80% from the peak PCT level, or to an absolute value of less than 0.5 µg/L. Antibiotic duration was the primary outcome. Patients were followed up for 28 days from the first sepsis event.

**Results**

For the first sepsis event the intervention group had a mean antibiotic duration of 9.3 days while the control group had a mean duration of 10.9 days (p = 0.10). The mean duration of treatment was 12.0 days for a second episode of sepsis in the control group and 9.6 days in the intervention group (p = 0.09). Clinician compliance to the PCT algorithm was 62.5%. The intervention group had more antibiotic-free days (7.8 days) compared to the control group (3.9 days) (p = 0.004). The length of ICU stay and length of hospital stay for the two groups were similar. The in-hospital mortality was reduced in the intervention group (15%) compared to the control group (30%).

**Conclusion**

Our data supports the use of PCT-guided algorithms for antibiotic stewardship in surgical trauma patients. Clinician compliance would likely increase the benefits observed in our study.

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**ABSTRACT TITLE:** INDEPENDENT RISK FACTORS ASSOCIATED WITH CANDIDA AURIS CANDIDAEMIA IN SOUTH AFRICA – AN ANALYSIS OF NATIONAL SURVEILLANCE DATA, 2016–2017

**Introduction**

*Candida auris* is a globally-emerging, multi-drug-resistant invasive fungal pathogen. We aimed to determine risk factors for *C. auris* candidaemia to inform prevention and management.

**Methods**

Culture-confirmed cases of candidaemia were detected through active, national laboratory-based surveillance at all public- and private-sector hospitals, January 2016 – June 2017. We defined a case of candidaemia as any patient with *Candida* cultured from blood. Fungal isolates were identified using Bruker mass spectrometry or ITS region sequencing. We used multivariable logistic regression to assess clinical/demographic factors associated with *C. auris* candidaemia versus other *Candida* species from patients admitted to 25 enhanced surveillance sites.

**Results**

Among 2044 cases of candidaemia, 206 (10%) were caused by *C. auris*, with a median age of 54 years (IQR: 36–64) versus 26 years (IQR: 0–57) among non-*C. auris* cases. Most cases in both groups were male (*C. auris*: 60% [122/203]; non-*C. auris*: 53% [960/1806]). *C. auris* cases predominantly occurred in Gauteng Province (191/205, 93%) and in private-sector hospitals (154/206, 75%). Among 50 *C. auris* cases with clinical data, crude in-hospital mortality was 49% (23/47), versus 42% (333/788) for non-*C. auris* cases (p = 0.368). Both groups had high proportions with invasive devices and intensive care unit admission. Twenty-nine per cent (12/41) and 85% (35/41) of *C. auris* cases had received prior antifungal/ antibiotic therapy. Being admitted to a private-sector hospital increased the odds of *C. auris* candidaemia three-fold (aOR 3.6; 95% CI 1.62-7.77). Other risk factors included older age (aOR 1.01 for every year; 95% CI: 1.01–1.03) and longer hospital stay before first positive blood culture (aOR 1.01 for every day admitted; 95% CI: 1.01–1.03).

**Conclusion**

Older patients with prolonged hospitalisation and admission to private-sector facilities had increased odds of *C. auris* candidaemia, possibly owing to exposure to multiple invasive devices and prior antimicrobial therapy.
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ABSTRACT TITLE: RISK ASSESSMENT FOR THE TRANSMISSION OF INFECTIOUS DISEASES IN THE SOUTH AFRICAN POLICE SERVICES (SAPS)

Introduction
In 2012, the Constitutional Court made a landmark judgment against the Minister of Correctional Services where it was found that the Department of Correctional Services (DCS) failed to take reasonable steps to prevent transmission of tuberculosis (TB). Infection prevention and control (IPC) is a collective term for activities, policies and procedures designed to prevent and control the transmission of infectious diseases within various environments (e.g. healthcare centres, detention and correctional centres, other institutions and the community).

The continuum of an offender’s journey starts at the crime scene and arrest, transportation between the police station (SAPS) holding cells and the courts until admission to a correctional centre. Infection transmission risks occur at each stage of the continuum and stringent IPC measures are critical to prevent infection spread.

Methods
An IPC audit tool was developed for SAPS. Five assessments were done. The assessment included process mapping, a questionnaire, observations, interviews and measurement of ventilation in SAPS cell blocks. Common IPC non-conformances were identified.

Results
- Ventilation issues were identified – variation between 25 L/s per person to 50 L/s/per person which is insufficient for a medium-risk population.
- Aggregation and overcrowding are significant risks for TB transmission.
- Improvement opportunities included open window enforcement, medical screening, separation protocols, hand hygiene, environmental hygiene, medication management and occupational health and safety.
- IPC procurement under the current supply chain system is problematic.

Conclusion
The risk assessment highlighted the need for the implementation of an IPC programme in the SAPS. A Provincial SAPS Committee-approved training programme was commenced in June 2017.

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ABSTRACT TITLE: RABIES VIRUS DOWN-REGULATES THE EXPRESSION OF ACTIN AND MICROTUBULE-ASSOCIATED PROTEINS IN NEURONS

Background
Rabies virus (RABV) causes neuronal dysfunction, and alters the structural morphology of dendritic spines by inducing changes in cytoskeleton which is further comprised of actin, microtubule and intermediate filaments. For microtubule, EB3 is a microtubule plus-end binding protein that stabilises microfilaments and belongs to the third family member of RP/EB group. The p140cap is an interacting partner of EB3 that, together with EB3, plays a vital role in postsynaptic density, dendritic spine functions and morphology.

Methods
The gene expression and respective protein contents of EB3 and p140cap were compared in neuronal cells under the effect of fixed and street strain of RABV. Furthermore, gene expression levels of important actin binding, microtubule and synapse-related proteins were also studied. In this study, immunofluorescence, western blot and real time PCR were carried out to compare the gene expression and corresponding protein contents of different cytoskeleton related proteins.

Results
Both strains of RABV significantly reduced the gene expression and protein contents of essential cytoskeleton-related proteins like EB3 and p140cap. However, the street strain considerably inhibited the transcription level of p140cap, but had no significant effect on its protein level. The fixed strain produced fractured microtubules in fixed neurons, and down-regulated different microtubule and actin-associated proteins, while up-regulated the level of Tesk2 gene. In neuronal cells, the fluorescence localisation of EB3 protein was random and varied at 48 hour and 98 hours of post-infection.

Conclusions
Rabies virus possibly alters the neuronal structures by changing the gene expression levels of integral protein binding partners of microtubules and actin. Most importantly, EB3 and p140cap are vital in maintaining the morphology of dendritic spines. These results could help us to understand the intricate relationship between RABV and microfilaments.

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ABSTRACT TITLE: MOLECULAR CHARACTERISATION OF SALMONELLA TYPHI ISOLATES FROM SOUTH AFRICA, 2012–2014

Introduction
Typhoid fever is a grave systemic infection caused by the bacterium Salmonella enterica serovar Typhi (Salmonella Typhi). It has been estimated that Salmonella Typhi causes approximately 720 illnesses per 100 000 populations annually, in Africa. Of more concern is the emergence and spread of the Salmonella Typhi H58 haplotype from Asia. Salmonella Typhi H58 is highly clonal and has been associated with multi-drug resistance to first-line antimicrobials and reduced susceptibility to fluoroquinolones, the current treatment regimen for patients with typhoid fever. The aim of this study was to use molecular techniques for the characterisation of Salmonella Typhi isolates from patients in South Africa, 2012–2014.

Methods
For the period 2012–2014, 195 Salmonella Typhi isolates were selected from the culture collection at the Centre for Enteric Diseases (CED). These isolates were subjected to conventional PCR in order to screen for the presence of the Salmonella Typhi H58. Isolates were also analysed using a multiple-locus variable-number tandem-repeats analysis (MLVA) assay that targeted five highly polymorphic variable-number tandem-repeats loci. The assay included PCR amplification of fluorescently labelled VNTR loci and analysis of amplicons by capillary electrophoresis.

Results
Fifty-four percent of isolates (105/195) were identified as Salmonella Typhi H58, while 46% (95/195) were identified as non-H58 Salmonella Typhi strains. MLVA analysis of the 195 isolates revealed 155 MLVA subtypes; the most common subtypes were: STyMT-121, STyMT-114, STyMT-102, STyMT-130, STyMT-132, STyMT-136 and STyMT-139. The MLVA subtyping tool successfully discriminated among Salmonella Typhi H58 isolates and identified MLVA types that were exclusively associated with Salmonella Typhi H58.

Conclusion
The dissemination of the Salmonella Typhi H58 haplotype within South Africa is concerning. Molecular subtyping tools, such as MLVA, can be very useful in disease tracking and for the investigations of typhoid fever outbreaks caused by the Salmonella Typhi H58.

ID: 8539
Category: Sexually Transmitted Diseases (STDSSA)
Permission: Yes
**ABSTRACT TITLE:** COMPARING COMMERCIAL PROBIOTIC AND VAGINAL LACTOBACILLUS STRAINS TO EVALUATE WHETHER PROBIOTICS MARKETED FOR VAGINAL HEALTH IN SOUTH AFRICA ARE SUITABLE FOR BACTERIAL VAGINOSIS TREATMENT

**Introduction**

Bacterial vaginosis (BV) is prevalent globally and associated with increased risk of sexually transmitted infections, including HIV. Antibiotic treatment of BV has high recurrence rates (~50% within 6 months). Adjunctive treatment with probiotics improves efficacy and durability of treatment. We compared characteristics of Lactobacillus strains from commercially-available vaginal probiotics to those isolated from vaginal samples.

**Methods**

Lactobacillus spp. were isolated from commercial vaginal probiotics (10 isolates from 5 products), and from cervico-vaginal fluid (60 isolates from healthy South African women). Species were identified using a MALDI-TOF biotyper. Bacterial size, growth kinetics, effect on pH and on BV associated vaginal species (clinical and ATCC strains of Gardnerella vaginalis and Prevotella bivia) and antibiotic profiles were measured.

**Results**

Lactobacillus spp. isolated from healthy women were L. crispatus, L. jensenii, L. gasseri, L. mucosae and L. vaginalis which were absent from all probiotic products (which contained L. reuteri, L. rhamnosus and L. acidophilus). Clinical isolates grew better at lower pH (pH 2 – pH 6.0) than probiotic strains, and lowered pH more effectively (3.6 vs. 4.4). Culture supernatants from both probiotic and clinical strains promoted on ATCC G. vaginalis-growth, while inhibiting clinical G. vaginalis and all P. bivia strains. All strains were resistant to metronidazole, the most common BV treatment, while the majority of probiotic (8/10) and clinical strains (58/60) were sensitive to clindamycin. All isolates were sensitive to rifamycins (used in TB treatment) and other common antibiotics (including penicillin and amoxicillin). However, probiotic strains were more sensitive to rifamycins and less sensitive to penicillin and amoxicillin than clinical isolates.

**Conclusion**

Characteristics of commercial probiotic Lactobacillus strains are highly variable and did not contain the predominant species isolated from the healthy female genital tract. Clinical L. crispatus isolates had comparable characteristics, and a better probiotic profile than commercial probiotic strains.
and classified as de-escalation (stopping or changing to a narrower-spectrum agent) or administration of a more appropriate agent (if current therapy was inactive against the organism).

**Results**

Of 231 patients, 16 had MRSA, 41 had MSSA and 174 had CoNS based on phenotypic methods. Clinical information was obtained for 184 patients. The Xpert system showed a sensitivity and specificity of 100% (95% CI 93.7–100% and 97.8–100% respectively) for *S. aureus* identification. The sensitivity for detecting methicillin resistance in *S. aureus* was 93.8% (95% CI 71.7–98.9%); specificity was 100% (95% CI 91.4–100%) after resolution of a single discrepancy. The error/invalid rate was 2.58%. Potential impact included a reduction in time-to-final result by 30.5h, with more timely de-escalation in 29 patients and administration of an appropriate antistaphylococcal agent in 17 patients.

**Conclusion**

The Xpert MRSA/SA BC assay performs well in differentiating MRSA, MSSA and CoNS in positive blood cultures. It has potential to improve antibiotic utilisation in approximately 25% of patients. These benefits must be balanced against the assay’s cost in resource-constrained settings. A formal clinical trial may provide further evidence to determine the benefit of the assay.

**ID: 8548**

**Category:** Clinical Microbiology (SASCM)

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**ABSTRACT TITLE:** INTRODUCTION OF DRUG RESISTANT TB – REFLEX TESTING IN KWA-ZULU NATAL – A DESCRIPTIVE ANALYSIS OF MTBDRSl RESISTANT ISOLATES OVER A 9 MONTH PERIOD

**Introduction**

Following WHO recommendations for both the shortened multi-drug resistant (MDR) treatment regimen and use of the GenoType MTBDRsl Line Probe Assay (SL-LPA), the South African Department of Health together with NHLS developed a “DR-TB reflex” testing algorithm. The aim of this is to identify those MDR- or rifampicin-resistant TB patients who can be commenced on the shortened MDR-TB treatment regimen.

**Methods**

874 GenoType MTBDRsl assays performed directly on sputum samples as part of DRTB reflex testing over a four-month period were analysed. The DR-TB reflex testing set comprises a panel of tests: Auramine-O, MGIT960 culture, LPA–first line, SL-LPA and where necessary phenotypic drug susceptibility testing.

**Results**

Of the 874 samples processed, 109 (12.5%) showed resistance to either fluoroquinolones, SL injectable drugs (SLID) or both. Fluoroquinolone-resistant isolates showed mutations in the gyrA gene, with an absent gyrA WT2 and a corresponding positive gyrA mutation1 (MUT1) as the commonest pattern. The second commonest was an absent WT2 with corresponding MUT3C. The least common pattern was an absent WT3 with a corresponding MUT3C. Of note, eight isolates had mutation bands with no corresponding absent WT (6 MUT3C, 1 MUT3B, 1 MUT1). No resistance was detected in the gyrB gene.

**Conclusion**

Preliminary data show 12.5% prevalence of drug resistant TB in the tested period. GyrA and rrs gene mutations are the commonest mutations.
**Introduction**

Antimicrobial resistance (AMR) may be the next global health crisis. We aimed to describe South African patients’ and prescribers’ AMR knowledge, attitudes and perceptions (KAP).

**Methods**

We conducted a cross-sectional KAP survey among a convenience sample of prescribers and patients in South Africa. We used logistic regression to examine associations between knowledge and antibiotic use, beliefs, or behaviours.

**Results**

Mean patient (n = 403) knowledge scores (out of 14) were higher in females (p = 0.0005) and those with more education (p = 0.015); 76% believed the human body, rather than the infectious agent, becomes resistant to antibiotics. After adjusting for education and sex, a 1-unit increase in patient knowledge was associated with increased odds of the following beliefs: (i) important to finish the antibiotic course (aOR 1.34; 95% CI:1.16, 1.66); (ii) over-use impacts AMR (aOR 1.3; 95% CI:1.18, 1.43), but reduced odds of (iii) demanding antibiotics should be given (aOR 0.84; 95% CI:0.74,0.94); (iv) feeling relieved (aOR 0.89; 95% CI:0.81, 0.97) or happy (aOR 0.905; 95% CI:0.82, 0.99) when prescribed antibiotics.

Prescribers (n = 175) > 55 years old had lower median knowledge scores than younger prescribers (p = 0.0005). Those who infrequently prescribe antibiotics when unnecessary had higher knowledge scores (p = 0.01); 70% of prescribers feel pressure from patients to prescribe antibiotics.

Prescribers with higher knowledge scores were more likely to believe the following: (i) to decrease AMR, narrow spectrum antibiotics should be used (aOR 1.41; 95% CI:1.03, 1.92), (ii) explaining to patients disease features which should prompt follow-up (aOR 1.76; 95% CI:1.01, 30.74). Prescribers with higher knowledge scores were less likely to report that antibiotics cannot harm the patient if not needed and less likely to prescribe when not necessary (aOR 0.55; 95% CI: 0.33, 0.91).

**Conclusion**

The association between knowledge and behaviour/perceptions suggests that increasing patient and prescriber knowledge could influence antibiotic use behaviours.
ST307 belonged to two clades. Clade I was associated with K. pneumoniae and rapid PCR identification were used to track the emergence of clones in a real-time fashion. Whole genomic sequencing (WGS) and track the emergence of high-risk antimicrobial resistant (AMR) especially in developing countries, with the ability to recognise measures to curb the spread of antibiotic resistance.

Conclusion
This study described the rapid emergence over a six-year period of K. pneumoniae ST307 clade II with blaOXA-181 in South Africa and highlighted the importance of using WGS to develop molecular surveillance methods for tracking emerging AMR clones in a rapid fashion. Such methods will aid with the implementation of measures to curb the spread of antibiotic resistance.

ABSTRACT TITLE: KLEBSIELLA PNEUMONIAE ST307: A SILENT PATHOGEN ON THE RISE IN SOUTH AFRICA

Introduction
There is a desperate need for molecular surveillance systems, especially in developing countries, with the ability to recognise and track the emergence of high-risk antimicrobial resistant (AMR) clones in a real-time fashion. Whole genomic sequencing (WGS) and rapid PCR identification were used to track the emergence of a novel OXA-181-producing K. pneumoniae clone ST307 in South Africa.

Methods
Illumina WGS was performed on a collection (2011 to 2016) of clinical K. pneumoniae with blaOXA-48-like carbapenemases from South Africa and used this information to design PCR primers for the identification of ST307 and its association with blaOXA-181 on IncX3 plasmids.

Results
Whole genomic sequencing on 28 isolates showed that K. pneumoniae ST307 belonged to two clades. Clade I was associated with blaCTX-M-15 and corresponded to ST307 sequences deposited in Genbank (n = 9). A novel ST307 clade II (that differs from clade I in approximately 100 SNPs) contained blaOXA-181 and was associated with IS 3000 on IncX3 plasmids. Subsequent PCR screening of K. pneumoniae with blaOXA-48-like showed that ST307 clade II with blaOXA-181 was first detected during 2012 in Johannesburg. In 2013 and 2014, 20/135 (15%) of K. pneumoniae with blaOXA-48-like tested positive for ST307 clade II with blaOXA-181 and was found in Alberton, Johannesburg and Pretoria. During 2015 and 2016 the numbers of ST307 clade II increased exponentially; 297/499 (60%) of K. pneumoniae with blaOXA-48-like belonged to ST307. This clade had subsequently spread to 10 other cities (i.e. Bloemfontein, Emalahleni, Ermelo, East London, Klerksdorp, Krugersdorp, Mbombela, Polokwane, Secunda, Vanderbijlpark) across six different provinces (i.e. Eastern Cape, Gauteng, Free State, Limpopo, Mpumalanga, North West).

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ABSTRACT TITLE: EVALUATION OF METHODS FOR GENERATION OF IN VITRO MUTANTS RESISTANT TO BEDAQUILINE, CLOFAZIMINE AND LINEZOLID FROM MYCOBACTERIUM TUBERCULOSIS REFERENCE STRAINS
**Introduction/Aim**

The introduction of bedaquiline and repurposing of linezolid and clofazimine for the management of drug-resistant TB has resulted in improved cure rates. However, there is a paucity of information regarding *Mycobacterium tuberculosis* resistance acquisition mechanisms. We sought to determine the optimum methodology for resistance induction to these drugs and their resistance-associated variants (RAVs) using Whole Genome Sequencing (WGS).

**Methods**

ATCC reference strains with unique resistance profiles were used to compare two in vitro resistance generation approaches; serial passaging through increasing antimicrobial pressure (five strains) and spontaneous generation (two strains). This was done for bedaquiline, clofazimine and linezolid. Generated mutants were confirmed with MGIT 960 drug susceptibility testing (DST). WGS and bioinformatic analysis were performed to identify induced RAVs, which were then classified as known or novel variants. In addition, clofazimine-induced resistant mutants were subjected to bedaquiline MGIT 960 DST to assess cross resistance.

**Results**

Mutants were successfully generated for both approaches in all strains investigated. Generated mutants for each drug using both approaches showed up to an eight-fold increase compared to currently proposed critical concentration values. Spontaneously generated mutants showed a broader range of MIC values (from 2–8 µg/ml) for each drug and were obtained within a month from initiation. Mutants generated through serial passaging exhibited high MIC values (> 8 µg/ml) and these were obtained after at least five passages (> 3 months). RAVs occurred in the rplC (linezolid), rv0678 (bedaquiline and clofazimine) and/or atpE (bedaquiline) regions. We identified both known and novel mutations for bedaquiline and clofazimine within respective genetic targets. Additionally, clofazimine resistant mutants (from both approaches) were cross resistant to bedaquiline without any previous drug exposure.

**Discussion/Conclusion**

Both approaches were capable of mutant generation from various ATCC strains with the spontaneous generation approach being preferred as it is rapid. Mutations identified correspond to known mutations, particularly for linezolid. In addition, for bedaquiline and clofazimine, novel mutations were identified, which warrant further study.

**ID: 8687**

**Category:** Antimicrobial Resistance (SAASP)

**Permission:** Yes

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**ABSTRACT TITLE:** ANTIMICROBIAL RESISTANCE IN A TERTIARY HOSPITAL IN THE EASTERN CAPE
DISCREPANCIES BETWEEN PHENOTYPIC AND MOLECULAR METHODS FOR DETECTION RIFAMPICIN SUSCEPTIBILITIES FROM A TERTIARY DIAGNOSTIC LABORATORY IN PRETORIA: TOWARDS CONSENSUS IN RPOB MUTATIONS

Introduction
Multi-drug resistant tuberculosis (MDR-TB) caused by resistance to at least rifampicin (RIF) and isoniazid (INH) remains a major health concern worldwide. To curb the spread of MDR-TB, GenoType®MTBDRplus was endorsed for the accurate detection and treatment initiation of TB cases whilst waiting for phenotypic drug susceptibility test (DST) results. Discrepancies between phenotypic DST and GenoType®MTBDRplus have been observed. It is thus important to correctly characterise different rpoB mutations causing different RIF susceptibilities. We investigated discrepancies between GenoType®MTBDRplus and phenotypic DST for RIF and sequenced the rpoB gene in susceptible and resistant isolates.

Methods
Forty isolates provided by the NHLS-TB Laboratory were characterised. The isolates were cultured and tested for drug susceptibility, phenotypically using BACTEC™MGIT™ (RIF critical concentration = 1µg/ml) and genotypically with GenoType®MTBDRplus according to manufacturer’s instructions. A PCR amplification of the rpoB gene was performed and the amplicons were sequenced. The sequences of susceptible isolates were used to develop the consensus sequence in silico and used for comparison with the resistant strains to query results obtained with the H37Rv reference strain.

Results
There was 100% agreement (kappa = 1.0) between phenotypic drug susceptibility testing and Sanger sequencing. A relatively lower agreement of 90% (kappa = 0.8) was observed between phenotypic DST and GenoType®MTBDRplus assay. Five mutations were detected and occurred at different frequencies with S531L (50%) occurring most often followed by D516V (20%), H526S (10%), H526D (10%) and a TTC insertion at position 1297 (10%). The consensus sequence constructed gave similar results to H37Rv reference strain.

Conclusion
Discrepancies occur because molecular tests detect resistance at a molecular level and not the ultimate phenotypic expression. Sequencing could help minimise phenotypic and genotypic discrepancies found in most diagnostic laboratories. Similar results between H37Rv and the developed consensus sequence highlight better performance of Sanger sequencing for DST in comparison to GenoType®MTBDRplus.

ABSTRACT TITLE: #CAUTI MUST FALL

Introduction
In 2013, the Catheter Associated Urinary Tract Infection (CAUTI) rate in an internationally renowned 346-bed private hospital was of concern at 3.62 per 1 000 catheter days. Of the 346 beds, 109 are specialised ICU and High Care beds and many of the patients are catheterised. Urinary catheterisation of patients is required for close output monitoring of critically ill patients, selective surgery and urethral obstruction. CAUTI is the most common healthcare-associated infection. CAUTIs result in prolonged hospitalisation, increased use of antimicrobials, with the potential to develop multidrug resistant organisms and increased hospital and patient costs.

Methods
In 2013 a number of CAUTIs were present shortly after admission. A Standard Operating Procedure (SOP) was drafted for ‘on admission’ urinalysis on all patients. The SOP was rolled out to all units, training was given and CAUTI audits conducted. A CAUTI Champion was appointed in late 2013 and a multidisciplinary CAUTI committee was formed, which included an Urologist, Infection Control, Nursing Management and Unit Managers, and regular meetings were held. In addition, CAUTI Unit representatives were identified and trained. A CAUTI Workshop presented by the Urologist was arranged and CAUTI rounds were conducted in all units and gaps identified were
discussed with possible solutions. A campaign for the early removal of "awake patients" catheters was developed and driven by IPC and unit CAUTI champions.

Results
The 2014 CAUTI rate/1 000 catheter days was 2.81 down from 3.62 in 2013. In 2015 the CAUTI rate fluctuated to 2.85 and required further focus. By 2016, the CAUTI rate per 1 000 catheter days decreased to 2.04.

Conclusion
Collaborative team efforts, constant training, reminders to do admission urine tests, and audits is required to prevent CAUTIs and sustain a reduction. The nature of the patients at the hospital is such that many will require urine catheters, but management and prompt removal is of paramount importance.

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ABSTRACT TITLE: MENINGOCOCCAL CARRIAGE AMONGST FIRST-YEAR STUDENTS ON ENTERING UNIVERSITY: IS THERE A NEED IN SOUTH AFRICA FOR MENINGOCOCCAL PREVENTION STRATEGIES?

Introduction
Although peak incidence of invasive meningococcal disease (IMD) is in infants, a second peak is seen in adolescents due to their social interactions and high carriage of Neisseria meningitidis (NM). Adolescent carriage prevalence ranges from 3% in the African meningitis belt to 23% in the United Kingdom. IMD is a severe disease associated with outbreaks. Prevention strategies should be sought even with low IMD incidence.

Methods
To determine the NM carriage prevalence and carriage risk factors amongst first-year students upon entering university, we collected oropharyngeal swabs and questionnaires from students during registration week at two large universities in Western Cape (WC) and Gauteng Provinces (GA) in 2017. Swabs were placed directly into Todd Hewitt broth and incubated for 24 hours; plated onto Thayer Martin media and incubated for a further 24–48 hrs in 5% CO 2. NM colonies were confirmed using MALDI-TOF mass spectrometry. Risk factors for carriage were analysed using STATA version 14.0.

Results
Two-thousand-one-hundred-and-twenty-one students completed the questionnaire and swab, 901 (42%) in WC and 1 220 (58%) in GA. Median age was 18 years, 41% (876) were male and 0.8% (16/1 985 with known HIV status) HIV-infected.

Sixty-six students (3%, 95% confidence interval (CI) 2.5–3.9%) were NM carriers, of which 52% (34/66, p = 0.09) were male and all were HIV-uninfected.

When adjusting for home province (KwaZulu-Natal (KZN) (Odds Ratio (OR)3.3, CI1.5–7, p = 0.003) and WC (OR3.4, CI1.8–6.2, p = 0.004) compared to GA), smoking (OR3.1, CI1.9-5, p < 0.001), smoke exposure (OR4.7, CI1.3–6.7, p < 0.001), intimate kissing partners (OR3.8, CI2.2–6.5, p < 0.001), party (OR2.1, CI1.3–3.4, p = 0.003), nightclub (OR3.7, CI2.6–6.1, p < 0.001) or pub (OR3, CI1.8–4.9, p < 0.001) attendance in the last two weeks – home province (Eastern Cape adjusted – OR3.1, CI1.1–8.8, p = 0.04; KZ aOR3.7, CI1.7–8.3, p = 0.01; WC aOR2.6, CI1.4–5.0, p = 0.004, compared to GA), exposure to smoke (aOR2.9, CI1.3–6.7, p = 0.01) and intimate kissing partners (aOR2.7, CI1.5–4.8, p = 0.001) remained significant risk factors for carriage.

Conclusion
NM carriage in adolescents entering university was 3%. Significant NM carriage risk factors were home province, secondary smoke exposure and intimate kissing. Targeting adolescent smoke exposure is a practical health promotion activity that could prevent meningococcal circulation.

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ABSTRACT TITLE: CORONAVIRUS DIVERSITY IN SOUTH AFRICAN BATS – A RISK FOR HUMAN HEALTH?

Introduction
Emerging infectious diseases pose a major threat to human health. Bats are recognised as reservoir hosts for many important viruses with zoonotic potential, including coronaviruses (CoV). Following the emergence of SARS in 2002/03 and the subsequent identification of *Rhinolophus sinicus* as the likely ancestral SARS-CoV source, a wide diversity of bat CoV has been described worldwide. Our research, in transdisciplinary collaborations with ecologists and zoologists, aims to better understand CoV diversity and ecology in South African bats.

Methods
We conducted general surveillance and species-specific studies of bats across South Africa. Samples – including faecal pellets, saliva and urine swabs, and from voucher specimens sacrificed for museum collections, also blood and organs – were screened for CoV genome by a widely used Pan-CoV PCR assay and 816bp RGU fragment sequences (Drexler et al., 2010) used to construct ML trees in MEGA v7.

Results
An improved screening method greatly increased the CoV detection rate: 92 of 686 samples from 9 bat species were screening-positive: 66 for α-CoV, 19 for β-CoV, and 7 for both. Most sequences are α-CoVs, with ~20% prevalence in *Neoromicia capensis* as the likely ancestral SARS-CoV source. Preliminary analyses of partial RdRp, nucleocapsid and spike gene fragments of novel β-CoV identified in *Neoromicia* and *Pipistrellus* bats show their close relatedness to BiCoV PML-PHE1/RSA/2011 (NeoCoV), previously found by us in *N. capensis* and belonging to the same viral species as the recently emerged MERS-CoV, responsible for the ongoing outbreak in the Arabian Peninsula.

Conclusions
Extensive sampling allowed detection of α- and β-CoV from a wide range of bat species across large parts of South Africa. An improved screening approach yielded significantly more positives. There is substantial CoV diversity in southern African bats. Most importantly, our finding of additional MERS-CoV-related CoV will help address the unresolved question of the origin of this zoonotic pathogen.
isolates subjected to WGS, most common vaccine serotypes were 1, 5, 14, 6A and 6B (26%, 13%, 8% and 8% respectively). Non-vaccine serotypes were rare and included serotypes 24 (8%) and 35D (3%).

Most pneumococcal isolates tested were susceptible to a broad range of antibiotics. A majority of serotypes were resistant to both trimethoprim-sulfamethoxazole and tetracycline. Further genomic analyses to uncouple these observations are in progress.

**Conclusion**
The occurrence of a pneumococcal serotype 1 and 5 meningitis outbreak three years after the introduction of PCV13 is alarming and calls for strengthening of meningitis surveillance and a re-evaluation of the current vaccination programme in high risk countries.

**ID: 8688**

**Category:** Paediatric Infectious Diseases (SASPID)

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**ABSTRACT TITLE:** TREATMENT OUTCOMES IN PERINATALLY-HIV-INFECTED CHILDREN AND ADOLESCENTS AFTER 10 YEARS ON ANTIRETROVIRAL THERAPY

**Introduction**
The burden of paediatric HIV in South Africa has shifted to older children and adolescents. Nevertheless, information on long-term treatment outcomes of perinatally-HIV (PHIV) infected children is limited. We examined long-term immunologic and virologic outcomes of children who remained in care for ≥ 10 years after starting antiretroviral therapy (ART).

**Methods**
We included PHIV-infected children who initiated ART at a Cape Town clinic between 2002 and 2005, and had follow-up for ≥ 10 years from ART initiation date. CD4 counts and viral loads (VL) were analysed for each successive year on ART.

**Results**
The median follow-up among 127 patients included was 12.2 years (interquartile range [IQR] 11.1–13.0). At ART initiation, median age was 2.6 years (IQR 1.3–4.9) and mean CD4 percentage was 13.7% (95% Confidence Interval [CI] 13.6–13.9). The first ART regimen was non-nucleoside reverse transcriptase inhibitor (NNRTI)-based (64%) or protease inhibitor (PI)-based (36%). After 10 years since ART initiation, 49.6% of patients were on 1st-line, 43.3% on 2nd-line, 6% on non-nucleoside reverse transcriptase inhibitor (NNRTI)-based and 3.1% on lamivudine monotherapy and 3.9% on no ART. After 10 years on combined ART, 87.5% had CD4 > 500 cells/µl (98% of those on 1st-line vs 79% of those on 2nd-line ART; p = 0.003). In those who had VL performed after 10 years, 74.3% (81/109) had VL < 400 copies/ml (81.5% of those on 1st-line vs 69.2% on 2nd-line
ART; \( p = 0.16 \). The 10-year probability of experiencing viral failure was 56.7% (95% CI 48.3–65.5), and was higher in children on NNRTI-based regimens vs PI-based regimens (65.8% vs 43.1%; logrank test, \( p = 0.04 \)). The 10-year probability of switching to 2nd-line ART was 45.7% (95% CI 37.5–54.8).

**Conclusion**

Virologic and immunologic outcomes are good overall in children who remain in care for ≥10 years, but >40% of children were on 2nd-line ART with poorer immunologic outcomes.

**ID: 8307**

**Category:** Sexually Transmitted Diseases (STDSSA)

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**ABSTRACT TITLE:** STRONG ASSOCIATION BETWEEN HIGHER RISK SEX AND HIV PREVALENCE AT REGIONAL LEVEL: AN ECOLOGICAL STUDY OF 27 SUB-SAHARAN AFRICAN COUNTRIES

**Introduction**

It is unclear why HIV prevalence varies by nearly two orders of magnitude between regions within countries in sub-Saharan Africa. In this ecological study, we assess if HIV prevalence by region is associated with any of four markers of higher risk sexual behaviour: lifetime number of partners, multiple partners in past year, higher risk sex (defined as sex with non-cohabiting, non-marital partners) and age at debut.

**Methods**

We performed Pearson’s correlation between the four behavioural risk factors and HIV prevalence by region in 47 nationally representative surveys from 27 sub-Saharan African countries, separately by gender. In addition, principal components analysis was used to reduce the eight risk factors (four for each gender) to two principal components (PCs). Mixed effects linear regression was used to assess the relationship between the resulting two PCs and HIV prevalence after controlling for the prevalence of male circumcision.

**Results**

HIV prevalence varied by a median 3.9-fold (IQR 2.9–8.5) between regions within countries. HIV prevalence was strongly associated with higher risk sex and, to a lesser extent, the other risk factors evaluated. Both PCs were strongly associated with HIV prevalence when assessed via linear regression.

**Conclusion**

Differences in sexual behaviour may underpin the large differences in HIV-prevalence between subpopulation within sub-Saharan African countries.

**ID: 8677**

**Category:** Infectious Diseases (IDSSA)

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**ABSTRACT TITLE:** PRELIMINARY EVALUATION OF DRIED BLOOD SPOTS TO TEST FOR HEPATITIS C VIRUS (HCV) ANTIBODIES AMONG PEOPLE WHO INJECT DRUGS, SEX WORKERS AND MEN WHO HAVE SEX WITH MEN

Hepatitis C virus infection is a significant cause of morbidity and mortality worldwide (Mahd Hanafiah et al., 2013). It has been estimated that 130–170 million people are chronically infected with HCV and 3–4 million new infections per year (Bennette S et al., 2012). The highest prevalence of HCV infection was reported in Asia and Africa (Mohd Hanafiah et al., 2013). The prevalence of 2.1%–2.8% has been reported in sub-Saharan Africa (Lyden et al., 2014) and 0.1%–1.7% sero-prevalence reported in South Africa (Abuehassan, 2012). However, HCV infection still remains as the most under-diagnosed virus leaving those populations that are infected unaware of their infection status (Tuaillan E et al., 2009) especially in South Africa. It has been reported that the most-at-risk population are unaware of their HCV infection status including the high-risk groups (Bennett et al., 2012). This could be due to the stigma and discrimination they are faced with when accessing healthcare facilities and especially when they test HCV positive.

In order to diagnose HCV infection, patients have to undergo several phlebotomy procedures for multiple tests. This can pose a challenge for PWIDs, who are commonly referred to as ‘hard sticks’ and may be restricted for testing altogether (Dokubo EK et al., 2014). In some international settings, blood draws are not culturally accepted; as a result the testing of HCV is not a priority. Testing protocols may also limit HCV diagnosis; in other circumstances venipuncture is not a convenient method that is readily available and in other parts of the world diagnostic technology is limited therefore making the diagnosis of HCV in resource-limited settings a challenge (Dokubo EK et al., 2014).

There have been numerous HCV diagnostic tests which have been invented throughout the years, most of which are used in
centralised laboratory facilities. These sample matrices are cheaper, reliable and faster tests. These tests are easy to handle, store and can be transported from distant clinics to the centralised laboratory without refrigeration and retain the integrity of the specimen (Vinikoor et al., 2015). Various studies have validated the use of these sample matrices such as the Dried Blood Spot (DBS) and found a sensitivity of 93.8–100% and a specificity of 94.0–100% (Bennett et al., 2012).

DBS offers less invasive measure of sample collection and they also have a potential to facilitate the detection of HCV antibodies (Dokubo EK et al., 2014). This could potentially lead to early diagnosis, reduced risk behaviours and better treatment effects (Dokubo EK et al., 2014). This will scale up testing for HCV infection among high-risk groups in South Africa, more especially in PWIDs group which has been reported to be the population with the highest HCV prevalence (Dokubo EK et al., 2014). It can also be expected that HCV prevalence data in South Africa among the most-at-risk population will be generated also have a positive improvement to the current South African healthcare outcomes.

This study is conducted at NICD and is part of a larger collaboration on hepatitis C infection in key populations (people who inject drugs (PWID), sex workers (SWs) and men who have sex with men (MSM)) accessing HIV testing services in Cape Town, Durban, Pietermaritzburg, Mthatha, Port Elizabeth, Johannesburg and Pretoria. Currently, I do not have any funding for attending the conference. It would be of great pleasure if I could attend the conference. It would be of great pleasure if I could attend the conference because my study (research) offers new insights about sample matrixes for HCV testing to the larger population (research scientist clinicians and the general population) and hopefully will improve the current patient care treatment and HCV database for HCV in South Africa.

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ABSTRACT TITLE: EPIDEMIOLOGY OF INFLUENZA - AND RESPIRATORY SYNCYTIAL VIRUS (RSV)-ASSOCIATED HOSPITALISATION IN CHILDREN AGED

Introduction
Few data are available describing the spectrum of influenza- or respiratory syncytial virus (RSV)-associated hospitalised illness in children aged < 5 years in sub-Saharan Africa.

Methods
During 2011–2016, we conducted prospective, hospital-based surveillance for severe respiratory illness (SRI) in children aged < 5 years at two sites. In infants aged 2 days to < 3 months, SRI was defined as a diagnosis of suspected sepsis or physician-diagnosed acute lower respiratory tract infection, irrespective of signs and symptoms and duration of illness. In children aged 3 to 59 months, SRI was defined as a physician-diagnosed lower respiratory tract infection, irrespective of signs and symptoms and duration of illness. Nasopharyngeal aspires were tested for influenza and RSV by real-time reverse transcription polymerase chain reaction. We estimated rates of influenza- and RSV-associated hospitalised SRI. We examined clinical and epidemiological differences between...
children who tested positive for influenza versus RSV using multivariable penalised logistic regression.

**Results**

Among 3,657 hospitalised children, 200 (5.5%) tested positive for influenza viruses, 877 (24.0%) for RSV, and 11 (0.3%) for both (excluded from further analyses). The median age of children hospitalised with influenza was 14 months versus 4 months for RSV ($p < 0.01$). Influenza-associated hospitalisation rates per 100,000 were highest among infants aged 6–11 months (524, 95% Confidence Interval (CI) 389–677), while RSV-associated hospitalisation rates were highest in infants aged 0–2 months (6,557, 95% CI 5,972–7,261). HIV-exposure was associated with increased incidence of influenza- and RSV-associated hospitalisation in infants aged 0–5 months, relative risk (RR) 2.2 and 1.5 respectively. HIV-infection was associated with increased incidence of influenza- and RSV-associated hospitalisation in all age groups, RR range 2.2–6.7.

**Conclusion**

Influenza- and RSV-associated hospitalisation were common among South African infants. HIV-infection and HIV-exposure increased risk of influenza- and RSV-associated hospitalisation in South African children.

**ID: 8638**

**Category:** Paediatric Infectious Diseases (SASPID)

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**ABSTRACT TITLE:** CULTURE-CONFIRMED TUBERCULOSIS IN INFANTS LESS THAN THREE MONTHS OF AGE: A CLINICAL DESCRIPTION OF 71 CASES

**Introduction**

Young infants are particularly at risk for disseminated and severe pulmonary tuberculosis (TB). Few studies address TB disease in infants less than three months of age. The aim of this study was to describe clinical and radiological characteristics of TB in a cohort of infants in this age group.

**Methods**

We conducted a retrospective, descriptive study of all infants below three months (91 days) of age (on date of admission to hospital) with culture-confirmed TB on any clinical specimen. The infants were identified using a database of prospectively collected *Mycobacterium tuberculosis*-positive culture results for all children < 13 years who presented to Tygerberg Hospital, a tertiary referral hospital in South Africa, from 1 March 2003 through 30 June 2011. Infants with mycobacterial disease caused by *M. bovis* BCG and non-tuberculous mycobacteria were excluded.

**Results**

Seventy-one infants met the inclusion criteria. Nineteen infants (27%) were classified as having congenital TB. Sixty-six (92%) had pulmonary TB, of which 22/66 (33%) also had extrapulmonary TB; four (6%) infants had only extrapulmonary TB. Twenty-seven infants (38%) were HIV exposed and 10 infants (14%) were HIV-infected. Five (7%) infants had drug-resistant TB. Cough, loss of weight or failure to gain weight and history of fever were the most common presenting symptoms and were found in 41 (58%), 29 (41%) and 24 (34%) of infants respectively. Eighteen (25%), 15 (21%) and 6 (8%) infants presented with respiratory distress, wheezing and stridor, respectively. Chest X-ray abnormalities included: mediastinal lymphadenopathy (39; 55%), large airway compression (39, 55%), alveolar/bronchopneumonic opacification (47; 66%), miliary pattern (14; 20%) and cavities (9; 13%) infants. Of the 39 with airway compression, 21 (54%) underwent bronchoscopy and 21 (54%) infants required lymph node decompression. Ten (14%) infants died.

**Conclusion**

Infants under three months of age presented with severe pulmonary manifestations and also disseminated disease.

**ID: 8294**

**Category:** Sexually Transmitted Diseases (STDSSA)

**Permission:** Yes

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**ABSTRACT TITLE:** PREVALENCE AND MOLECULAR ANALYSIS OF MYCOPLASMA GENITALIUM STRAINS ISOLATED FROM PREGNANT WOMEN AT AN ACADEMIC HOSPITAL IN PRETORIA, SOUTH AFRICA
Introduction

*M. genitalium* is sexually transmitted causing urethritis in men and cervicitis in women. Treatment is syndromically and detection of antimicrobial resistance and typing of strains relies on DNA sequence data. High levels of sequence variability between clinical isolates are seen which may be associated with antimicrobial resistance. This study was done to determine the prevalence and the molecular characteristics of *M. genitalium* strains from pregnant women attending the Dr George Mukhari Academic Hospital (DGMAH).

Methods

Endocervical swabs were collected from 100 pregnant women attending DGMAH. The specimens were screened for *M. genitalium* using a real-time PCR assay. Genotypic resistance markers for macrolide and fluoroquinolones were determined by sequence analysis of the V-region of the 23S rRNA, gyrA, and parC genes respectively. The strains were typed using mgpB single-nucleotide polymorphism typing (SNP) and MG309 variable number tandem (VNTR) analysis.

Results

The prevalence of *M. genitalium* was 7.0%. In two isolates macrolide resistance-associated mutations were seen. No resistance-associated mutations were seen in the gyrA genes, but a parC fluoroquinolone resistance-associated mutation was seen in one of the macrolide resistant isolates. SNP typing revealed four sequence types, and four different types were also seen using MG309 VNTR analysis. Typing assigned *M. genitalium* to two major clusters with macrolide and fluoroquinolones resistance in the same cluster.

Conclusion

*Mycoplasma genitalium* is frequently undiagnosed. This study reported the first multidrug resistant *M. genitalium* strain in South Africa. As azithromycin was included in the national syndromic treatment guidelines in 2015, it is alarming to already find resistance-associated genes.

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**ABSTRACT TITLE: EVALUATION OF TWO COMMERCIAL NAATS FOR DETECTION OF C. TRACHOMATIS, N. GONORRHOEAE AND T. VAGINALIS IN ENDOCERVICAL SWABS COLLECTED FROM PREGNANT WOMEN ATTENDING THE DGMAH**

Introduction

Each year about 500 million new cases of sexually transmitted infections occur in men and women worldwide. Nucleic acid amplification tests offer a sensitive and specific diagnostic approach for the detection of STIs. The aim of the study was to evaluate two commercial nucleic acid amplification tests for the detection of *Chlamydia trachomatis* (CT), *Neisseria gonorrhoeae* (NG) and *Trichomonas vaginalis* (TV) in endocervical swabs.

Methods

One hundred endocervical swab samples that were collected from women attending the termination of pregnancy and ante-natal clinics at DGMAH were used. Multiplex detection of NG, CT and TV was done using the Real-time NG/CT/TV (Sacace, Italy) and Seegene Seeplex ACE Detection (Seegene, Korea) assays according to the manufacturers’ instructions as well as by an in-house PCR. An expanded gold standard was used to evaluate the assays.

Results

Using the Sacace real time m-PCR, NG was detected in 5%; CT in 24% and TV in 7% of the samples. Using Seegene Seeplex ACE Detection assay, NG was detected in 4%, CT in 22% and TV in 4% of the specimens while using in-house PCR, NG was detected in 6%, CT in 18% and TV in 9% of the samples. When applying the expanded gold standard the prevalence of the pathogens was as follows: *N. gonorrhoeae* 4%; *C. trachomatis* 22% and *T. vaginalis* 5%. The Seegene Seeplex assays performed the best in terms of detecting *N. gonorrhoeae* and *C. trachomatis*, followed by the Sacace real time assay. The in-house PCR was the most sensitive and specific in detecting *T. vaginalis*.

Conclusion

The three molecular assays used in this study were cost-effective and fairly sensitive and specific for the three pathogens. The two commercial assays were more successful at detecting *N. gonorrhoeae* and *C. trachomatis* while the in-house PCR best detected *T. vaginalis*.

**ID: 8399**

**Category:** Sexually Transmitted Diseases (STDSSA)

**Permission:** Yes

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**ID: 8585**

**Category:** Infection Control (ICSSA)

**Permission:** Yes

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**ABSTRACT TITLE:** EVALUATION OF THE BD MAX FOR THE MULTIPLEX DETECTION OF MRSA, CRE AND ENTEROPATHOGENS

**Introduction**

Multiplex qPCR has been introduced into the diagnosis of bacterial pathogens. Such assays produce rapid, sensitive and specific diagnosis and can identify multiple organisms simultaneously as compared to culture. In detecting multi-drug resistant organisms such as MRSA and CRE and in diagnosis of Enteric diseases, a rapid diagnosis allows for timeous infection control strategies and a better public health response. The BD MAX™ System is an open fully automated real time PCR instrument which can also run different assays simultaneously. This study will evaluate 3 BD Max assays: the BD Max MRSA XT and Extended Enteric Bacterial Panel and the BioGx Assay for VIM, GES, OXA, NDM, KPC and mcr-1 (CRE Assay).

**Methods**

Positive and negative cultures for each assay were spiked into swabs or stool samples at 1x10^4 CFU/ml. Results were compared to culture (MRSA and Enteric) or a Roche LightMix real time PCR assay (CRE). Limits of detection were determined through 1:10 serial dilution of two positive samples per assay.

**Results**

All assays had accuracy, sensitivity, specificity, positive and negative predictive values of 100%. The limits of detection for each assay were: 1x10^4 4 CFU/ml (Enteric), 1x10^3 3 CFU/ml (MRSA) and 1x10^2 2 CFU/ml (CRE) respectively. Preparation of samples took 2-10 minutes and in total the assays took +/- 2.5 hours to run.

**Conclusion**

The BD Max is a simple, rapid, highly sensitive and specific system for detection of multiple bacterial pathogens with minimal expertise required. The MRSA assay also eliminates false positives in methicillin-resistant coagulase negative Staphylococcus and false negatives in mecC containing S. aureus. The CRE detects presence of several carbapenemases and colistin resistance but culture is still required for identification. Being an open system additional assays can be designed to run on one instrument. The BD Max is therefore a suitable platform especially in resource-limited laboratories.

**ID: 8692**

**Category:** Infectious Diseases (IDSSA)

**Permission:** Yes

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**ABSTRACT TITLE:** OUTBREAK PREPAREDNESS AND THE NICD 24-HOUR HOTLINE – A REVIEW OF CALLS MADE TO THE HOTLINE, AUGUST 2016 TO JULY 2017

**Introduction**

The International Health Regulations (IHR) Core Capacity Monitoring framework requires public health services to maintain 24-hr availability for outbreak alert and response. The NICD hotline
cellphone number is widely distributed to facilitate investigation of communicable diseases with public health importance. We review calls made to the NICD 24-hr hotline to assess adherence to IHR, technical content of calls made, and role of the hotline in South African outbreak responses.

Methods
Calls to the NICD hotline (082-883-9920) are documented using a standardised data collection tool by each doctor on call. Post-call alerts are defined as outbreaks/potential outbreaks, and risk assessment is graded using WHO Mass Gatherings risk characterisation as minimal, minor, moderate, major or severe. Summary statistics are generated for number and origin of alert/call, type of enquiry, caller details, clinical details, diagnosis and turnaround time.

Results
From 1 July 2016 to 30 June 2017, 1,083 calls were documented, with an average of 90 calls/month (standard deviation (SD) 3.57). Six-hundred-and-thirteen (57%) originated from the private sector. The majority of callers were situated in Gauteng (409, 38%), KwaZulu-Natal (294, 27%) and Western Cape provinces (167, 15%) respectively. Calls pertained to rabies post-exposure prophylaxis (PEP) (484, 45%), requests for assistance with diagnosis of infectious diseases in individual patients (180, 16%), PEP for non-rabies conditions (139, 13%), and advice regarding public health management for laboratory-confirmed conditions (117, 11%) or other reason (162, 15%). Of 878 calls with a risk characterisation, 50 (6%) alerts were identified as ‘outbreaks’, 18 (2%) events attained ‘moderate’ risk status and 15 deaths were reported. Food-borne illness were the most common outbreak leading to an alert, (21/50) followed by viral haemorrhagic fever (5).

Conclusion
The NICD 24-hr hotline contributes to adherence to IHR, and facilitates outbreak alert and response activities.

ABSTRACT TITLE: PREVALENCE AND CHARACTERISATION OF FOOD AND WATERBORNE BACTERIAL PATHOGENS IN RIVER WATER, MEAT FROM INFORMAL ABATTOIRS, AND STOOL FROM CHILDREN WITH DIARRHOEA IN NYANGA TOWNSHIP, CAPE TOWN, SOUTH AFRICA

Introduction
Morbidity and mortality due to diarrhoeal diseases is heaviest among children in sub-Saharan Africa and south-east Asia, especially in areas with poor hygiene. Informal settlements face hygiene challenges and have high infant mortality rates; yet little is known about the contribution of foodborne diarrhoeic pathogens. We aimed to determine the prevalence and characteristics of bacterial pathogens in the stool of children with diarrhoea, river water, and meat from informal abattoirs in Nyanga. Also, to assess the level of faecal contamination of the river water.

Methods
Samples were collected between September 2015 and May 2016 in Nyanga, Cape Town. Clinical and epidemiological data were collected using a questionnaire. A duplex real-time PCR assay, gel-based PCR, and CHROMagarTME STEC were used to screen Tryptic Soy Broth enrichments of the samples for stx, rfbE and wbdl, and STEC respectively. The enrichments (or sample) were also tested for selected foodborne pathogens using standardised methods. The resultant bacterial pathogens were serotyped (Salmonella and Shigella), tested for virulence genes, and antibiotic susceptibility determined.

Results
A total of 64, 66, and 85 water, stool, and meat samples respectively were collected. Diffusely adherent *E. coli* (DAEC-18% [95% Confidence Interval = 11–29%]), and *Shigella flexneri* (17% [95% CI = 9–27%]), were most prevalent in the stool of children (mean age = 14.9 months) that presented with acute diarrhoea (mean duration = 2.5 days) with one or more pathogens detected in 73% (48/66) of children. Other pathogens in stool included *Salmonella* (6%), *Plesiomonas* (9%), *Aeromonas* (3%), and *Campylobacter* (5%). Meat carried *Salmonella enterica* (5%), *Aeromonas sobria* (3%), *Campylobacter jejuni* (5%), and diarrhoeic *E. coli* (3%). Water was highly contaminated (mean CFU/ml = 1.11E+06 –2.74E+05) and carried *Plesiomonas shigelloides* (17%), *Shigella flexneri* (5%), *Vibrio vulnificus* (9%), and diarrhoeic *E. coli* (7%). Trimethoprim-sulphamethoxazole resistance was observed amongst all pathogens, irrespective of their source.

Conclusion
Foodborne pathogens were prevalent in human and non-human sources in this informal settlement. Local and international public health standards should be implemented, and further larger scale research is needed to fully define the prevalence of foodborne pathogens in similar settings in South Africa.
**ID: 8363**

**Category:** Infectious Diseases (IDSSA)

**Permission:** Yes

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**ABSTRACT TITLE:** THE DEVELOPMENT AND APPLICATION OF A HIGH THROUGHPUT METHODOLOGY TO DETERMINE MICs OF MYCOBACTERIUM TUBERCULOSIS ISOLATES AGAINST ANTIMICROBIAL AGENTS

**Introduction**

Drug susceptibility testing of Mycobacterium tuberculosis is time consuming and expensive. Multi-point inoculation offers the advantage of testing multiple isolates on a series of solid media with a single break-point concentration of a drug in each plate or a series of different drug concentrations of one drug. The former provides the resistance profile of the isolates tested while the latter results in an MIC. This has the potential to reduce consumables and labour costs for break-point susceptibility testing as well as MIC determination. Although multi-point inoculation had been performed with other bacterial species, it has hardly been used with M. tuberculosis since its slow growth and clumping has been thought to be factors to influence reproducibility of this method adversely. We aimed to determine the reproducibility of MIC determination for anti-TB drugs of M. tuberculosis isolates using agar dilution with multi-point inoculation, and thereafter validating the results by comparing it to classic agar dilution on quadrant plates and the MTT assay.

**Methods**

Thirty M. tuberculosis isolates were grown in Middlebrook 7H9 broth with 20% Tween until mid-log phase was reached. Agar dilution MICs were determined on Middlebrook 7H10 agar for 11 anti-TB drugs at concentrations ranging from 128 to 0.125 mg/L. The agar plates were inoculated using a multi-point inoculation device with 36 points each delivering 1 micro litre of a suspension of 1X10^4 cfu/ml. For the quadrant plate method and the MTT assay 100 micro litre of the same suspension was used. All tests were done in triplicate.

**Results**

Agar dilution with multi-point inoculation was found to be reproducible within the 11 anti-TB drugs tested and correlated well with agar dilution on quadrant plates and the MTT assay for the three anti-TB drugs tested.

**Conclusion**

The multi-point inoculation method has potential for wide scale application in break-point drug susceptibility testing as well as MIC testing of M. tuberculosis isolates.

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**ID: 8396**

**Category:** Infectious Diseases (IDSSA)

**Permission:** Yes

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**ABSTRACT TITLE:** THE MANAGEMENT AND OUTCOMES OF STAPHYLOCOCCUS AUREUS BACTERAEMIA AT A SOUTH AFRICAN REFERRAL HOSPITAL: A PROSPECTIVE OBSERVATIONAL STUDY

**Introduction**

*Staphylococcus aureus bacteraemia* (SAB) is an important cause of mortality and morbidity. Limited evidence exists on the management and outcomes of SAB in a South African setting. The aim of this study is to describe a local cohort of patients with SAB, assess outcomes, and explore the factors associated with complications and death.

**Methods**

We conducted a prospective observational study of all patients over the age of 13 years admitted to Groote Schuur Hospital with SAB. The following data were collected: demographics and medical co-morbidities, management, and 90-day outcomes. We summarised patient characteristics and examined factors associated with poor outcomes using multivariate logistic regression.
Results

One hundred consecutive distinct SAB infection episodes, in 98 patients, were included between November 2013 and January 2015; median age 50 years (interquartile range (IQR) 33 to 63), males 72%, HIV infection 21.5%. Median time to notification of culture results was 51.5 hours (IQR 41.0 to 66.8). SAB was healthcare-associated in 68.4%, with 44.7% linked to drip-site infection; 24.0% of cases were caused by methicillin-resistant S. aureus (MRSA). Ninety-day mortality was 47%, with 83.3% of deaths attributable to SAB. Independent prognostic factors for 90-day mortality were MRSA infection (OR 3.1; 95% confidence interval (CI) 1.1 to 9.7) and the presence of co-morbidities (OR 4.8; 95% CI 1.5 to 19.8); the risk for complicated infection was higher with suboptimal or inadequate antibiotic therapy (OR 7.2; 95% CI 1.6 to 39.9). Definitive antibiotic therapy was suboptimal or inadequate in 22.6% of all cases, and in 35.3% of those with MRSA.

Conclusion

Most episodes of SAB were healthcare associated and related to drip-site infection, suggesting that many of these infections are preventable. MRSA rates and overall SAB mortality were higher than in high income countries. Our findings suggest that there is a need for improved management.

ID: 8443

Category: Infectious Diseases (IDSSA)

Permission: Yes

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ABSTRACT TITLE: INCIDENCE OF BACTERIAL INFECTIONS AND COLONISATION IN PATIENTS ADMITTED TO A TUBERCULOSIS HOSPITAL

Introduction

Hospital-acquired infections are a worldwide problem, which poses an extreme threat to public health. Patients with drug-resistant tuberculosis (TB) are treated with multiple antibiotics including moxifloxacin, linezolid, and meropenem. Broad spectrum antibiotic exposure and prolonged hospital admission puts them at greater risk for colonisation by multi-drug resistant (MDR) bacteria. The aim of the study was to determine the spectrum of bacterial colonisation in TB patients upon admission and during hospitalisation.

Method

Nasal, groin and rectal swabs (for the detection of ESBLs, carbapenem-resistant Enterobacteriaceae (CRE), vancomycin-resistant enterococci (VRE) and MRSA) were analysed from a cohort of patients (n = 37: 28 (community), nine (healthcare facilities)) admitted to a TB hospital upon admission and at four-week intervals thereafter during hospitalisation. Identification and antimicrobial susceptibility testing of bacterial isolates (n = 64) were determined at National Health Laboratory Services (NHLS) by the VITEK-MS system. PCR and DNA sequencing were used for detection of carbapenem resistance genes.

Result

Patients (n = 13/37; 35%) were colonised by an MDR pathogen (ESBL, MRSA) on admission. Colonisation rates were lower in patients admitted from the community (9/28; 32%) compared to those transferred from other healthcare facilities (4/9; 44%). All patients from the community (17/37; 46% of total patients) became colonised by MDR bacteria within one month of admission, mostly with ESBL producing Enterobacteriaceae. Only two patients had MRSA colonisation at admission. Among ESBL, Enterobacteriaceae, Escherichia coli (42/64; 65.6%) and Klebsiella pneumoniae (15/64; 23.4%) predominated. Two Enterobacteriaceae isolates with reduced carbapenem susceptibility did not contain carbapenemase-encoding genes. Nineteen percent (7/37) of patients demised during their hospitalisation, however no deaths could be attributed to drug-resistant organisms.

Conclusion

This study provides insight into the spectrum of bacterial pathogen colonisation in hospitalised TB patients and suggests a review of infection control programmes and practices at the TB hospital.

ID: 8696

Category: Infectious Diseases (IDSSA)

Permission: Yes

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ABSTRACT TITLE: LESSONS LEARNT FROM AN ACTIVATION OF THE NATIONAL EMERGENCY OPERATIONS CENTRE, SOUTH AFRICA, 2017

Introduction

In response to the western African Ebola crisis of 2014, the World Health Organization recommended that all State Parties should develop, strengthen and maintain their capacity to respond promptly and effectively to public health risks and public health emergencies of international concern. To address public health emergencies national and subnational health authorities require efficient and coordinated responses from competent and skilled staff in equipped facilities, supported by appropriate protocols and training. Via a memorandum of agreement (MOA) with the National Department of Health, an Emergency Operations Centre (EOC) was developed at the National Institute for Communicable Diseases (NICD).

Methods

To test the preliminary protocols and systems of the EOC – in the absence of a large-scale infectious disease outbreak – the NICD accepted the responsibility of activating the EOC in response to a public health emergency of a non-infectious nature. The activation was governed by an addendum to the MOA stipulating that the NICD should assist a provincial health department in managing the public health emergency by providing operational support. For this purpose the NICD implemented an Incident Management System (IMS).

Results

The NICD provided support in the form of incident management, financial administration, logistics, human resources, data management and communications. Challenges identified through the activation resulted in the following key lessons learnt: response teams must be flexible, knowledgeable and willing to work within an IMS; decision-makers to be available and involved to manage risks; establishment of clear lines of communication helps define roles and responsibilities; strong leadership within the IMS is required to coordinate teams.

Conclusion

Overall, the NICD satisfied the requirements of the MOA. However, in preparation for a public health emergency of an infectious nature, so as to rapidly reduce morbidity and mortality, it is recommended that universal training of infectious disease control practitioners on IMS and its application be prioritised.

ID: 8530

Category: Sexually Transmitted Diseases (STDSSA)
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ABSTRACT TITLE: PROFILING VAGINAL MICROBIOTA USING SENSITIVE QUANTIFICATION ASSAYS

Introduction

Bacterial vaginosis (BV) is the most common cause of vaginal discharge syndrome affecting up to 60% of women of childbearing age in parts of South Africa. Diagnosis of BV in resource-poor settings is not prioritised and relies on semi-quantitative microscopy algorithms such as Amsel’s criteria and Nugent score (NS). Here, we evaluated a quantitative real-time PCR (qPCR) assay to detect bacterial communities.

Methods

Vaginal swabs from 251 South African women presenting with STI symptoms were evaluated for BV using NS. By qPCR, we analysed DNA from vaginal swabs for six BV-associated bacteria, using ThermoFisher vaginal microbiota panel: Gardnerella vaginalis(GV), Prevotella bivia (PB), BV-associated-bacteria-2 (BVAB2), Megaspheara 1(M-1), Lactobacillus crispatus(LC) and Lactobacillus jensenii (LJ), and determined quantitative bacterial levels. The sensitivity and
specificity of each qPCR assay, compared to NS, was measured using ROC analysis. Cut-offs were calculated based on sensitivities and specificities. A logistic regression model was used to determine the strongest predictors of BV status for all bacterium.

**Results**

Nugent scores from 251 women showed 30.2% normal flora (NS 0–3), 34.2% intermediate flora (NS 4–6) and 36.3% BV (NS > 7). The prevalence of BVAB2, GV, M-1, PB, GBM 46.2%, 68.1%, 31.4%, 47.4% and 60.1%, respectively. Combined titres of GV, BVAB2 and M-1, (GBM) generated a greater Area under curve (AUC), 0.9, compared to individual organisms (0.68, 0.74 and 0.79, respectively). The cut-off, sensitivity and specificity were as follows (listed respectively): BVAB2 (> 0.006; 72%; 73%), GV (> 12.43; 95%; 70%), M-1 (> 0.18; 64%; 92%), PB (> 0.02; 52%; 67%), GBM (> 514; 94%; 77%) (all p < 0.05). Furthermore, an increase in GBM levels (p = 0.034), decrease in LC (p < 0.0001) and LJ (p = 0.026) were identified as the strongest predictors of BV status.

**Conclusion**

We developed an accurate assay, which has the potential to be used as a BV diagnostic and monitoring tool.

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**ID: 8505**

**Category:** Antimicrobial Resistance (SAASP)

**Permission:** Yes

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**ABSTRACT TITLE:** GENOTYPIC CHARACTERISATION OF AMPC β-LACTAMASES AMONG KLEBSIELLA PNEUMONIAE ISOLATED FROM PATIENTS WITH BACTERAEMIA IN SOUTH AFRICA

**Introduction**

AmpCβ lactamases are cephalosporinas produced by Enterobacteriaceae and confer resistance to β-lactam antibiotics. The distribution of plasmid-mediated AmpC resistance in South Africa is unknown. AmpC-producing Klebsiella pneumoniae acquire resistance through the plasmid-mediated ampC gene or hyper-production of an inducible chromosomal AmpC enzyme. Due to limited specificity of phenotypic methods in detecting AmpC enzymes, there is a need to use molecular methods to accurately characterise AmpC-mediated resistance. The aim of the study was to investigate the presence of plasmid-mediated AmpC β-lactamases in multi-drug resistant Klebsiella pneumoniae clinical isolates.

**Methods**

Klebsiella pneumoniae isolates were obtained from sentinel site laboratories as part of a surveillance programme from July 2010–July 2012. Identification and antimicrobial susceptibility testing (AST) patterns of organisms were confirmed on VITEK®II system and the MicroScan®Walkaway system, respectively. Isolates that were non-susceptible to cefoxitin were selected for screening of genes coding for plasmid-mediated AmpC β-lactamase production (blaMOX, blaFOX, blaDHA, blaACC, blaEBC and blaCIT) using a conventional PCR assay.

**Results**

Of 2 774 Klebsiella pneumoniae isolates, 114 (4.1%) were phenotypically resistant to cefoxitin and eligible for AmpC screening. AST results showed resistance to multiple antibiotics, including ceftazidime (94%), cefepime (92%), cefotaxime (89%), cefoxitin (87%), amoxycillin-clavulanate (89%), trimethoprim-sulfamethaxazole (91%), gentamycin (93%) and ciprofloxacin (82%), but high susceptibility to imipenem and meropenem. Plasmid-mediated AmpC β-lactamases were present in 8.8% (10/114) of the isolates, with 5.3% of the resistance mediated by blaCIT (n = 6), and 3.5% by blaDHA (n = 4). A high percentage of isolates did not express the plasmid-mediated AmpC genes tested for (91.2% n = 104).

**Conclusion**

Plasmid-mediated AmpC β-lactamases were present in 8.8% of Klebsiella pneumoniae isolates; these consisted of blaCIT and blaDHA. Further investigations are underway to characterise chromosomal hyper-producers in these isolates.

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**ID: 8510**

**Category:** Antimicrobial Resistance (SAASP)

**Permission:** Yes

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**ABSTRACT TITLE:** PREVALENCE OF CARBAPENEMASE PRODUCING PSEUDOMONAS AERUGINOSA CIRCULATING IN PRETORIA, SOUTH AFRICA

**Background**

Pseudomonas aeruginosa is an important nosocomial pathogen. Carbapenems are used for treating infections caused by multidrug-resistant P. aeruginosa. The extensive use of carbapenems has led to the emergence of carbapenem resistance in this bacterium. Carbapenemases hydrolyse carbapenems efficiently and are considered the most clinically significant mechanism of carbapenem resistance in P. aeruginosa isolates. The aim of this study was to determine the rate of carbapenem resistance and virulence in P. aeruginosa circulating in the Pretoria region.

**Material and Methods**

A total of 237 consecutive non-repeat isolates were collected from the diagnostic division of the Department of Medical Microbiology (TAD, NHLS) from May to September 2016. Identification and susceptibility were determined by the VITEK® automated system. All imipenem-resistant isolates were screened for the presence of metallo-beta-lactamases using the MBL E-test. The following carbapenemases were detected using multiplex PCR assays: Verona integron encoded metallo-beta-lactamase, Oxaclillinase-48, Adelaide imipenemase, German imipenemase, Sao Paulo metallo-beta-lactamase, Imipenem metallo-beta-lactamase, New Delhi metallo-beta-lactamase and Guiana extended spectrum genes and selected virulence genes.

**Results**

Sixty (24%) of the 237 P. aeruginosa isolates collected were reported as imipenem resistant. Fifteen (15/60) (25%) imipenem-resistant isolates were confirmed as MBL positive. The blaVIM gene was detected in eight (53%) of the fifteen MBL positive isolates. The remaining seven (47%) MBL positive isolates harboured the GES gene. The NDM gene was detected in fourteen (93%) of the 15 MBL positive isolates. The OXA-48 and GES genes were detected from the imipenem-resistant isolates with percentages of 22% and 78%, respectively. A high prevalence of quorum-sensing genes i.e. rhl (94%), lasL (93%) and lasR (92%) were detected, while the effector enzyme (exoY) and phenazine (phzI) genes were detected at 0.4%

**Conclusions**

The high prevalence of carbapenem resistant and carbapenemase producing P. aeruginosa isolates found in this study highlights the need for alternative treatment options. Quorum sensing is important for regulating antimicrobial production and biofilm formation, enabling persistence and spread of resistant P. aeruginosa. There is an urgent need for the implementation of active surveillance systems and adequate infection control measures in South African hospitals to prevent and limit the spread of resistant bacteria.
Conclusion
There was a high prevalence of STIs among this group of men, especially in the extra-genital sites. As MSM are at a high risk for STDs, including HIV infection, they should be counselled about repeated STI screening (including extra-genital sites) and safer sex. Public health messages aimed at MSM need to emphasise safe insertive as well as receptive sexual practices.

ID: 8463

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ABSTRACT TITLE: NOVEL MONO, DI AND TRI-FATTY ACID ESTERS BEARING β-ALANINE T-BUTYL ESTER HEAD GROUPS ENHANCE THE TRANSDERMAL DELIVERY OF TENOFOVIR

Introduction
Oral antiretroviral (ARV) drug therapy has significantly improved the treatment of HIV and AIDS. Innovative drug delivery strategies such as novel drug delivery systems and alternate routes of drug administration can overcome the limitations of oral ARV therapy. The use of chemical permeation enhancers (CPEs) has widened the pool of drugs that can be delivered via the transdermal route. This study explored the synthesis and characterisation of novel mono, di and tri-fatty acid (FA) esters bearing β-alanine t-butyl ester head group as permeation enhancers for the transdermal delivery of tenofovir (TFN).

Methods
The novel esters were characterised using FTIR, 13C NMR, 1H NMR and HRMS. The biosafety of the novel FA esters was established using in vitro cell culture. In vitro transdermal permeation studies were performed using rat skin and Franz diffusion cells. Transepithelial electrical resistance (TEER) was used to establish the integrity of the treated skin. Hematoxylin and eosin (H&E) stained images were used to histologically evaluate the treated skin samples.

Results
The synthesised compounds were non-toxic to mammalian cells confirming their safety for pharmaceutical applications. All the synthesised derivatives displayed better transdermal permeation enhancement capabilities as compared to their respective individual FAs. The mono oleate derivative (MOAPE) displayed the greatest ER for TNF (5.87) at 1% w/w. Histological investigations of skin treated with MOAPE revealed fluidisation of the stratum corneum. Histological and TEER studies corroborated with the findings of the in vitro permeation experiments and revealed that there was no significant change to the viable epidermis of the skin after 1% MOAPE exposure. The TEER findings also suggested that the permeation enhancement effects of MOAPE were not permanent, with the results indicating a return towards original skin integrity after removal of the enhancer formulation.

Conclusion
This study reported novel FA esters showing promise as effective permeation enhancers. The findings show that the novel mono ester derivative of OA (MOAPE) adds to the pool of CPEs available to formulation scientists and can be safely incorporated into TNF TDD systems.

ID: 8541

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ABSTRACT TITLE: MOLECULAR CHARACTERISATION OF CHLAMYDIA TRACHOMATIS ISOLATES USING SEQUENCE VARIATION IN THE MAJOR OUTER MEMBRANE PROTEIN GENE (OMP1)

Introduction
We have applied High Resolution Melting Analysis (HRMA) for the genotyping of the Chlamydia trachomatis (CT) and applied
it specifically to the 14 sexually transmitted infection-related genotypes: A-C, D-K, and L1-L3. Based on the genotype of the OMP1 (Outer Membrane Protein) gene CT is grouped into different serovars, which present in different clinical manifestations; with type A, B, Ba, and C causing trachoma, D-K cause urogenital infections and L1, LII & LIII associated with lymphogranuloma venereum (LGV). The OMP1 is one of the conserved genes found in CT. The aim of this study is to evaluate the prevalence of the above mentioned serovars.

Methods

Two hundred and sixty-five Eswab™ clinical samples were screened for CT using Anyplex™ II STI-7 Detection. We confirmed the presence of the OMP1 gene with the conventional PCR. HRMA was performed to identify the CT serovars on a Quantstudio 5 qPCR instrument and CDC controls were included in the analysis. HRM analysis was done on the High-Resolution Melt Software v3.1.

Results

Using HRM we identified the following serovars: A, B, C, D, E, F, G, I, J, L3. The highest prevalent serovars were serovar F = 29%; E = 16.1%; I = 12.9%; D and G both = 9.7% and the rest below 6.5%. No serovars: H, K, L1, L2 were identified. These are the preliminary results which should be validated using sequencing to confirm each genotype.

Conclusion

In conclusion, serovars from D-K in this study dominate, with F being the most prevalent one. This serovar set has been associated with urogenital infections. HRM genotyping could be used to differentiate multiple concurrent urogenital CT infections in the clinical samples.

ID: 8205

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ABSTRACT TITLE: ANTIMICROBIAL RESISTANCE AND ANTIBIOTIC STEWARDSHIP: KNOWLEDGE, ATTITUDES AND PERCEPTIONS AMONGST FINAL-YEAR UNDERGRADUATE HEALTH PROFESSIONAL STUDENTS IN A SOUTH AFRICAN UNIVERSITY

Introduction

Antimicrobial resistance (AMR) is a major public health threat, with the World Health Organization and South African Department of Health identifying the education and training of healthcare professionals on AMR and antimicrobial stewardship (ABS) in the Global Action Plan and National Strategy Framework respectively. This study describes the knowledge, attitudes and perceptions (KAP) of AMR and ABS amongst final year medical, pharmacy and nursing students at a single University in Durban, South Africa.

Methods

The study was a cross-sectional questionnaire-based survey on the KAP of final-year medical, pharmacy and nursing students at a South African University.

Results

A total of 132 questionnaires were completed (response rate 33%), with individual response rates of 63% (n = 63), 86% (n = 46) and 9% (n = 23) for pharmacy, nursing and medical students respectively. The mean correct knowledge score was 88.9%, with significantly lower scores seen for nursing students when compared to the other two groups. The perceived seriousness of AMR at international, national and local levels was significantly lower amongst nursing students. Only a third of all students and 45% of nursing students agreed that use of antibiotics contributes to AMR. Several nursing and medical students reported taking antibiotics for viral illnesses whilst almost a quarter of all students sampled did not consult a doctor before starting an antibiotic.

Conclusion

Several gaps in knowledge were identified, with key differences between the student groups. Attitudes and perceptions also differed substantively indicating the need for curriculum review on AMR and ABS content as suboptimal KAP may lead to negative patient outcomes.

ID: 8380

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The proportion of cross resistance among rifamycins were: across all three (216/300; 72%), between RIF and RFP (292/300; 98%) and to RFB (50/52; 96%).

The proportion of cross-resistance among rifamycins were: across all rifamycins, high levels of cross resistance across all rifamycins, however 28% of MDR/XDR-TB cases could potentially benefit from rifabutin as a substitute drug to the failing regimen. The use of LPA and rpoB mutations specifically S531L and D516V can be beneficial in rapidly differentiating phenotypic susceptibility to rifabutin according to this study.

ID: 8381

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ABSTRACT TITLE: CROSS-RESISTANCE AMONG RIFAMYCINS IN MYCOBACTERIUM TUBERCULOSIS CLINICAL ISOLATES

Introduction

High level cross-resistance among rifamycins in Mycobacterium tuberculosis isolates is commonly inferred. However, previous studies reported that the minimum inhibitory concentrations (MICs) of rifabutin (RFB) among rifampicin (RIF)-resistant M. tuberculosis carrying rpoB mutations varies depending on the mutation position.

Objective

To determine the proportion of cross-resistance among rifamycins and to assess the use GenoType MTBDRplus assay in predicting differential phenotypic susceptibility to rifamycins in M. tuberculosis isolates.

Method

A total of 300 clinical isolates of rifampicin-resistant M. tuberculosis collected between June 2015 to April 2016 during routine surveillance activities for rifampicin drug resistance in high burden districts of South Africa were included. Drug susceptibility testing for RIF, RFB and rifapentine (RFP) was performed using the BD MGIT960 instrument and the Sensitrile MycoTB plate was used to determine MICs for RIF and RFB. To determine rpoB mutations, all the isolates were tested by LPA method and isolates with inconclusive results were sent for rpoB gene sequencing.

Results

The proportion of cross resistance among rifamycins were: across all three (216/300; 72%), between RIF and RFP (292/300; 98%) and RIF and RFB (217/300; 72%). The S531L mutation was the mostly associated with cross resistance to all rifamycins (144/153; 94%), while the D516V mutation associated with differential susceptibility to RFB (50/52; 96%).

Conclusion

The results show high levels of cross resistance across all rifamycins, however 28% of MDR/XDR-TB cases could potentially benefit from rifabutin as a substitute drug to the failing regimen. The use of LPA and rpoB mutations specifically S531L and D516V can be beneficial in rapidly differentiating phenotypic susceptibility to rifabutin according to this study.
DST. Sensitivity, specificity and diagnostic accuracy of MTBDRsl VER 2.0 were determined with BACTEC® MGIT 960 DST as a reference method, using WHO 2013 recommended critical concentrations for DST.

Results

Directly comparable results between LPA and DST were available for 134 clinical samples. Sensitivity, specificity and diagnostic accuracy were calculated and are as follows respectively:

- FLQ (OFX) 76.8% (CI: 63.6–87.0%), 89.7% (CI: 80.8–95.5%) and 84.2%;
- FLQ (MFX), 79.2% (CI: 65.0–89.5%), 84.9% (CI: 75.5–91.7%) and 82.7%;
- SLI (KAN) 78.9% (CI: 67.6–87.7%), 90.3% (CI: 80.1–94.6%) and 84.2%;
- SLI (AMK), 85.9% (CI: 75.0–94.3%), 89.7% (CI: 79.9–95.8%), and 87.9%;
- SLI (CAP) 86.2% (CI: 75.3–93.3%), 91.2% (CI: 81.8–96.7%) and 88.7%.

Conclusion

MTBDRsl VER 2.0 LPA performed well on clinical samples with good sensitivity and specificity for FLQ and SLID. The results are comparable with data published in the WHO 2016 Policy Guidance Document. The inclusion of this assay as an initial direct test in the SA National TB Programme will enable faster diagnosis and laboratory-based treatment options for DR-TB patients.

ID: 8424

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ABSTRACT TITLE: AMIKACIN-RESISTANT ACINETOBACTER SPECIES ARE MEDIATED BY THE APHA6 GENE AT AN ACADEMIC COMPLEX HOSPITAL IN DURBAN, KWAZULU-NATAL, SOUTH AFRICA

Introduction

Drug resistant Acinetobacter species (Acinetobacter spp.) present serious therapeutic and infection control policy challenges globally. Although aminoglycosides have played a crucial role in the treatment of MDR Acinetobacter spp, recent reports indicated that Acinetobacter isolates are developing resistance to aminoglycosides around the globe. This study investigated the minimum inhibitory concentrations (MICs) of amikacin against Acinetobacter spp. and genes associated with its resistance. The association between amikacin resistance and clinical outcomes of patients was also determined.

Method

Clinical information from 107 patients cultured with Acinetobacter spp. was recorded during clinical ward rounds, including clinical outcomes, history of antibiotics prescribed and microbiological investigations. The 107 Acinetobacter spp. were investigated for susceptibility to antimicrobial agents in use at local hospitals. Resistant genes related to amikacin (aphA6 and aacA4 gene) were investigated by polymerase chain reaction (PCR) and sequencing. Analysis was performed on the relationship between clinical outcomes and antimicrobial resistant patterns, as well as on the amikacin MICs in resistant isolates (n = 6) versus their PCR results.

Results

Amikacin resistance was observed in six isolates (5.6%). The MICs were 32 (n = 3) and ≥ 64 µg/mL (n = 3) for the amikacin resistant isolates. The aphA6 gene (797 bp) was detected in all amikacin-resistant isolates. However, the aacA4 gene (489 bp) was not present in these isolates. While the majority [5/6 (83%)] of cases were discharged, mortality rates were [1/6 (17%)]. No underlying clinical factors were significantly associated with clinical outcome.

Conclusion

In this local setting, amikacin is commonly used with piperacillin-tazobactam as a second-line treatment option in general antibiotic policy. The majority, 101 (94.4%) of the tested 107 Acinetobacter spp. isolates were susceptible to amikacin which underscores the crucial role of this antibiotic in the treatment of MDR Acinetobacter spp. All the six amikacin-resistant isolates were extensively drug resistant which is of serious concern.

The study was approved by the Ethics Committee of University of KwaZulu- Natal (Ethic approved: BE 283/12).

ID: 8450

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ABSTRACT TITLE: ANTIMICROBIAL SURGICAL PROPHYLAXIS FOR CAESAREAN SECTION, MAKING THE RIGHT CHOICE THROUGH COLLABORATIVE INTERVENTIONS
Introduction

Surgical site infections (SSI) are the second most common cause of hospital-acquired infections (HAI). SSIs lead to increased cost and poor patient outcomes. Appropriate antibiotic surgical prophylaxis (ASP) has been shown to reduce the occurrence of SSIs. The aim of the study was to measure compliance in terms of drug choice, dose, timing and duration of therapy for C-section.

Methods

One-thousand-and-thirty-nine patient records were reviewed prospectively over a 78-week period in the maternity ward at a private hospital in Gauteng, South Africa. Data collected from patient peri-operative documents were entered onto a standardised Microsoft Excel™ spread sheet and analysed. Problem areas were identified and addressed with individual prescribers and presented to the Antimicrobial Stewardship (AMS) Committee.

Results

Over the first 28 weeks of the study 71% (n = 266/375) of patients reviewed received the right choice of antibiotic, 50% (n = 191/375) correct dose, 7% (n = 26/375) were compliant to the right time prior to incision (30–60 minutes) and 100% (n = 375/375) received duration less than 24 hours. Data analysis revealed ceftriaxone utilisation as ASP. The prescriber was approached by the pharmacist to consider changing practice to a narrower spectrum alternative. This intervention failed. The data was then presented to the hospital AMS committee in week 28, and a decision was taken to remove ceftriaxone from all theatres. As a result, this lead to 100% (n = 460/460) compliance to the appropriate ASP agent measured post intervention. Under-dosing of cefazolin according to weight was also identified and compliance following feedback improved to 69% (n = 321/460). The average time of antibiotic administration prior to incision was 14 minutes.

Conclusion

Further intervention is needed to improve compliance to correct antibiotic dose and appropriate time of administration prior to incision. The AMS committee plays an important role in a hospital to drive improvement and communicate difficulties identified to medical practitioners.
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ABSTRACT TITLE: ANALYSIS OF THE ANTIBIОGRAM FROM AN ORTHOPAEDIC DEPARTMENT AT A TERTIARY HOSPITAL IN JOHANNESBURG

Introduction
Changing patterns of microbial aetiology and increasing antibiotic resistance negatively impact on initiation of early effective treatment of orthopaedic infections. This may result in increased morbidity and the need for additional antibiotics and surgery. A unit-specific antibiogram assists in guiding empiric antibiotic choices.

Methods
For the period 1 July 2015 to 30 June 2017, results of all bacterial pathogens from the orthopaedic department were reviewed.

Results
There were 407 non-duplicate isolates with a similar number of Gram-positive and Gram-negative organisms.

Staphylococcus aureus made up 70% of Gram-positives. This was followed by enterococci (17%). The remaining Gram-positives comprised of streptococci, coagulase-negative staphylococci and anaerobes.

Twenty-three percent of S. aureus isolates were methicillin-resistant. Enterococcus faecalis, all ampicillin-susceptible, was the predominant enterococcal species. There were no vancomycin-resistant enterococci.

The Gram-negatives comprised of Enterobacteriaceae (63%), non-fermenters (34%) and anaerobes (4%).

The predominant Enterobacteriaceae were Enterobacter species, Escherichia coli and Proteus species. Rates of susceptibility to all antibiotic groups varied by genus with Enterobacter species showing the lowest susceptibility rates and Proteus species the greatest. Overall rate of susceptibility to piperacillin-tazobactam, ceftazidime and cefepime for these Enterobacteriaceae was 80%, 81% and 83%, respectively.

The non-fermenters comprised of equal numbers of Pseudomonas aeruginosa and Acinetobacter baumannii. Eighty-five percent of P. aeruginosa isolates were susceptible to ciprofloxacin and 91% to amikacin. Of the β-lactams, piperacillin-tazobactam and cefepime demonstrated the greatest activity.

Overall A. baumannii isolates showed very low rates of susceptibility to all antibiotic groups. Sixty-six percent were multidrug-resistant (MDR).

Conclusion
Tracking and reporting of resistance patterns is an important component of antibiotic stewardship. Antibiograms have many uses, including being a valuable antibiotic-prescribing tool. The antibiogram must be used in conjunction with clinical (infection type: septic arthritis, osteomyelitis, implant or surgical site infection) and epidemiological (community- or hospital-acquired infection and individual patient risk factors for MDR infection) data.

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ABSTRACT TITLE: THE EVALUATION OF THE APPROPRIATE USE OF ECHINOCANDINS FOR THE TREATMENT OF INVASIVE CANDIDIASIS IN A PRIVATE HOSPITAL SETTING

Introduction
Invasive candidiasis (IC) is a major contributor to morbidity and mortality in non-neutropenic patients in the ICU setting. Echinocandins are the first choice of therapy for IC but their appropriate use remains a challenge. The aim of this study was to evaluate the appropriate use of echinocandins utilising the START Candida Score and the (1–3)-β-D-Glucan Assay (Fungitell assay).

Methods
This retrospective study was conducted between October 2016 and June 2017 in a private hospital. All patients who received an echinocandin, as well as their potential risk factors for IC were identified using the electronic billing database. Laboratory data were collected from the antibiotic stewardship and infection prevention database. Total parenteral nutrition (TPN), surgery, multifocal Candida colonisation and severe sepsis were coded as ‘0’ if absent, and present ‘1’ (severe sepsis coded ‘2’). A Candida score of ≥ 3 selected patients at high risk for invasive candidiasis.

Results
Thirty patients received an echinocandin. Thirteen (43%) patients had a positive Candida score ≥ 3 of which seven (53%) had a positive blood culture (BC). Two patients with a negative Candida score had a positive BC. The two most prevalent Candida species isolated from blood cultures were Candida parapsilosis (55%) and Candida albicans (33%). A Fungitell assay was performed in four patients of whom only one was positive. The mean duration of echinocandin therapy was 12.9 days. Repeat blood cultures were only performed on two patients.
Conclusion

More than half of the patients (57%) received an echinocandin with a negative START Candida score. The Fungitell assay, as an adjunctive diagnostic tool, was greatly underutilised. Duration of therapy was not guided by follow-up blood cultures. The focus point for the antimicrobial stewardship committee going forward is to promote awareness and correct implementation of current guidelines regarding the diagnosis and management of IC.

ID: 8478

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ABSTRACT TITLE: ANTIMICROBIAL SUSCEPTIBILITY PROFILE OF STAPHYLOCOCCUS AUREUS FROM ORTHOPAEDIC INFECTIONS

Introduction

Staphylococcus aureus is the most commonly isolated microorganism in osteomyelitis, and more than 30% of these isolates may be methicillin resistant Staphylococcus aureus (MRSA). A study of S. aureus bacteraemia in Johannesburg showed a MRSA rate of 23%. Drugs such as trimethoprim-sulfamethoxazole and rifampicin may be used as suppressive therapy in chronic osteomyelitis. Infections caused by MRSA respond poorly to empiric cloxacillin therapy. This study aimed to identify the resistance profile of S. aureus from orthopaedic infections in order to guide management.

Methods

The study included 84 consecutive, non-duplicate Staphylococcus aureus isolates obtained from pus swabs taken in the Orthopaedic ward at King Edward VIII Hospital. Data was extracted retrospectively from the laboratory electronic database for the period 1 January to 31 December 2016. All swabs and isolates were processed according to standard laboratory operating procedure.

Results

77/84 (93%) of Staphylococcus aureus isolates were susceptible to cloxacillin; 76/84 (90%) to clindamycin; 75/84 (89%) to erythromycin; and 82/84 (98%) to moxifloxacin. All isolates were susceptible to vancomycin, linezolid and teicoplanin. Of the seven methicillin-resistant staphylococcus aureus isolates, 43% were susceptible to clindamycin, 29% susceptible to erythromycin, 14% susceptible to trimethoprim-sulfamethoxazole, 71% susceptible to moxifloxacin, 100% susceptible to vancomycin, teicoplanin, tigecycline and linezolid. Of note none of the MRSA isolates were susceptible to ciprofloxacin, rifampicin and tetracycline. The average age of patients was 34.1 years.

Conclusion

Cloxacillin is a reasonable empiric choice for the treatment of staphylococcal infections, however, drugs such as trimethoprim-sulfamethoxazole and rifampicin, should not be used without susceptibility results. Specimens should be taken and therapy guided by culture results as there are serious consequences if resistance is missed and infection progresses. Vancomycin remains a good option for the treatment of MRSA.

ID: 8484

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ABSTRACT TITLE: KLEBSIELLA PNEUMONIAE PREVALENCE AND ANTIMICROBIAL SUSCEPTIBILITY PROFILE: TELLING THE DR GEORGE MUKHARI ACADEMIC HOSPITAL’S STORY.

Introduction

Both locally and globally antimicrobial resistance is a general public health concern which threatens the effective treatment of severe infections. Klebsiella pneumoniae is the most commonly encountered Gram-negative bacteria globally. It has been implicated in outbreaks with high mortality in neonatal and high-care units. This study was aimed at determining the prevalence and antimicrobial susceptibility profile of K. pneumoniae isolated from blood culture specimens taken at selected high-care wards of Doctor George Mukhari Academic Hospital in 2015.

Methods

Data on microbiology results of 2015 was obtained from the NHLS laboratory data information system (TrakCare) using Microsoft Excel. Blood culture results from high-care wards were filtered, double checked for repeats and categorised according to isolated microorganisms. A description of the susceptibility trends to selected antimicrobial agents commonly used against the organism in osteomyelitis, and more than 30% of these isolates may be methicillin resistant staphylococcus aureus isolates were susceptible to ciprofloxacin, rifampicin and tetracycline. The average age of patients was 34.1 years.

Conclusion

Cloxacillin is a reasonable empiric choice for the treatment of staphylococcal infections, however, drugs such as trimethoprim-sulfamethoxazole and rifampicin, should not be used without susceptibility results. Specimens should be taken and therapy guided by culture results as there are serious consequences if resistance is missed and infection progresses. Vancomycin remains a good option for the treatment of MRSA.

ID: 8484

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ABSTRACT TITLE: KLEBSIELLA PNEUMONIAE PREVALENCE AND ANTIMICROBIAL SUSCEPTIBILITY PROFILE: TELLING THE DR GEORGE MUKHARI ACADEMIC HOSPITAL’S STORY.

Introduction

Both locally and globally antimicrobial resistance is a general public health concern which threatens the effective treatment of severe infections. Klebsiella pneumoniae is the most commonly encountered Gram-negative bacteria globally. It has been implicated in outbreaks with high mortality in neonatal and high-care units. This study was aimed at determining the prevalence and antimicrobial susceptibility profile of K. pneumoniae isolated from blood culture specimens taken at selected high-care wards of Doctor George Mukhari Academic Hospital in 2015.

Methods

Data on microbiology results of 2015 was obtained from the NHLS laboratory data information system (TrakCare) using Microsoft Excel. Blood culture results from high-care wards were filtered, double checked for repeats and categorised according to isolated microorganisms. A description of the susceptibility trends to selected antimicrobial agents commonly used against the organism in osteomyelitis, and more than 30% of these isolates may be methicillin resistant staphylococcus aureus isolates were susceptible to ciprofloxacin, rifampicin and tetracycline. The average age of patients was 34.1 years.

Conclusion

Cloxacillin is a reasonable empiric choice for the treatment of staphylococcal infections, however, drugs such as trimethoprim-sulfamethoxazole and rifampicin, should not be used without susceptibility results. Specimens should be taken and therapy guided by culture results as there are serious consequences if resistance is missed and infection progresses. Vancomycin remains a good option for the treatment of MRSA.
100%, imipenem: 99.7% and ertapenem: 99.7%. There was a mixed level of susceptibility and resistance ratio to other agents: 64% to 36% for amoxicillin-clavulanic acid and 84% to 16% for ciprofloxacin. Out of the 359 K. pneumoniae isolated, 109 (30%) were extended spectrum beta lactamase (ESBL) producers.

Conclusion
The level of resistance to antimicrobial agents indicated in this report affirms the need for an ongoing surveillance system that gives regular updates on antimicrobial resistance trends and to aid the update of antimicrobial policies within DGMAH.

ID: 8502
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ABSTRACT TITLE: MANAGEMENT OF UNCOMPLICATED URINARY TRACT INFECTIONS IN THE EMERGENCY DEPARTMENT: A POTENTIAL OVERLOOKED CONTRIBUTOR TO ANTIBIOTIC RESISTANCE

Introduction
Uncomplicated urinary tract infections (UTIs) are one of the most common infections in the primary healthcare setting, more commonly in adult non-pregnant women. Approximately 20% of all antibiotic prescriptions issued by general practitioners are for patients with uncomplicated UTIs and treated empirically without conducting a urine culture or susceptibility testing. This is of concern as Escherichia coli (E. Coli), which is the most prevalent organism isolated from urine cultures, has an emerging increase in resistance to quinolones.

The study aimed to assess the empiric choice of antibiotics prescribed for an uncomplicated UTI in the emergency department and to determine whether urine cultures were requested, the most prevalent organism and its sensitivity pattern.

Methods
A retrospective record review of patients that consulted in the emergency department with suspected UTI in the time period of 1 January 2014 until 29 of February 2016 was conducted. Patients admitted to the hospital, complicated UTIs and recurrent UTIs were excluded.

Results
Three hundred records were reviewed. The majority of the study population was female (88%), with a mean age of 33.3 years. Urine cultures were requested in only 56% (n = 168) of the patients presenting with signs and symptoms of an uncomplicated UTI. Cultures were positive in 77% (n = 131) of cases and EColi was the most prevalent organism isolated (38.82%, n = 100), with Klebsiella pneumonia accounting for 4.71% (n = 8) of the isolates.

Ciprofloxacin was prescribed in the majority of cases (37.9%), followed by amoxicillin/clavulanate (29%); 23.3% (n = 70) of patients received two or more antibiotics, either in class of antibiotic or route of administration, with n = 54 of these patients receiving intravenous antibiotics.

Conclusion
Antibiotic stewardship principles need to be applied in the emergency department, with special emphasis on double antimicrobial cover as well as establishing an ethos to perform cultures prior to antibiotic prescribing.

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**ABSTRACT TITLE:** PREVALENCE OF THE RIFAMPICIN RESISTANT DETERMINANT AT CODON 491 IN SOUTH AFRICA, UNDETECTED BY COMMERCIAL PHENOTYPIC AND GENOTYPIC METHODS

**Introduction**

The commercial WHO-endorsed phenotypic and molecular assays are incapable of detecting rifampicin resistance in *Mycobacterium tuberculosis* clinical strains harbouring the resistance conferring Ile491Phe mutation found outside the Rifampicin Resistance-Determining Region (RRDR). Of concern to South Africa, is a publication showing that 30% of MDR-TB cases in Swaziland, partially nestled within South Africa, harbour this mutation. We aimed to determine the prevalence of this mutation by retrospective analysis of Whole Genome Sequenced isolates representing two provinces, in close proximity to Swaziland, from the 2014 National TB Drug Resistance Survey (DRS).

**Methods**

Survey isolates from two provinces were sequenced using Next-Generation Whole Genome Sequencing methodology. The provinces included Gauteng, the financial hub which attracts foreign migrants, and KwaZulu-Natal, the province with the highest burden of TB disease bordering Swaziland's eastern perimeter. Isolates were further subjected to drug susceptibility testing on the MGIT960 to first and second-line anti-tuberculosis drugs.

**Results**

Of the 1 535 genomes sequenced from the TB DRS, 1 (0.07%) isolate (Gauteng) harboured the Ile491Phe as a minor variant (2% frequency). This isolate was fully susceptible to both first- and second-line drugs. The crude unadjusted prevalence of resistance conferring mutations within the RRDR occurred at 5.15% (79/1535). Inclusion of the Ile491Phe mutation adjusts the prevalence to 5.21% (80/1535).

**Conclusion**

Despite the prevalence of this mutation at high rates in Swaziland, the prevalence was only found to be 0.07% in the two provinces or 0.12% when considering Gauteng only. Due to the clinical implications of not detecting this mutation, surveillance activities have been enhanced to improve detection of this mutation. It would be impractical to screen all isolates based on the total number of TB diagnosed annually in South Africa, therefore, clinicians should bear in mind the occurrence of this mutation in cases with poor clinical outcomes despite adherence to therapy.

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**ABSTRACT TITLE:** PREVALENCE OF MINORITY HIV-1 DRUG RESISTANT MUTATIONS IN CHILDREN WITH VIROLOGIC FAILURE IN A RURAL KWAZULU-NATAL COHORT

**Introduction**

Missed minority drug resistance mutations (DRMs) may pave a way to therapy failure in a short period of time. Therefore, assays have been developed including next generation sequencing (NGS) that are able to identify a larger proportion of mutations including those bearing minority DRMs within a patient’s viral population. The aims of this study were to (1) describe the prevalence of minority HIV DRMs in paediatric population with virologic failure in a rural KwaZulu-Natal (KZN) paediatric cohort using NGS technology, and (2) to compare the genotypes generated using Sanger sequencing with NGS.

**Methods**

Thirty-four patients were genotyped using Sanger sequencing and NGS. A 1.3 kb region of the Pol gene was genotyped using Sanger sequencing. The whole 9.7kb HIV genome was sequenced using NGS. All electropherograms were analysed using the Geneious V8.0.5 software system for the presence of DRMs including minority variants. Sequences were assembled against an HIV-1 subtype C reference sequence from South Africa. For NGS a reference sequence was annotated with known HIV resistance mutations within the protease and RT genes.

**Results**

NGS was able to detect minority drug resistance mutations in eleven (32.3%) samples which were missed by Sanger sequencing. NGS also detected an additional three (8.8%) specimens that harboured DRMs but were found to be susceptible by Sanger sequencing.

**Conclusion**

The presence of minority DRMs among children is likely to obstruct the use of ART and consequently predispose patients to therapy failure. We noted that children on PI-based regimen, while at a lower prevalence, still harboured DRMs that remained undetected.
by conventional Sanger sequencing. Finally, this study emphasised the need to apply more sensitive assays to accurately distinguish patients failing due to the emergence of minority DRMs from those that are non-adherent in order to maximise the efficacy of the limited range of anti-retroviral drugs currently in use in South Africa.

**ID: 8237**

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**ABSTRACT TITLE:** PREVALENCE OF HUMAN PAPILLOMAVIRUS INFECTIONS IN ANORECTAL CANCERS AT THE DR GEORGE MUKHARI ACADEMIC HOSPITAL FROM 2005 TO 2012  

**Introduction**  
Human papillomavirus (HPV) accounts for 94% of anorectal cancer cases globally. The study aimed at investigating the prevalence of HPV-genotypes in anorectal cancers and reports on the prevalence of HPV-infections and HPV-genotypes in different histological sites of anorectal cancers.  

**Methods**  
HPV-DNA from 148 formalin-fixed paraffin-embedded (FFPE) tissue samples was extracted using NucliSENS® easyMAG® following manufacturer’s protocol with an off-board pre-lysis step. All DNA extracts were screened for HPV-DNA with a nested-PCR assay (MY/ GP primer-set) targeting the L1-gene. Genotyping was carried out using Linear Array HPV genotyping assay. All HPV-positive samples were tested for HPV-E6/E7 mRNA expression using APTIMA HPV Assay. Descriptive data analysis on Epi Info version 7.1.5 was then used to calculate frequencies of categorical data.  

**Results**  
Overall HPV prevalence of 23/148 (16%) was found. Of these, the majority 14/23 (61%) were males compared to 9/23 (35%) females. Only eight HPV genotypes (HPV-6, 11, 16, 18, 33, 82, 84, and 66) were detected. Non-oncogenic HPV-6 was predominant in 8/23 (35%). Adenocarcinoma of the rectum region was frequently observed in 133/148 (90%). Anorectal cancer was more prevalent in patients older than age 50 (mean age 54.4; std. Dev 17.9). There was only 1/23 (4%) of the samples (HPV-33) that expressed HPV-E6/E7 mRNA.  

**Conclusion**  
Non-oncogenic HPV-types observed in the majority of adenocarcinoma of the rectum may not necessarily imply any aetiological role in anorectal cancer development. Since the majority of the advanced-aged population present with higher anorectal cancer prevalence, caution should be taken in younger-aged population to prevent cancer development at a later life-stage, especially in males.

**ID: 8340**

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**ABSTRACT TITLE:** ARBOVIRUS LABORATORY INVESTIGATION OF A CLUSTER OF CASES PRESENTING WITH FEVER AND RASH IN JOHANNESBURG, 2017  

**Introduction**  
An investigation into generalised maculopapular rash reported by dermatologists in a cluster of cases in Johannesburg, was initiated by the Outbreak Response Unit of the NICD at the end of January 2017. Arthropod-borne virus (arbovirus) infection was considered as a possible differential diagnosis based on clinical presentation of the patients and the season.  

**Methods**  
Major endemic arboviruses were included in the laboratory investigation at the Centre for Emerging and Zoonotic Diseases: West Nile, Sindbis, Chikungunya and Rift Valley fever. Differential laboratory diagnosis included a battery of assays targeting specific antibodies (haemagglutination inhibition assay, ELISA, virus neutralisation test), viral RNA (PCR, sequencing) and infectious virus by attempted isolation in suckling mice or in tissue culture.  

**Results**  
A total of 47 suspected cases were referred to the NICD for testing. IgM antibodies against Sindbis virus were detected in 27 serum samples, indicative of recent Sindbis virus infection. In addition, recent infection was confirmed in 11 patients by the testing of paired sera, taken two weeks apart to detect a significant rise in antibody titre. Sindbis virus RNA could not be detected in any of the patients, presumably due to blood specimens taken post viremic stage of infection – Sindbis virus is causing typically transient and low viraemia.  

**Conclusion**  
Sindbis virus is a mosquito-borne arbovirus which is maintained in a mosquito-bird transmission cycle. Sindbis virus infection in humans has a short incubation period of less than seven days and is usually a mild and self-limiting febrile illness, but chronic joint manifestations may occur in some cases. The cluster of cases was rare compared
to the number of Sindbis cases usually reported in the past. This is, however likely a reflection of the enhanced passive surveillance and the increased awareness of rash cases reported by dermatologists from January 2017.

**ID: 8581**

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**ABSTRACT TITLE:** EVALUATION OF COBAS E601 SYPHILIS ASSAY FOR THE DETECTION OF SYPHILIS

**Introduction**

*Treponema pallidum* is a causative agent of syphilis infections, which can lead to severe complications if not diagnosed and treated rapidly. The diagnosis as well as screening of syphilis often require assays with high specificity and sensitivity. The Cobas E601 assay is an automated immunoassay which detects antibodies against *T. pallidum*. The test is intended as an aid in the diagnosis of syphilis infection with total assay duration of about 20 minutes. This study was done to evaluate the diagnostic performance of E601 Syphilis cobas assay for the detection of syphilis by comparing with Serodia TPPA assay.

**Methods**

A total of 67 human residual serum samples that were sent for routine laboratory screening for syphilis were collected. Performance of Cobas E601 Syphilis assay was evaluated by determining its sensitivity and specificity compared to Serodia TPPA assay.

**Results**

Of 67 samples tested, 12/67 (18%) were found to be reactive using the Cobas E601 Syphilis assay and 55/67 (82.0%) were found to be nonreactive. The overall sensitivity and specificity of the Cobas E601 Syphilis assay as compared to the Serodia TPPA assay was 98.2% and 100%, respectively.

**Conclusion**

Cobas E601 Syphilis assay demonstrated good diagnostic performance when used to detect syphilis from serum or plasma samples. It can be used as an alternative to Serodia TPPA assay. The assay also has an advantage of less hands-on time and shorter TAT.

**ID: 8464**

**Category:** Sexually Transmitted Diseases (STDSSA)

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**ABSTRACT TITLE:** THE EFFECT OF ANTIMICROBIALS USED FOR GENITAL DISCHARGE DISEASE ON TRICHOMONAS VAGINALIS

**Introduction**

Trichomoniasis is the most common sexually transmitted infection caused by the protozoan, *Trichomonas vaginalis*. *T. vaginalis* infection is often asymptomatic. This infection causes vaginal discharge in women and urethritis in men. It has been reported that trichomoniasis is associated with serious health complications and it increases the risk of HIV acquisition and transmission. Metronidazole has been the standard treatment for trichomoniasis. Multiple cases of metronidazole-resistance have been reported since 1962. Syndromic management of STIs is used to treat multiple infections simultaneously based on the signs and symptom with which the patient presents. In South Africa, the vaginal discharge syndrome is managed with ceftriaxone, azithromycin and metronidazole.

**Methods**

Ten *T. vaginalis* isolates were tested. Each isolate was tested against six combinations of two antimicrobials by the checkerboard method, four combinations of three antimicrobials and two combinations of four antimicrobials. The results obtained from the checkerboard of two antimicrobials were used to design the experiments for three and four antimicrobials combinations.

**Results**

The MICs for metronidazole ranged between 0.25–1 µg/ml and for doxycycline, they ranged between 64–128 µg/ml. Ceftriaxone and fluconazole showed no antitrichomonal activity. All combinations tested had an indifferent effect. However, in the combination of metronidazole and doxycycline a decrease in the MICs for these antibiotics was observed.

**Conclusion**

Combinations of metronidazole and antimicrobials used in syndromic management including fluconazole has no effect against *T. vaginalis*. The decrease in MICs of metronidazole and doxycycline in their combination suggests that these two antimicrobials have
HIV infected can be challenging to heal. This case along with more referral to a competent tertiary centre. Chronic HSV ulcers in the failure of syndromic management of genital ulcers should prompt Discussion exudating lesion, which had grown to > 20 cm², with a new satellite a PI based regimen due to failure. High dose IV acyclovir for two positive on PCR and histology. Her antiretrovirals were changed to with a recurrence of a similar lesion on her right labia minora, HSV-2 lesion, histologically HSV, which was excised. She presented in 2008 Durban, in 2007 with a six-month history of non-healing peri-anal A female HIV positive patient presented to King Edward VIII hospital, Case report genital ulcer disease. Herpes simplex virus (HSV) infection is common in HIV-positive populations and can cause debilitating chronic ulcers. A short course of acyclovir is part of the national treatment guidelines for genital ulcer disease. Case report

A female HIV positive patient presented to King Edward VIII hospital, Durban, in 2007 with a six-month history of non-healing peri-anal lesion, histologically HSV, which was excised. She presented in 2008 with a recurrence of a similar lesion on her right labia minora, HSV-2 positive on PCR and histology. Her antiretrovirals were changed to a PI based regimen due to failure. High dose IV acyclovir for two weeks, followed by three weeks of IV foscarnet failed to shrink the exudating lesion, which had grown to > 20 cm², with a new satellite lesion developing on right perineum. Treatment with topical 1% cidovovir cream reduced the amount of exudation and led to re-epithelialisation of a small area. Unfortunately, the donated stock ran out and topical cidovovir had to be stopped. The lesion continued to grow over the next weeks. Off-label use of topical imiquimod lead to marked improvement within two weeks and near complete healing over three months.

Unfortunately, due to shortage of stock on recurrence, imiquimod could not immediately be resumed and her lesion grew larger. When restarting topical imiquimod, she managed to auto-inoculate her left thumb with herpetic whitlow. Both lesions were scraped and growth of HSV 2 obtained in cell culture. Both isolates were extremely acyclovir resistant. Over the course of several months both lesions responded to imiquimod.

Discussion Failure of syndromic management of genital ulcers should prompt referral to a competent tertiary centre. Chronic HSV ulcers in the HIV infected can be challenging to heal. This case along with more than five others treated in the King Edward VIII hospital Infectious diseases unit, illustrate the success with topical imiquimod.

ID: 8551
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ABSTRACT TITLE: CHRONIC HSV-2 ULCER IN HIV+ WOMAN FROM GENITAL AREA TO FINGER

Introduction
Herpes simplex virus (HSV) infection is common in HIV-positive populations and can cause debilitating chronic ulcers. A short course of acyclovir is part of the national treatment guidelines for genital ulcer disease.

Case report
A female HIV positive patient presented to King Edward VIII hospital, Durban, in 2007 with a six-month history of non-healing peri-anal lesion, histologically HSV, which was excised. She presented in 2008 with a recurrence of a similar lesion on her right labia minora, HSV-2 positive on PCR and histology. Her antiretrovirals were changed to a PI based regimen due to failure. High dose IV acyclovir for two weeks, followed by three weeks of IV foscarnet failed to shrink the exudating lesion, which had grown to > 20 cm², with a new satellite lesion developing on right perineum. Treatment with topical 1% cidovovir cream reduced the amount of exudation and led to re-epithelialisation of a small area. Unfortunately, the donated stock ran out and topical cidovovir had to be stopped. The lesion continued to grow over the next weeks. Off-label use of topical imiquimod lead to marked improvement within two weeks and near complete healing over three months.

Unfortunately, due to shortage of stock on recurrence, imiquimod could not immediately be resumed and her lesion grew larger. When restarting topical imiquimod, she managed to auto-inoculate her left thumb with herpetic whitlow. Both lesions were scraped and growth of HSV 2 obtained in cell culture. Both isolates were extremely acyclovir resistant. Over the course of several months both lesions responded to imiquimod.

Discussion Failure of syndromic management of genital ulcers should prompt referral to a competent tertiary centre. Chronic HSV ulcers in the HIV infected can be challenging to heal. This case along with more than five others treated in the King Edward VIII hospital Infectious diseases unit, illustrate the success with topical imiquimod.
expression of HPV mRNA. Overall HPV mRNA was present in 69.9% of all the HPV DNA samples.

**Conclusion**

Almost half the patients attending gynaecology clinics at DGMAH harbour HPV. HPV mRNA expression was detected in a high number of these samples, these may indicate an increased risk of patients to develop pre-cancerous cervical lesions.

**ID: 8564**

Category: Sexually Transmitted Diseases (STDSSA)

Permission: Yes

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Conclusion

These data suggest that STI-associated inflammation may be eased on STI clearance. However, further studies are warranted to identify other factors driving genital inflammation in the absence of STIs.

**ID: 8051**

Category: Antimicrobial Resistance (SAASP)

Permission: Yes

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**ABSTRACT TITLE**

**MULTILOCUS SEQUENCE TYPING OF CARBAPENEM-RESISTANT PSEUDOMONAS AERUGINOSA ISOLATES FROM PATIENTS PRESENTING AT PORT ELIZABETH HOSPITALS, SOUTH AFRICA**

**Background**

*Pseudomonas aeruginosa* is an important nosocomial pathogen that exhibits multiple drug resistance with increasing frequency, especially to carbapenems making patient treatment difficult. Carbapenem-resistance may be caused by porin gene mutations, active drug efflux, and carbapenemase production. This study evaluated the incidence of genes responsible for carbapenemase production in carbapenem-resistant *Pseudomonas aeruginosa* and assessed the genetic relatedness of the isolates by multi-locus sequence typing (MLST).

**Materials and Methods**

Identification and antimicrobial susceptibility testing of *P. aeruginosa* isolates (n = 234) by the VITEK 2 system detected 81 carbapenem-resistant *P. aeruginosa* isolates. PCR and DNA sequencing were used to screen isolates for three metallo-β-lactamase encoding genes. MLST included amplification of seven housekeeping genes and sequence type alignment using the online *P. aeruginosa* MLST database.
Results

Only the blaVIM-2 gene was detected in 15 of the 81 carbapenem-resistant isolates. MLST indicated six different novel sequence types among the blaVIM-2 positive P. aeruginosa isolates with the majority of the isolates (9/15) containing identical allelic profiles of the sequence type allocated ST1 ( provisionally assigned sequence type, awaiting addition of new sequence types to PubMLST database). Five of these ST1 isolates were from patients and an environmental sample in the same hospital ward suggesting an environmental reservoir. Carbapenem-resistance in the blaVIM-2 negative isolates may be due to other mechanisms.

Conclusion

The incidence of genes responsible for carbapenemase production in carbapenem-resistant Pseudomonas aeruginosa and genetic relatedness of these isolates in public healthcare facilities within the Port Elizabeth area is of concern and requires further investigation.

ID: 8275

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ABSTRACT TITLE: EVALUATION OF RESISTANCE PATTERN OF VARIOUS CANDIDA SPECIES

Introduction

Because of the increased incidence of resistant strains of Candida infections, to the antifungal agents, it is imperative to know the effective antifungal susceptibility pattern in a timely manner for the treatment of patients. The purpose of this study was to determine antifungal resistant pattern among different Candida species so that appropriate antifungals could be administered to the patients with Candida infections.

Methods

A total of thirty-five individual isolates of various Candida species isolated from blood and body fluid specimens of patients admitted to the University of Texas Medical Branch, Texas from November 1, 2012 to April 30, 2013 in were included in this study. All the isolates were subjected for Epsilometer Test (Etest). The minimum Inhibitory Concentration (MIC) end points were determined for four antifungal agents such as fluconazole, itraconazole, voriconazole, and micafungin.

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ABSTRACT TITLE: COMPARISON OF THE MODIFIED CARBAPENEM INACTIVATION METHOD (MCIM) WITH THE RAPIDEC® CARBANP TEST (BIOMERIEUX, FRANCE) FOR THE PHENOTYPIC CONFIRMATION OF CARBAPENEMASE PRODUCTION IN ENTEROBACTERIACEAE

Introduction
The rapid laboratory detection of carbapenemase producing Enterobacteriaceae (CPE) is of utmost importance to guide both treatment and infection control interventions. Various phenotypic confirmatory tests have been developed to aid this. In this study we have compared the performance of the modified carbapenem inactivation method (mCIM) and the Rapidec® CarbaNP test (bioMérieux, France).

Methods
A total of 68 stored Enterobacteriaceae were evaluated using the mCIM as outlined in document M100-S27 of the Clinical and Laboratory Standards Institute (CLSI) guidelines as well as the Rapidec® CarbaNP test according to manufacturer instructions. These 68 isolates were stratified as follows: 39 genotypically confirmed CPE (5 NDM, 8 OXA-48-like, 8 KPC, 10 VIM and 8 IMP) and 29 ESBL and AmpC producing Enterobacteriaceae. The sensitivity and specificity of the two tests under evaluation were determined.

Results
The sensitivity and specificity of the mCIM test was 97.4% and 100% respectively. The OXA 163 positive isolate was not detected by the mCIM method. The sensitivity and specificity of the Rapidec® CarbaNP test was 84.6% and 100% respectively. Five of the eight OXA-48-like positive isolates (OXA-181, OXA-232, OXA-370, OXA-244, and OXA-163) were not detected by the Rapidec® CarbaNP test. The Rapidec® CarbaNP test also failed to detect the VIM 23 positive isolate.

Conclusion
The mCIM test performed satisfactorily for the phenotypic confirmation of carbapenemase production. The Rapidec® CarbaNP test had poor sensitivity (84.6%) for the detection of carbapenemase production, in addition the sensitivity for the detection of OXA-48-like producing Enterobacteriaceae was 37.5%, making it unsuitable for routine use in settings with high prevalence of OXA-48 like producers. The specificity of both tests was 100%. The low cost and ease of performance of the mCIM test combined with the excellent sensitivity and specificity makes it ideal for the detection of CPE in our resource-constrained setting.

ID: 8421
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ABSTRACT TITLE: IMPROVING ANTIBIOTIC UTILISATION IN SURGERY WARDS

Introduction
The discovery of antibiotics has transformed modern medicine by playing a critical role in the management of infectious diseases. However, rapid development of resistance by bacteria is gradually affecting this initial success. Antimicrobial stewardship (AMS) programmes along with infection and prevention control measures have been shown to reduce the burden of antimicrobial resistance in hospitals. Evidence of the impact of AMS programmes in a surgical setting is limited.

Methods
An AMS ward round was implemented in two surgical wards of a tertiary South African hospital with two stages – a retrospective baseline and a prospective intervention stage. The appropriateness of antibiotic utilisation was determined using a guideline developed by Gyssens and colleagues. Additionally, the volume of antibiotic consumption was determined by DDDs/1 000 patient's bed-days.

Results
In both stages of the study amoxicillin/clavulanic acid was the most frequently used agent. A significant reduction in the duration of antibiotic therapy for two days and more was observed from 4.74 ± 4.58 days in the baseline stage compared to 3.96 ± 2.04 days in the intervention phase (p = 0.001). There was a significant reduction in the volume of antibiotic consumption from a total of 739.30 DDDs/1 000 patient days in the baseline stage to 564.93 DDDs/1 000 patient days in the intervention period (p = 0.038). There was a significant reduction of inappropriate antibiotic utilisation from 35% in the baseline stage to 26% in the intervention stage (p = 0.006). This was associated with a reduction in the average antibiotic cost in one of the study wards. Gram-negative bacteria were the most prevalent pathogens in both stages of the study.

Conclusion
Implementation of an AMS ward round showed an improvement in the appropriateness of antibiotic utilisation, along with reduction in consumption and cost of antibiotics in an in-patient surgical setting.
ID: 8448

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**ABSTRACT TITLE:** CHARACTERISATION OF INHIBITOR RESISTANT TEMONIERA (IRT) PRODUCING STRAINS OF ESCHERICHIA COLI AND KLEBSIELLA PNEUMONIAE IN PRETORIA, SOUTH AFRICA

Introduction

Beta-lactam/Beta-lactamase inhibitors are extensively used in the treatment of various infections both in the hospital and outpatient settings. Inhibitor resistance may lead to clinical failures in these patients. The aim of this study was to characterise Inhibitor Resistant Temoniera (IRT) producing strains of *Escherichia coli* and *Klebsiella pneumoniae* in Pretoria, South Africa.

Methods

Sixty non-repeat isolates of *Escherichia coli* and *Klebsiella pneumoniae* demonstrating inhibitor resistance on the Vitek®2 automated system were collected from 1 December 2013 to 31 July 2015. Manual susceptibility testing for amoxicillin-clavulanate was performed using the Kirby Bauer disk diffusion method (CLSI M100S25) and an E-test (bioMérieux), whilst only the Kirby Bauer disk diffusion method (CLSI M100S25) was used for piperacillin-tazobactam. Genomic DNA was extracted using an automated extraction method. A conventional multiplex PCR assay was performed to detect the Temoniera (TEM), Sulfhydryl Variable (SHV), and Oxacillinase (OXA) genes as described previously. The identity of the amplicons was confirmed using Sanger sequencing.

Results

Twenty-five (42%) of the isolates showed a categorical agreement between the E-test and Vitek®2 susceptibility results for amoxicillin-clavulanate. The categorical agreement between Vitek®2 and disk diffusion testing for piperacillin-tazobactam was found to be 47% (28/60). None of the beta-lactamase genes were detected in thirteen (22%) of the isolates. The TEM gene was detected in 47 (78%) of the isolates. Sequencing confirmed the TEM genes as TEM-1.

Conclusion

Most of the strains in this study revealed the presence of the TEM-1 gene. No Inhibitor Resistant Temoniera genes were detected in the isolates. Poor categorical agreement (< 50%) was identified between the Vitek®2 automated system antimicrobial susceptibility testing and manual susceptibility testing. This research gives us some insight into the local epidemiology and resistance mechanisms involved in inhibitor resistance in the Pretoria region.

ID: 8487

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**ABSTRACT TITLE:** COMPARISON OF MANUAL VS. RAPID AUTOMATED LABORATORY METHODS FOR THE IDENTIFICATION OF ANAEROBIC BACTERIA ISOLATED FROM CLINICAL SAMPLES

Introduction

Laboratory methods used for the identification of anaerobic bacteria isolated from clinical samples have evolved from labour-intense manual methods to more advanced rapid automated systems that have the ability to generate results within minutes as opposed to overnight incubation. The methods used in our laboratory have changed from the manual Finegold and API® rapid ID 32A testing methods used until October 2016, to the automated Vitek® 2 ANC card, for two months only, and then Vitek® MS (bioMérieux, France) from November 2016 until the present.

Method

All anaerobic organisms isolated from clinical samples submitted to our laboratory for identification, from June 2015 to July 2017 were stored in Robertson’s cooked meat medium, for a study looking at antimicrobial susceptibility patterns. Anaerobic identification was confirmed using two MALDI-TOF systems, namely the Vitek® MS (bioMérieux, France) and Bruker MALDI Biotyper® (Bruker Daltonics).
Results
Of 304 anaerobic organisms stored during the study period, 164 isolates were found to be viable. So far, 139 isolates that were tested using manual methods have been retested using the rapid automated systems for confirmation of identification. Preliminary results show a 94% concordance. Mixed cultures, with more than one anaerobic organism isolated, were found to be the main cause of discordant results. As a result, Gram staining and further testing will be carried out on these isolates. Final results will be reported once the study is completed.

Conclusion
The preliminary data show that the rapid automated identification systems introduced recently into our laboratory provide satisfactory results, translating into quicker results for patient management.

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**ABSTRACT TITLE:** AN INVESTIGATION OF TETRACYCLINE-RESISTANT UREAPLASMAS COLONISING PATIENTS AT THE DR GEORGE MUKHARI ACADEMIC HOSPITAL IN 2012 AND 2014

Introduction
Infections of the female urogenital tract and the foetus/newborn are associated with ureaplasmal colonisation. The tetracycline derivative, doxycycline is prophylactically administered to pregnant women at the Dr George Mukhari Academic Hospital. Resistance to tetracycline was reported among these isolates (SIR Mycoplasma kit) however, no data for MIC values to tetracycline are available.

Methods
Twenty-six tetracycline-resistant ureaplasmas (isolated from women presenting for termination of pregnancy in 2012 and 2014) were revived from microbank beads in U9 broth. Positive cultures were inoculated onto A2 agar medium for PCR analysis (speciation; tetM resistance gene). The MIC to tetracycline was done by broth microdilution.

Results
Of the 26 Ureaplasma-resistant isolates, only eight could have been revived for further testing. PCR amplification have shown dual infection with *Mycoplasma hominis* and the presence of the tetM gene in all eight specimens. Broth microdilution results were obtained for four of the 2014 isolates only. Isolates L9 and L72 have shown MIC values of 16 µg/ml and 4 µg/ml respectively. Isolates L18 and L34 were fully susceptible to tetracycline.

Conclusion
Dual infection with *Mycoplasma hominis* was documented for all isolates tested. The promiscuous nature of Tn916 carrying the tetM gene may cause Ureaplasma and *Mycoplasma hominis* in combination to exhibit an increased resistance to tetracycline. Lack of expression of the tetM gene in isolates L18 and L34 may be clarified by searching for mutations on the structural gene itself or on the leader orf12 peptide. Similar studies on a larger scale should be done to review management strategies for preventing complications in pregnant women across the globe.
Results
In this study mutation, Arg463Leu was responsible for INH resistance in 14% (14/100) of our MDR-TB cultures in the absence of codon 315 mutation. This could mean that we are currently missing about 14% of INH resistance when diagnosing MDR-TB patients. The lineage type of thirteen of the fourteen isolates with R463L mutations was Beijing and one was X3.

Conclusion
Discriminatory power will be achieved if this target is considered for inclusion in new rapid molecular tests for improved molecular detection of MDR-TB. Mutation Arg463Leu was more associated with the virulent Beijing strains in this study. This was the first study in South Africa to investigate the association of katG codon 463 mutations with INH phenotypic resistance in MDR-TB strains from Western Cape and Gauteng Provinces.

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ABSTRACT TITLE: ENSURING IMPROVEMENT IN ANTIMICROBIAL HANG TIME THROUGH A MULTI-DISCIPLINARY APPROACH

Introduction
Hang time is measuring the time elapsed from a physician prescribing an antimicrobial agent, until administration. According to Kumar et al., 2006, each hour of delay in the administration of antimicrobials following a doctor’s prescription is associated with an average decrease in patient survival of 7.6% per hour in patients with sepsis. Administration of antimicrobials within the first hour is associated with 80% increased possibility of survival. The aim of the study was to determine improvement in hang time compliance, over a period of 52 weeks involving pharmacy, nursing, doctors and hospital management.

Method
Hang time on first dose antimicrobials was measured across all wards, during daily ward rounds from 4 July 2016 to 2 July 2017. The ward pharmacists collected data on patients who were on specific target antibiotics, using the antimicrobial prescription charts. Data was captured on an electronic surveillance system.

Weekly hang time compliance reports were compiled for the hospital and per unit. The reports were distributed on a monthly basis to the nursing manager and pharmacy manager. The nursing manager distributed the hang time compliance graphs to each unit manager.

Commonly used antimicrobials were made available as ward stock in each unit. Hang time posters were placed in each unit, to create awareness and a SOP on antimicrobial hang time was formulated and approved.

Result
A total of 2 039 cases were assessed for compliance over a period of 52 weeks and 76.56% of the cases received their initial dose of antimicrobials within an hour following a prescription. The median hang time compliance increased from 72.15% in week eight to 89.57% in week 52.

Conclusion
This antimicrobial stewardship project needs constant monitoring, commitment and a multi-disciplinary approach to ensure sustainability. Improvement work is required to increase compliance to ensure all patients receive their first antimicrobial dose within an hour following a prescription.

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ABSTRACT TITLE: THE EPIDEMIOLOGY OF GRAM-NEGATIVE BACTERAEMIA AT TYGERBERG HOSPITAL

Introduction
E. coli and K. pneumoniae are common causes of bacteraemia. β-lactam antibiotics are used to treat these infections. Common cephalosporin and carbapenem-resistance mechanisms include extended spectrum β-lactamases (ESBLs) and carbapenemases respectively. ESBLs belong to three major families: TEM, SHV and CTX-M. There is limited local data describing ESBL distribution. This is important for surveillance and evaluating infection prevention practices.
Methods

Bacteriemic E. coli (n = 70) and K. pneumoniae (n = 70) isolates were collected from the NHLS Microbiology laboratory between April 2015 and March 2016. Identification and antibiotic susceptibility testing (AST) were performed as part of routine testing. Carbapenemase and ESBL genes were detected by PCR, and CTX-M genes characterised by PCR. Isolates were typed using rep-PCR, and images analysed using Gel compar.

Results

Thirty-one percent (44/140) of patients were paediatric. Patients were located in wards throughout the hospital. Fifty-four percent (38/70) of K. pneumoniae and 16% (11/70) of E. coli were cephalosporin resistant (presumed ESBL). None were carbapenem resistant. There was good correlation between phenotypic AST and ESBL PCR. No isolates contained carbapenemase genes. TEM and SHV were the commonest β-lactamases in E. coli (49/70; 70%) and K. pneumoniae (59/70; 84.3%) respectively. Multiple genes were present in 42/70 (60%) of K. pneumoniae isolates, and only 4/70 (5.7%) of E. coli isolates. Of the 46 CTX-M genes, 43 were group 1, two were group 9, and one was untypeable. Strain typing showed substantial diversity among E. coli and K. pneumoniae, with minimal clustering. There was no association between clusters and hospital wards or ESBL type.

Conclusion

While the commonest β-lactamases were TEM- and SHV-related, additional work is needed to further classify these genes. The lack of clustering suggests multiple clones in the hospital. Differentiation of hospital and community acquired infection, and additional molecular typing is planned to further investigate this.

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ABSTRACT TITLE: AN APPROACH TO DETERMINING A BASELINE SOUTH AFRICAN ANTIMICROBIAL STEWARDSHIP CURRICULUM FOR THE BACHELOR OF PHARMACY PROGRAMME

Introduction

Antimicrobial stewardship (AMS) is a rapidly developing healthcare practice which aims to ensure judicious use of antimicrobial agents, to minimise the development of antimicrobial resistance. During a workshop at the 37th annual general meeting of the Academy of Pharmaceutical Sciences of South Africa, AMS training for South African pharmacy students was discussed. This study evolved from that workshop and involves the development of a proposed AMS curriculum, suitable for implementation in undergraduate pharmacy programmes, in South Africa.

Method

The study aims to ensure that the proposed curriculum is in-line with international and national AMS recommendations, and incorporates input from AMS professionals in South Africa. The methodology involves several phases, including: a desktop review; consultation with academic pharmacists from South African universities; and consultation with AMS experts in South Africa (infectious disease specialists, clinical microbiologists and AMS pharmacists). Collation of the results from these study phases will allow for the development of a final proposed AMS curriculum.

Result

The phase one desktop evaluation results have revealed the content and structure of international AMS curricula, as well as the principles which govern AMS in South Africa. Frequency counts have been performed to determine the incidence of trends in the literature. The results indicate that international AMS curricula consist of ten to thirty-hour programmes, which utilise mixed pedagogies and multiple evaluation techniques. The content of these international curricula are in-line with international and national AMS principles. Identification of South African AMS strategic objectives, enablers and principles has further contextualised the initial proposed content for the curriculum being developed.

Conclusion

The findings from the desktop evaluation will form the groundwork of the AMS curriculum to be developed. Collation of these findings with future study phases will allow for the formation of the final proposed AMS curriculum, suitable for incorporation into South African undergraduate pharmacy programmes.

ID: 8598

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ABSTRACT TITLE: AN APPROACH TO DETERMINING A BASELINE SOUTH AFRICAN ANTIMICROBIAL STEWARDSHIP CURRICULUM FOR THE BACHELOR OF PHARMACY PROGRAMME
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ABSTRACT TITLE: SPECTRUM AND SUSCEPTIBILITY PROFILE OF MICROORGANISMS CAUSING PERITONITIS IN CRITICALLY ILL PATIENTS

Introduction
The incidence of drug resistant infections in critically ill patients is reported to be increasing worldwide, including South Africa. Data has shown that patients undergoing re-laparotomy have increased rates of drug resistant bacteria.

The aim of this study was to describe the susceptibility profile of microorganisms found during laparotomy for peritonitis in a group of critically ill patients.

Methods
The study included 35 isolates obtained from critically ill patients in King Edward VIII Hospital ICU. Peritoneal fluid data was extracted retrospectively from the laboratory electronic database for the period 1 January to 31 December 2016. To describe the spectrum of organisms only one of the same isolate per patient was counted.

Results
A total of 35 isolates were obtained from 24 patients (n = 34 from relaparotomy). The most common abdominal pathologies ranged from secondary peritonitis following trauma (n = 7), peptic ulcer disease (n = 5) and iatrogenic bowel injury (n = 5).

The predominant microorganisms isolated were Gram-negative bacteria (n = 19, 54.3%) followed by Gram-positive cocci (n = 12, 34.3%) and Candida species (n = 4, 11.4%). The commonest Gram-negative bacteria were Klebsiella pneumoniae (n = 5, 14.3%), Acinetobacter baumannii (n = 5, 14.3%) and the commonest Gram-positive was Enterococcus species (n = 6, 17.1%). There were high rates of resistance in the enterobacteriaceae isolated, with only 8% being susceptible to amoxicillin-clavulanate; 75% susceptible to imipenem and amikacin; 42% susceptible to piperacillin-tazobactam and gentamycin. 50% of Enterococci were susceptible to ampicillin. 75% of yeasts isolated were non-albicans Candida species. Acinetobacter species isolated were multidrug resistant with only 40% and 20% susceptibility to gentamycin and amikacin respectively.

Conclusion
The high rates of drug resistant bacteria would impact on morbidity of patients. Broad spectrum cover for Gram-positive, Gram-negative bacteria and possibly antifungal therapy would be required for these patients as empiric therapy. The high incidence of non-albicans Candida species is a cause for concern. Final treatment would need to be guided by culture results.

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ABSTRACT TITLE: TEMPORAL GENE EXPRESSION OF CHLAMYDIA TRACHOMATIS IN KERATINOCYTES AT 37OC VERSUS 33OC

Introduction
Keratinocytes are the first port of entry for Chlamydia trachomatis of the lymphogranuloma venereum (LGV) biovar which causes LGV. However, LGV pathogenesis and gene expression studies are usually performed in cells which are not the native host cells.

Methods
To investigate the effect of temperature on the expression of chlamydial genes, HaCaT cells were grown in 12 well plates and inoculated with chlamydia (L2, E and US151) and incubated at 33oC or 37oC. At 2, 12, 24, 36 and 48 hpi (also 60 and 72 hpi for cells incubated at 33 °C), total RNA was harvested and converted to cDNA which was then analysed by RT PCR for the expression of groEL-1, incB, pyk, hctA and omcB. TEM was performed to confirm the stage of the developmental cycle that the organism was at inside the keratinocytes.

Results
RT-PCR analysis showed that the early cycle genes were expressed throughout the cycle at 37oC and 33oC as expected. At 37oC, the mid- and late-cycle genes were expressed throughout the cycle, with different patterns between the isolates tested. At 33oC, the mid-cycle genes also showed different patterns of expression between the tested isolates. However, the expression of late cycle genes in L2 and E was in keeping to that in the published literature.

Conclusion
This study confirms that mid- and late-cycle chlamydial gene expression levels are different to the published research and that temperature has an effect on the level of chlamydial gene expression when grown in keratinocytes.

ID: 8417
Category: Sexually Transmitted Diseases (STDSSA)
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Introduction

Bacterial vaginosis (BV) is the most prevalent dysbiosis that is characterised by a shift from a normal vaginal microbiome (mainly lactobacilli producing lactic acid and hydrogen peroxide) to a polymicrobial mixture of pathogens. Bacterial vaginosis (higher prevalence in black women) has negative impacts on maternal (e.g. increased risk for human immunodeficiency virus acquisition) and foetal health (e.g. premature rupture of the membranes). The vaginal microbiome can be profiled using next generation sequencing platforms like Illumina MiSeq. The aim of this study was to determine the alpha diversity (richness and evenness) of the vaginal microbiome of African women with and without BV and to identify the risk profiles associated with BV and HIV.

Methods

Bacterial vaginosis status was diagnosed using Nugent scoring (categories: healthy, intermediate and BV) of Gram-stained slides. Forty-one samples were selected for targeted 16S metagenomics (categories: healthy, intermediate and BV) of Gram-stained slides. Bacterial vaginosis status was diagnosed using Nugent scoring. HPV infection resulting in persistent HPV infection.

Results

A total of 2 489 715 high quality reads was generated from all samples with an average of 60 724.756 Operational taxonomic units (OTUs) per sample. Lactobacillus iners was the most dominant species in BV negative women while the Prevotella species (spp.) was the most dominant in BV positive women.

Conclusion

Lactobacillus spp. are known to produce hydrogen peroxide, which prevents vaginal colonisation by pathogens. Prevotella spp. such as Prevotella bivia is an opportunistic organism known to invade epithelial cells causing inflammatory responses and endometritis. Therefore, its presence in the vagina warrants efforts to treat and eradicate it.

ABSTRACT TITLE: VAGINAL MICROBIOME PROFILING OF WOMEN WITH AND WITHOUT BACTERIAL VAGINOSIS USING ILLUMINA TARGETED SEQUENCES

ABSTRACT TITLE: HPV INFECTIONS AND CO-INFECTION WITH SELECTED SEXUALLY TRANSMITTED INFECTIONS IN WOMEN ATTENDING DGMAH, PRETORIA, SOUTH AFRICA

ID: 8549

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cytology (LBC) samples are collected thus far; results for 64 patients are analysed. HPV detection was performed using Abbott m2000rt system and detection of other STIs was performed with Anyplex II STI-7 that identifies seven major STIs: Chlamydia trachomatis (CT), Trichomonas vaginalis (TV), Mycoplasma hominis (MH), Mycoplasma genitalium (MG), Ureaplasma urealyticum (UU), Ureaplasma parvum (UP), Neisseria gonorrhoeae (NG).

Results

Only samples positive for HPV DNA were tested for the other STIs. In self-collected samples, 51.8% were positive for UP, 57.4% were MH, 11.1% were TV, 24.1% were UU, 5.6% were CT, 1.9% were MG, 10 of the self-collected samples were not matched with doctor-collected samples. In the doctor-collected LBC samples, 47.3% were positive for UP, TV, 45.5% for MH, 10.9% for TV, 18.2% for UU, 7.3% for CT and 5.5% for MG. STIs multiple infections were detected in 68.5% of the samples.

Conclusion

Almost half of the patients infected with HPV were infected with MH and UP. These two STIs were the common STIs in both the vaginal and cervical LBC samples. The least detected was MG in both types of samples. Co-infection with these STIs may provide an environment for HPV infection to persist.

Results

According to the defined search term criteria, Dengue had the most publications during the study period (6,403 articles), followed by Chikungunya (1,040 articles), Yellow Fever (746 articles) and Zika (417 articles). While the number of Dengue research articles has increased steadily during the study period (14% per year), Yellow Fever research articles have remained relatively static since 2000. Chikungunya research articles show two peak publication periods: 2005–2008 and 2013–2016. The first Zika research articles were published in 2008 and 91% of all Zika research articles were published in 2016. For each illness, the USA contributed the greatest number of research articles (24–51%). While developing countries with an endemic illness burden such as Brazil and India also feature prominently, there is a distinct lack of research output from Africa.

Conclusion

While overall research output for each illness appears to be related to disease prevalence, other factors such as recent outbreaks, increased public awareness and global access to research funding may influence research trends.

ID: 8376

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ABSTRACT TITLE: A BIBLIOMETRIC ANALYSIS OF YELLOW FEVER, DENGUE, CHIKUNGUNYA AND ZIKA RESEARCH TRENDS FROM 2000 TO 2016

Introduction

A bibliometric analysis was conducted to investigate trends in Yellow Fever, Dengue, Chikungunya and Zika research output from 2000 to 2016. For each illness, this included a) description of the number of original research publications published per year; b) analysis of the geographical origins of the research during the defined study period; and c) identification of the top ten most cited articles per illness topic.

Methods

Using the Web of Knowledge Database (https://webofknowledge.com), we searched for any original research articles published between 2000 to present which contained “Yellow Fever”, “Dengue”, “Chikungunya” or “Zika” search terms in the title. The built-in analysis features were used to sort these results by year of publication, geographical origin and number of citations.

ID: 8322

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ABSTRACT TITLE: SIGNAL PEPTIDE PREDICTION SUGGESTS MYCOBACTERIUM TUBERCULOSIS CURLI PILIN SUBUNIT SECRETION VIA THE SEC PATHWAY MAY HINDER MTP OVEREXPRESSION IN ESCHERICHIA COLI

Mycobacterium tuberculosis curli pili (MTP) are novel, potential TB diagnostic biomarkers. However, the production of high quality recombinant transmembrane and secretory proteins may be challenging due to their secretion attributes. The signal peptide of MTP governed by the classical secretion pathway may hinder the purification of the protein in E. coli. In this study, the secretion...
characteristics of MTP were determined and cloning, expression and purification of MTP was attempted in E.coli. A fragment of MTP unique to MTBC was cloned into pet101 and pGEX-6P-1 vectors and confirmed by nucleotide sequencing. Expression of the protein was attempted at IPTG concentrations ranging from 0.1mM to 1mM and at temperatures between 25 °C to 37 °C. The pGEX-6P-1/mtp clone expressed protein was purified, yielding a MTP-GST fusion protein and a free GST band that were analysed by LC/MS mass spectrometry. Inclusion body preparation attempted from the pet101/mtp clone yielded two bands at 10 kDa and below 10 kDa, both of which were analysed by LC/MS mass spectrometry. Transcription activity clones were interrogated by real time PCR. Signal peptide and protein secretion characteristics were determined by bioinformatics analysis. Expression of the pGEX-6P-1/mtp clone demonstrated the expected fragment at approximately 35 kDa, confirmed by Western Blotting but mass spectrometry did not detect any MTP fragments. The bioinformatics analyses of MTP predicted a strong Sec regulated secretion pathway and the absence of non-classical “mycobacterial specific” secretion. Excluding the signal peptide region and using a GST tag greatly enhanced the expression of the protein in the soluble fraction. However, purification of the MTP peptide remained problematic. The predicted Sec regulated secretion pathway may play a role in the inhibition of MTP overexpression in E.coli. Alternatives to E. coli expression or more efficient purification strategies are required for the acquisition of high quality M. tuberculosis antigens.

ID: 8387

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ABSTRACT TITLE: COLISTIN, CARBAPENEM AND CEPHALOSPORIN-RESISTANT KLEBSIELLA PNEUMONIAE REPORTED FROM MISRATA, LIBYA

Introduction

Klebsiella pneumoniae is a significant human pathogen causing community and nosocomially-acquired infections. The spread of extended-spectrum β-lactamase (ESBL)-producing enterobacteriaceae has resulted in the emergence of carbapenem-resistant K. pneumoniae (CRKP). Although colistin is generally used to treatment CRKP infections, resistance to colistin has also been reported in several countries and become a major public health concern.

Method

Clinical samples were obtained from hospitalised (140) and non-hospitalised patients (60) in Misrata, Libya. Identification of the isolated species was achieved using the VITEK 2 compact system. Screening for carbapenem and cephalosporin-resistance was performed using the disk diffusion method with carbapenem 10 µg and cephalosporin 30 µg disks and minimum inhibitory concentrations (MIC) determined by VITEK 2. Colistin resistance was determined using both Sensititre Gram-negative Xtra plate format (GNX2F) and VITEK 2. Carbapenemase activity was detected using the RAPIDEC CARBA NP, Modified Hodge test, carbapenem inactivation methods, MAST Combi Carba plus kit (D73C) and meropenem combined disc test. ESBL production was confirmed using Sensititre ESBL confirmatory plates (ESB1F), modified double disc synergy test (MDDST) and MAST ESBL detection kit DS2C.

Result

Of the eighty K. pneumoniae isolates, 47 (58, 75%) demonstrated resistance to at least one of the four carbapenems, 16 (20%) were ESBL producers, and 2 (2.5%) were carbapenem and colistin resistant. Seventeen (21.25%) isolates were susceptible to all antibiotics tested except ampicillin and augmentin.

Conclusion

This study has revealed the high rate of CRKP among hospitalised patients in Libya with OXA-48 being the most prevalent. The emerging colistin resistance and the spread of ESBL producers in both the hospital and in the community requires urgent action such as the establishment of antibiotic stewardship programmes along with firm infection control practices to prevent further spread in both the clinical and non-clinical settings.

This study was funded by the Libyan government.

ID: 8455

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ABSTRACT TITLE: RESISTANCE TO SECOND-LINE ANTI TB DRUGS IN SOUTH AFRICA: ASSOCIATION BETWEEN MIC AND GENETIC RESISTANCE DETERMINANTS

Background
Genetic polymorphisms among Mycobacterium tuberculosis are known to confer differential levels of resistance. We investigated the association between minimum inhibitory concentration (MIC) levels and mutations in gyrA and gyrB (OFX and MOX) and rrs (AMI and KAN) genes.

Method
The presence of mutations in 210 drug-resistant M. tuberculosis isolates was determined by DNA sequencing and the MICs were determined by DNA sequencing and the MICs were determined by DNA sequencing.

Result
In total, 35 isolates were resistant to OFX, 30 to MOX at > 0.5 µg/ml breakpoint and only two at > 2 µg/ml breakpoint. The gyrA mutations with A90V, D94G, D94N, S91P were detected in 13, 11, 1, 2 and 7, 17, 1 of the OFX and MOX resistant isolates, respectively. Isolates with mutations at the codon 90 and 91 had MIC of 4–8 µg/ml for OFX and 1 µg/ml for MOX, respectively, and isolates with mutations at codon 94 had MIC of 8–32 µg/ml and 1–8 µg/ml for OFX and MOX, respectively. The L566F gyrB mutation were found in two MOX and MOX resistant isolates (MIC = 8 µg/ml). The two MOX-resistant isolates at >2µg/mlbreakpointhadmutationatthecondon94andwereassociated with high MIC (≥ 8). Thirteen isolates showed resistance to AMI and KAN, with 11 isolates showing cross resistance. The majority of AMK (n = 13) and KAN (n = 8) resistant isolates harboured a mutation in the rrs locus. All isolates with mutations at A1401G (6/6) and a double mutation (A514C_A1401G) were found to have a high MIC (16 µg/ml) for AMI as well as for KAN (40 µg/ml).

Conclusions
MTB MICs were found to be consistently lower for MOX than for OFX among isolates with the same gyrA mutation. The codon 94 gyrA and L566F gyrB mutation was associated with high-level resistance to both OFX and MOX. This finding indicates moxifloxacin may be used in the treatment of ofloxacin-resistant TB. Mutation at A1401G of the rrs gene was associated with high-level of resistance conferring cross-resistance between KAN and AMI.

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ABSTRACT TITLE: THE FIRST EVER REPORT ON HAEMOPHILUS INFLUENZAE CARRIAGE IN CHILDREN ATTENDING DGMAH: BIG BOOST FOR HIB VACCINATION

Introduction
In 1999, South Africa incorporated Haemophilus influenzae type b (Hib) vaccine in the infant routine immunisation programme which substantially reduced the rate of mortality and morbidity due to Hib in the country. However, prevalence of H. influenzae has not been evaluated at Dr George Mukhari Academic Hospital (DGMAH) post-introduction of the vaccine. Reports from other parts of the world have reported mixed findings on carriage and type replacement. Hence, this study aimed to determine the prevalence of H. influenzae carriage in the nasopharynx of children served by DGMAH.

Methods
A total number of 350 nasopharyngeal swabs were collected from children aged between two months to 14 years attending the pediatric clinic at DGMAH. The samples were processed in the laboratory using various methods. These included phenotypic traits identification such as colony morphology on chocolate agar, Gram staining, oxidase production and tests on growth factors requirement. Lastly, the identification of isolates was confirmed by conventional polymerase chain reaction targeting the omp6 gene region.

Results
Of the 350 samples processed, 50 isolates were Haemophilus species positive by phenotypic test methods. Of which, 28 (8%) isolates were confirmed positive for H. influenzae by PCR, whereas 322 (92%) were found to be negative for H. influenzae. The first ever electron microscopy pictures of H. influenzae isolates at DGMAH showed the characteristic pleomorphic shape of H. influenzae.

Conclusion
The study demonstrated an 8% H. influenzae carriage rate in children between the ages of two months to 14 years in DGMAH.

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ABSTRACT TITLE: BIG BOOST FOR HIB VACCINATION

Introduction
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Conclusion
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**ABSTRACT TITLE:** CHRONIC DIALYSIS THERAPY YIELDS A SIGNIFICANT INFECTIOUS BURDEN

**Introduction**
Patients receiving chronic dialysis therapy are vulnerable to infection because of regular contact with hospital environments, indwelling catheters for dialysis access and comorbidity such as diabetes.

**Methods**
A retrospective review was conducted of laboratory-confirmed infections in patients on peritoneal dialysis (PD) or haemodialysis (HD) from June 2015 to May 2016. The profile of Gram-positive, Gram-negative bacterial pathogens and fungal pathogens was described and antimicrobial susceptibility patterns were evaluated.

**Results**
Two-hundred-and-thirty-eight infections were reported, corresponding to an infection rate of 9.1/100 dialysis patient years (95% CI: 8.0–10.4). The infection rate for PD was 31.3/100 dialysis patient years (95% CI: 24.2–39.8) which was significantly higher than that for HD (7.2/100 dialysis patient years (95% CI: 6.2–8.3)) (p < 0.0001). Fifty-one bloodstream infections were reported in the HD group and 41 cases of peritonitis in the PD group. There were marked differences in infection rates when comparing regions and units across South Africa.

In those who developed infections, one third had a hospital admission in the last 90 days; 34% were diabetic; mean age at time of infection was 54.3 years. 47.9% (114) were hospitalised for the infection with a median length of stay of seven days and a 5.0% (n = 12) mortality. 49.0% of the patients had Gram-positive infections, 44.4% had Gram-negative infections, and two patients had fungal infections.

**Conclusion**
This is the first published study from a renal dialysis company in SA. The results demonstrate a significant burden of both Gram-positive and -negative organisms and half of these organisms exhibit antimicrobial resistance profiles.

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**ABSTRACT TITLE:** DETECTION OF HUMAN NOROVIRUSES AMONG CHILDREN FIVE YEARS AND UNDER IN KWAZULU-NATAL

**Introduction**
Noroviruses (NoV) are single-stranded RNA viruses belonging to the family caliciviridae. They cause gastroenteritis in all age groups but mostly in young children and the elderly. They are classified into seven genogroups (GI–GVII) and only GI, GII and GIV infect humans. They cause self-limiting infection that resolves in 10–51 hours after exposure and symptoms include diarrhea, vomiting, cramps, chill and headaches. The aim of this study was to detect NoV GI and GII in children ≤ 5 years in KwaZulu-Natal.

**Methods**
Stool specimen were collected in children aged five years and younger with diarrhoeal symptoms in King Edward VIII Hospital. The specimen was tested for NoV antigen using ELISA, and RT-qPCR was used to detect viral RNA. The result was compared to each other.

**Results**
One-hundred-and-eighty-two specimens were tested for NoV by ELISA and RT-qPCR method. Compared to RT-qPCR, the ELISA had a sensitivity of 24.39% (95% CI = 12.36–40.30%) and specificity of 93.62% (95% CI = 88.23–97.04%). The incidence of NoV when specimens were tested by RT-qPCR was 22.5%. Infection rate was higher in children between the age of 12–24 months and there was no statistical significant between females and males.

**Conclusion**
NoV was a common cause of diarrhoeal illness in children presenting to the King Edward Hospital. GII was the most prevalent genogroups with 100% of all positive cases belonging to GI. Taking this into consideration with the lower sensitivity of the ELISA test, the RT-qPCR would be more suitable to routinely testing stool specimens.

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**ABSTRACT TITLE:** INVESTIGATION OF THE MOLECULAR PATTERNS OF CARBAPENEM-RESISTANT PSEUDOMONAS AERUGINOSA AND ACINETOBACTER BAUMANII, AND THE EFFICACY OF THE PHENOTYPIC DETECTION METHODS USED FOR DETECTING CARBAPENEM RESISTANT PSEUDOMONAS AERUGINOSA

**Introduction**
Carbapenem resistant Acinetobacter baumanii and Pseudomonas aeruginosa has been reported worldwide. These pathogens are the
most common cause of nosocomial infections and they result in serious outbreaks in hospitals especially in ICU wards. Of concern is the multidrug resistance pattern of these pathogens resulting in challenges in their treatment. The resistant genes are plasmid-mediated which could result in hospital outbreaks, a challenge for infection control.

This study was done to compare the best method which can rapidly detect carbapenemase-producing organisms in Acinetobacter baumanii and Pseudomonas aeruginosa at the Dr George Mukhari Academic Hospital in 2015.

**Methods**

A total number of 35 Gram-negative isolates from National Health Laboratories Services (NHLS) at Dr George Mukhari Academic Hospital (DGMAH) and referral hospitals isolates with reduced susceptibility to carbapenems were collected from August to December 2015. Twenty-nine were Acinetobacter baumanii and 16 were Pseudomonas aeruginosa identified using the Vitek (BioMerieux). MICs for carbapenems were confirmed using an Etest. Resistant and intermediate isolates were further subjected to Rapidec Carba NP test. Conventional PCR was done to compare the sensitivity and specificity of the method. The PCR used targeted six common carbapenemase genes (BlaKPC, blaNDM, blaIMIP, blaOXA-48, blalMI, blaGES.)

**Results**

A total of 75.8% (22/29) Acinetobacter baumanii isolates were resistant and 3.4% were in the intermediate category for carbapenem. Twenty-one isolates (21/29) 72.4% were Rapidec Carba NP test positive. Rapidec Carba NP test had a sensitivity of 95% and a specificity of 100% for A. baumanii isolates. Eight (8/16) 50% Pseudomonas aeruginosa isolates had MIC ranging between 4 µg/ml to 32 µg/ml, 43.8% (7/16) were positive for Carba NP test and 56.3% (9/16) were positive for PCR.

**Conclusion**

Carba NP test is the most efficient phenotypic method for detecting carbapenemase-producing organisms, it is rapid and easy to perform. The circulating carbapenemase genes were blaGES for Acinetobacter baumanii and for Pseudomonas aeruginosa are blaVIM and blaKPC.

**ID: 8335**

**Category:** Infectious Diseases (IDSSA)

**Permission:** Yes

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**ABSTRACT TITLE:** COMPARISON OF MINIMAL INHIBITORY CONCENTRATION (MIC) VALUES PERFORMED BY MICROSCAN WITH SENSITITRE PANELS FOR TESTING OF ENTEROBACTERIACEAE

**Introduction**

Enterobacteriaceae are large group of Gram-negative, rod-shaped bacteria which are the most frequently isolated from clinical specimens. They may account for up to 80% of clinically significant Gram-negative bacilli and up to 50% of all clinically significant bacteria. Enterobacteriaceae cause both nosocomial and community-acquired infections and are increasingly resistant to antimicrobial agents. Laboratory identification and susceptibility testing are essential for diagnostic and antimicrobial stewardship programmes. The aim of the study was to compare antimicrobial susceptibility testing results by the MicroScan® with Sensititre MICs panels for Enterobacteriaceae.

**Methods**

Total number of 49 isolates from August 2015 to December 2016 were obtained from patients with clinically diagnosed bloodstream infections randomly selected and used to determine the MIC. Results of MicroScan testing were extracted from the access database. The same isolates stored in -70 were then processed using Sensititre® AIL™ panels and read using Vizion® instrument.

**Results**

Total of 49 isolates, including 38 Klebsiella pneumoniae, 4 Enterobacter cloacae and Escherichia coli, 1 Klebsiella oxytoxica, Klebsiella variicola and Serratia marcesens were processed. The following percentages of agreement was obtained for the antibiotics tested: ciprofloxacin 98%, cefotaxime 96%, ceftazidime 96%, gentamicin 98%, amikacin 98%, ertapenem 76%, imipenem 92% and tigecycline 100%. No errors were obtained for ceftazidime, amikacin and tigecycline.

**Conclusion**

The two instruments compared well for most of the antibiotics. Only ertapenem had decreased agreement and cannot be used as screening antibiotic for carbapenemases-producing Enterobacteriaceae.
ABSTRACT TITLE: ZOONOTIC AETIOLOGIES IN FEBRILE ADULTS IN THE MNISI COMMUNITY, MPUMALANGA, SOUTH AFRICA

Introduction
Zoonoses cause infectious diseases in humans who interact with livestock, domestic animals and vectors. A high prevalence of zoonotic infections was observed in a previous study at three public primary health clinics (PHCs) in Mnisi. Subsequently, a community health centre (CHC) in Mnisi was established as a sentinel site for the NICD surveillance programme which allows for seroprevalence monitoring of selected zoonotic diseases in an agro pastoral rural community.

Methods
From September 2014 to December 2016, consenting adult patients with fever (> 37.5 °C)/history of fever on whom a malaria rapid test was done were enrolled from one CHC and a questionnaire administered. Acute and convalescent blood samples were collected for laboratory testing for leptospirosis, Q fever, bartonellosis, brucellosis, arboviruses and rickettsia.

Results
In total, 70 patients were enrolled: 46% (32/70) did not return for follow-up bloods. Median age was 34 years (IQR 26–46 years); 60% were female. Median duration of illness was two days (IQR 1–3 days); 60% (40/67) received an antibiotic at the clinic and 11% (8/70) referred to the hospital. Twenty-four percent (17/70) of patients had no systemic symptoms, 59% (31/53) presented with only one symptom: muscle pain (67%) and respiratory symptoms (39%) were most common. On laboratory testing, 79% (55/70) of patients showed evidence of a recent or past infection/exposure for at least one of the zoonotic diseases: 63% (42/67) for tick bite fever, 19% (13/70) for Q fever, 4% (3/70) for arboviruses, 1/70 for leptospirosis and zero for bartonellosis or brucellosis.

Conclusion
Compared to the previous study at three PHCs, fewer patients at this one CHC, tested positive for Q fever (18.6% vs 37.8%; p-value 0.0110) and Bartonella (0% vs 9.5%, p-value 0.0358). Possible reasons for these differences may include: i) patients were enrolled from only one CHC, ii) the demographic may be different, iii) there was a drought which could decrease vectors, and iv) fewer patients returned for follow-up blood samples.

ID: 8404

ABSTRACT TITLE: CLOSTRIDIUM DIFFICILE INFECTION PATIENT CHARACTERISTICS AND ASSOCIATION WITH IN-HOSPITAL MORTALITY IN SOUTH AFRICA

Introduction
Clostridium difficile infection (CDI) is a global health problem resulting in severe diarrhoea and increased morbidity and mortality. The South African population includes a high prevalence of HIV and tuberculosis. These patients often have prior healthcare exposures and high risk for CDI. CDI has not been well studied in this population. This study aimed to identify baseline patient CDI characteristics and associations with patient outcomes.

Method
A retrospective review of adult patients with diarrhoea and a CDI test result was conducted at four secondary level public sector hospitals in Cape Town, South Africa. Patient characteristics, CDI management, and patient outcomes were collected. Univariate, logistic regression, and survival analyses were performed.

Result
A total of 116 positive and 148 negative CDI test results were reviewed. In-hospital mortality was more common in the CDI positive group (28% vs. 8%, P < 0.0001; hazard ratio 2.1, 95% CI 1.2–3.7). A logistic regression including CDI test result, prior hospitalisation (30-day and 90-day), critical care, tuberculosis, gender, multi-drug resistant tuberculosis, and hematochezia, found an independent risk of in-hospital mortality with a CDI positive test result (OR 4.4, P = 0.001). Clinically meaningful independent
variables associated with mortality included comorbid tuberculosis (OR 2.2, P = 0.035) and male gender (OR 2.5, P = 0.05). Tuberculosis was also significantly associated with a CDI positive test result (P < 0.05). Antibiotic exposure and hospitalisation in 30 days prior to CDI test were significantly associated with a CDI positive result (P < 0.05), but did not impact mortality.

Conclusion

Patients testing positive for CDI in South Africa have a significantly higher risk of mortality. Male patients are also at higher risk of in-hospital mortality, but there was no gender association with CDI test result. Initial findings suggest tuberculosis comorbidity may be considered a risk factor for CDI, in addition to known risk factors, prior antibiotic exposure and recent hospitalisation.

ID: 8475

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ABSTRACT TITLE: MYCOBACTERIUM TUBERCULOSIS HBHA AND MTP DELETION ELICITS UNIQUE CANONICAL PATHWAYS DURING EARLY INFECTION IN THP-1 DIFFERENTIATED MACROPHAGES

Introduction

*M. tuberculosis* was responsible for an estimated 10.4 million new cases and 1.4 million deaths in 2015 (WHO, 2016). Understanding host gene regulation provides vital information about pathogenesis that possibly translates into novel therapeutic targets. This study utilised THP-1 human macrophages infected with a hbhA-mtp knockout mutant of *Mycobacterium tuberculosis* to assess the effect of hbhA gene deletion in the Δmtp mutant on the host transcriptome.

Methods

In order to compare the transcriptome changes elicited by wild type (WT) and ΔhbhA-mtp infection at 4 hr post-infection, RNA was extracted from THP-1 differentiated macrophage monolayers and sequenced using Hiseq 2500 platform (Illumina). Reads were trimmed using Trimmomatic and mapped to the Homo sapiens reference genome Hg19 (UCSC) using TopHat (2.1.0) and Bowtie2. Differential gene expression patterns were identified using cuffdiff. Changes in canonical pathways were analysed by Ingenuity Pathway Analysis.

Results

WT infection enriched the TREM1 signalling pathway to a greater degree than ΔhbhA-mtp infection in macrophages. ΔhbhA-mtp infection enriched the dendritic cell maturation pathways to a greater degree than WT infection. ΔhbhA-mtp infection, but not WT, enriched Th1 and Th2 Activation, Th1, Th2, melatonin degradation, sumoylation, methylglyoxal degradation III, granzyme A signalling, PCP pathways.

Conclusions

The absence of HBHA and MTP proteins resulted in activation of host immune responses that are detrimental to the survival of *M. tuberculosis*, suggesting that these adhesins may assist the pathogen to evade the host immune responses during early infection.

ID: 8501

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ABSTRACT TITLE: PREVALENCE OF ANTIBIOTIC RESISTANCE AND VIRULENCE GENES OF ESCHERICHIA COLI O157: H7 ISOLATED FROM PORCINE IN GAUTENG, SOUTH AFRICA

Introduction

*Escherichia coli* (*E. coli*) is an emerging pathogen, harboured in the gastrointestinal tract of mammals. Most *E. coli* strains are harmless, however, some *E. coli* strains harbour virulence factors which cause intestinal and extra-intestinal diseases in animals and humans. Limited information is available on the molecular characteristics and antibiotic resistance of *E. coli* O157:H7 isolates from pigs in South Africa. The study’s aim was to identify presumptive *E. coli* O157:H7 isolates obtained from slaughtered pigs from five Gauteng abattoirs and to determine the prevalence of selected virulence and antibiotic resistance genes.
Methods

Isolates (n = 195) were collected from porcine rectal swabs. Methods used to confirm and identify presumptive E. coli O157:H7 isolates are: ChromID™ O157:H7 (bioMérieux, France) agar, MALDI-TOF (Daltonics, USA) analysis and a Multiplex PCR (M-PCR) assay [detecting O157 antigen (rfbE) and H7 flagella (fliC) genes]. M-PCR assays were performed to detect selected virulence [Shiga toxin 1 and 2 (Stx1 and Stx2), intimin (eae) and haemolysins (hlyA)] and antibiotic resistance [Sulfonamide (sul1), Tetracycline (tetA & tetB), Chloramphenicol (cmIA), Trimethoprim (dhfr1), Aminoglycoside (AadA)] genes.

Results

Isolates (n = 77) were identified as E. coli. Six isolates showed prevalence of fliC and rfbE genes, confirming E. coli O157:H7. Results shown prevalence 33.3% (2/6), 83.3% (5/6) and 100% of Stx 2, eae and hlyA virulence factors, respectively. No Stx 1 was detected. The tetA, dhfr1 and sul1 genes were detected in 50% (3/6) of the isolates. The tetB and cmIA genes were detected in 66.7% (4/6) of isolates. All isolates were positive for the aadA gene.

Conclusion

Detection of virulence genes (stx2, eae and hlyA) presents potential health risks to close human contacts (abattoir workers) and consumers of undercooked meat products. Antibiotic resistant genes (tetA, dhfr1, sul1, aadA and dhfr1) detected, confer resistance to different antibiotic classes. Continuous surveillance for E. coli O157:H7 is therefore important at the human-animal interface to prevent possible outbreaks.

ID: 8526

Category: Infectious Diseases (IDSSA)

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ABSTRACT TITLE: MOLECULAR EPIDEMIOLOGY OF LEGIONELLA PNEUMOPHILA IN SOUTH AFRICAN HOSPITALS, 2015–2016

Introduction

Healthcare facilities globally are often identified as sources of Legionnaire's disease (LD) due to the combination of large, complex plumbing systems, and potential exposure to individuals with increased risk of infection. Globally, serogroup (SG) 1, sequence type (ST) 1 is the most common strain identified by sequenced-based typing (SBT). Due to limited data, we aimed to describe the molecular epidemiology of L. pneumophila isolated from hospitals in South Africa.

Methods

From March 2015 through April 2016, 77 Legionella spp. isolates were isolated from routine water samples or site investigations from 17 private and public hospitals, representing four of the nine provinces [Eastern Cape (n = 4), Western Cape (n = 4), Gauteng (n = 8) and Mpumalanga (n = 1)] where there is an increased awareness for testing and notification of LD. Sequences of the 7 SBT genes were extracted from whole genome sequencing data, allele numbers and STs were assigned using the online L. pneumophila SBT database.

Results

Of the 77 isolates, 42 (55%) were L. pneumophila SG1, 31 (40%) were L. pneumophila SG2-14 and 4 (5%) were other Legionella spp. Of all L. pneumophila isolates, we identified 4 known STs: ST1 (59%; 43/73), ST421 (21%; 15/73), ST87 (1%; 1/73) and ST242 (1%; 1/73). SG1 comprised ST1 (91%; 30/33) and ST421 (9%; 3/33) while SG2-14 comprised ST1 (50%; 13/26), ST421 (42%; 11/26), ST87 (4%; 1/26) and ST242 (4%; 1/26). We identified 13 isolates with novel STs identified at multiple hospitals. L. pneumophila SG1 ST1 was the most common strain, detected in 59% (10/17) of hospitals. Three STs were identified in one Gauteng hospital [4/20 (20%) ST1, 15/20 (75%) ST421 and 1/20 (5%) novel ST].

Conclusions

SG1 ST1 was the most common strain detected in the water systems of hospitals, as has been described previously. Amongst SG2-14 isolates, two additional STs (ST87 and ST242) were identified compared to SG1 isolates. STs that have not previously been described were identified.
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ABSTRACT TITLE: MALARIA OUTBREAK INVESTIGATION – MOPANI AND VHEMBE DISTRICTS, LIMPOPO PROVINCE, SOUTH AFRICA, MAY 2017

Background
Malaria remains a major public health challenge in Vhembe and Mopani districts, Limpopo province – two of the five malaria endemic districts in SA. On 2 May 2017, the two districts reported unusual increase of malaria cases. Investigations were conducted to establish existence, magnitude and cause of the outbreak, to recommend control measures and future prevention.

Methods
A cross-sectional investigation was conducted. Discussions with key stakeholders were held. Malaria case data from April to May 2017 were reviewed. A case was defined as a person with malaria confirmation by microscopy or a rapid diagnostic test (RDT) in health facilities of the two districts from April–May 2017.

Results
A total of 5 662 cases, 55.3% in Mopani, were reported with a peak in May 2017. Investigation of likely origin revealed that 5 574 (98.4%) were local cases in both districts. Fifty-four malaria deaths were reported, the majority 42 (77.8%) in the Mopani district. The median age of the patients who died was 44 years (IQR: 35–60) with 0.95% case fatality rate (CFR). Indoor residual spraying (IRS) coverage was less than the WHO recommended 80% universal coverage for control. Environmental assessment revealed that the 2016/17 season recorded higher levels of rainfall compared to previous malaria seasons. Key stakeholder interviews revealed that the outbreak was associated with: late commencement of IRS due to late appointment of spray teams; reduced use of dichlorodiphenyltrichloroethane (DDT) replaced by pyrethroids (K-Othrine and deltamethrine); and stock outs of RDTs and antimalarial drugs in clinics.

Conclusion
The investigation established that the marked increase in malaria cases in the two districts was not a seasonal phenomenon but rather an outbreak. We recommended: an awareness programme through local radio stations on malaria prevention and treatment; an improved supply chain of RDTs and antimalarial drugs; adequate budget for procurement of DDT and advance appointment of IRS teams.

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ABSTRACT TITLE: CHARACTERISATION OF RIFAMPICIN AND ISONIAZID RESISTANCE OF MYCOBACTERIUM TUBERCULOSIS STRAINS FROM CLINICAL SPECIMENS SENT TO DR GEORGE MUKHARI TERTIARY LABORATORY IN SOUTH AFRICA

Introduction
Tuberculosis (TB) continues to be a global threat. A series of molecular characterisation of the organism have since been developed and introduced, which include the MTBDRplus assay that detects Mycobacterium tuberculosis (MTB) and susceptibility to rifampicin (RIF) and isoniazid (INH); and GeneXpert MTB/RIF assay that detects MTB and susceptibility to only RIF. This study was conducted to characterise the drug resistant patterns of M. tuberculosis strains.

Methods
This was a retrospective data analysis of MTB from the Laboratory Information System (LIS) TrackCare from 01 January 2016 to 31 December 2016. A total of 11 829 results were retrieved from specimens collected from different body sites sent to the National Health Laboratory Service, Dr George Mukhari (NHLS-DGM) Tertiary Laboratory, Medical Microbiology Department, during the study period for routine TB tests including microscopy, culture, and drug susceptibility testing using the MTBDRplus genotypic assays.

Results
Of the 11 829 clinical results retrieved, 6 591(56%) were MTB positive. In total, 5 356 (81.3%) were fully sensitive and 1 235 (18.7 %) were resistant to either RIF, INH or both drugs. A total of 415 (33.6%) were multidrug-resistant (MDR) TB; 422 (34.2%) were RIF monoresistant; 398 (32,1%) were INH monoresistant; overall, 510 (61.3%), RIF resistance was due to MUT3 in the rpoB gene. The prevalent INH resistance was due to katG gene mutation (MUT1) (49.9%).
Conclusion

This study shows that the most prevalent mutations in rpoB gene for RIF are caused by mutation in the MUT3 and MUT1 in the katG gene for INH.

ID: 8321

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ABSTRACT TITLE: CANDIDA SPECIES CARRIAGE IN DIABETIC PATIENTS IN MISRATA, LIBYA

Introduction

There is a paucity of studies describing the prevalence and antimicrobial profiles of Candida in Libya. Limited treatment choices in the antifungal armamentarium in public healthcare settings in the rest of Africa require a study of the prevalence and susceptibility of Candida species in Libya, where antifungals are not regularly distributed in public healthcare settings.

Methods

In this study, 170 diabetic mellitus type 2 (T2DM) patients were examined for Candida carriage in the oral mucosa, using differential/Fluka and Oxoid chromogenic media and API 32 ID C biochemical testing. Fluconazole susceptibility was investigated by disc diffusion on YNBG agar. Isolates were graded as susceptible, intermediate or resistance according to their inhibition zone measurements and microcolony scores.

Results

Thirteen species were identified from 182 isolates with a frequency of 69 C. albicans, 42 C. dubliniensis, 26 Chumicola, 20 C. glabrata, 5 isolates of C. krusei, C. tropicalis and C. kefyr respectively, 4 Csake, 2 C.parapsilopsis, 2C. magnoliae and 1 isolate each of C. guilliermondii, C. globsa and C. membranifaciens. Although largely susceptible to fluconazole, C. albicans, C. dubliniensis, C. hemicola and C. sake demonstrated an emerging resistance with intermediate to total resistance observed in all the other species except for C. magnolia and C. globsa which were both susceptible to fluconazole.

Conclusion

Early recognition and treatment of rare or resistant species which may be contributing to patient morbidity and mortality in Libya is imperative.

This study was funded by the Libyan government.

ID: 8393

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ABSTRACT TITLE: PERCEPTIONS AND PRACTICE OF INFECTION CONTROL AMONG MEDICAL STAFF AT FOUR MAIN HOSPITALS IN THE VHEMBE DISTRICT

Introduction

Recent studies have shown that nosocomial infections are often spread through devices used by healthcare professionals. The present study assessed the perceptions and practice of infection control (IC) by hospital staff in Vhembe District and evaluated the contamination of stethoscopes and cell phones.

Methods

In a cross-sectional study, questionnaires were administered to physicians and nurses. Stethoscopes and cell phones were swabbed for the isolation of microorganisms.

Results

A total of 178 participants aged between 23 and 65 years with a mean of 42.29 ± 9.76 years were recruited in the study. About 46% of staff cleaned their stethoscopes sometimes and handwashing was practiced by 52% of them. They were largely ignorant of other IC strategies such as antibiotic stewardship, surveillance or vaccinations. Many participants indicated that further education should be provided and more resources allocated for IC. About 88% of cell phones and 80% of stethoscopes were contaminated and yeast was the most commonly isolated microorganisms while Staphylococcus spp were isolated from 27% of the stethoscopes.

Conclusion

The present study showed that the practice of infection control by hospital staff is still poor and there is need for further effort and education to hospital employees in order to empower them on infection control strategies. Stethoscopes and cell phones are carriers of pathogens and their cleanliness should be emphasised among healthcare workers.
Variation and azole-resistance. Further studies will be useful in understanding the emergence of *Candida auris* bloodstream infections in the private sector in South Africa, exhibiting regional trends.

**Conclusion**

Echinocandins, respectively. The susceptibility to the antifungals: 97% (126/223) being > 50 years old. Fifty-nine percent of cases were aged ≥ 65. The most commonly used antibiotics were piperacillin-tazobactam (20%) followed by amoxicillin (12.37%), toxin A (tcdA) and toxin B (tcdB) was detected in all isolates, however the binary toxin was not detected.

**Conclusion**

*Candida auris*, first described in 2009 after isolation from the external ear canal of a patient in Japan has emerged as a global pathogen, including in South Africa. Phenotypically, it resembles *Candida haemulonii* and until recently, required molecular methods for identification. Currently, both matrix-assisted laser desorption/ionisation-time of flight (MALDI-TOF) mass spectrometry and the Vitek-2 systems can reliably identify this species. This organism has been associated with hospital outbreaks and multidrug resistance (MDR; resistance to two or more antifungal classes).

We report on the emergence of *Candida auris* bloodstream infections in a private healthcare setting in South Africa and describe the susceptibility patterns of this emerging pathogen.

**Methods**

All Candida species identified as *C. auris* or *C. haemulonii* from blood cultures from January 2015–June 2017 were analysed. Species identification was made using the MS or Vitek systems (bioMérieux, Marcy l’Etoile, France).

There are no agreed-upon interpretive breakpoints for *C. auris* for antifungal agents; however, conservative breakpoints, developed for other *Candida* species, have been applied to *C. auris*. The Vitek (bioMérieux, Marcy l’Etoile, France) or Sensititre YeastOne® (Trek Diagnostic Systems Ltd) were used to perform sensitivity testing.

**Results**

Of 1 625 *Candida* species blood culture isolates from January 2015 through to June 2017, 14% (223/1 625) were *Candida auris*/*haemulonii*, of which 98% (219/223) were from the Gauteng province. The age range was 0–94 years (median 53 years), 57% (126/223) being > 50 years old. Fifty-nine percent of cases were in ICU/high care wards. Susceptibility to the antifungals: 97% (216/223), 2% (4/223), 86% (31/223) and 98% (218/223) were susceptible to amphotericin B, fluconazole, voriconazole and the echinocandins, respectively.

**Conclusion**

*Candida auris*, the second most frequent cause of non-albicans candidaemia in the private sector in South Africa, exhibited regional variation and azole-resistance. Further studies will be useful in monitoring the evolution of the susceptibility patterns of *Candida auris* bacteraemia in South Africa.
**ABSTRACT TITLE: HYDATID CYST DISEASE IN KHUZESTAN, IRAN, DURING 2000–2015**

**Introduction**

Hydatid cyst disease (HCD) is a well-known parasitic disease globally. It develops in humans after ingestion of *Echinococcus granulosus* eggs. In order to better prevent and control HCD, it is crucial to identify the epidemiological aspects of this parasitic infection. The current study aimed to evaluate the epidemiology and features of HCD in a livestock-raising area in Khuzestan, southwest of Iran.

**Methods**

The present study was a descriptive-analytical study conducted on 360 patients from different areas of Khuzestan Province, southwest of Iran, with a diagnosis of HCD between 2000 and 2015, using their records. Demographic data, clinical features, and radiological data were collected. Data were summarised and analysed using descriptive and analytical statistical methods, respectively.

**Results**

56.1% of the patients were female. The mean age of the patients was 37.36 ± 15.2 years. Most patients were in the over-50-years-old age group [103 (28.6%)], and the less-than-10-years-old age group had the lowest number [19 (5.3%)]. Most of the cysts were detected in the liver [234 (65%)]. Regarding the finding, there was no statistically significant association between sex and residing area, animal contact, and the number of the cysts (p = 0.12, 0.36, and 0.95, respectively); however, a significant association was found between sex and the body organ involved (p = 0.007), so that liver involvement was mostly detected in females (79.9%), while involvement of the lung was mostly found in males (66.4%). No statistically significant association was found between age and the number of the cysts or the body organ involvement (p = 0.35 and 0.61, respectively).

**Conclusion**

Our study showed that HCD could be surprisingly common in apparently low-risk populations, such as those living in urban areas or without direct contact with dogs and farm animals. So, educating the community about the most common modes of acquisition could be helpful for the control and prevention of this disease.

**ABSTRACT TITLE: SEROPREVALENCE OF HEPATITIS E VIRUS INFECTION IN PATIENTS WITH LIVER DISEASE IN A REFERRAL HOSPITAL IN BOTSWANA**

**Introduction**

This study reports the seroprevalence of Hepatitis E Virus (HEV) infection in patients presenting to a referral hospital in Botswana.

**Methods**

A cross-sectional analysis of HEV infections among 331 patients presenting with clinical features of jaundice, elevated liver function tests, or a previous history of liver disease, was conducted at Princess Marina Hospital, Botswana from February 2015 to July 2016. Serum was screened for the presence of antibody to HEV IgM and IgG, using enzyme linked immunosorbant assay techniques (ELISA). Risk factors for HEV infection were assessed using logistic regression techniques (STATA version 13.1).

**Results**

Of the 331 participants, antibody to HEV was positive in 9.7% (n = 32). Anti HEV IgM was positive only in one (0.30%) of the 331 samples. Thirty-one (9.37%) samples were positive for anti HEV IgG. HEV antibodies were detected in 17 (53%) males and the median
age was 45 years (Q1:36, Q3:56). Seven (4.6%) patients with HEV positive serology were also HIV positive. Fifteen (48.4%) HBsAg positive samples were also found to be anti HEV IgG positive. HCV confection was reported in 2 (6.5%) samples. Associated symptoms included nausea and loss of appetite. Risk factors associated with HEV infection in this study were lack of proper hand washing, alcohol consumption, and HIV infection. People on HAART and sexually active people were also at risk of getting HEV infection. Low serum bilirubin and serum alkaline phosphatase were associated with HEV infection.

Conclusion
Prevalence of HEV infection is 9.7% in Botswana among patients with liver disease. This is the first study reporting the seroprevalence of HEV infection in Botswana. We recommend a routine hepatitis profile screening of all patients with clinical manifestations of liver disease as HEV infection can cause fulminant hepatitis.

**ID: 8580**

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**ABSTRACT TITLE:** THE INCIDENCE OF DISCORDANT RIFAMPICIN SUSCEPTIBILITY RESULTS BETWEEN GENOTYPIC AND PHENOTYPIC METHODS IN MYCOBACTERIUM TUBERCULOSIS COMPLEX ISOLATES AT DR GEORGE MUKHARI HOSPITAL TERTIARY LABORATORY

**Introduction**
Drug resistant tuberculosis increasingly compromises the global success of tuberculosis treatment. Rapid and accurate detection of Mycobacterium tuberculosis and drug susceptibility testing to rifampicin and isoniazid is the first requirement to treat these patients correctly, which will in turn limit transmission of drug resistant strains. Frequent discordant results between routinely used phenotypic and genotypic drug susceptibility testing methods at the National Health Laboratory service – Dr George Mukhari Tertiary Laboratory, are continuously observed, especially with rifampicin posing challenges and delay in tailoring correct treatment for patient management. This study aimed to determine the incidence of discordant rifampicin susceptibility results between the genotypic and phenotypic drug susceptibility testing methods.

**Methods**
From June 2013 to December 2015, we consecutively collected 50 Mycobacterium tuberculosis rifampicin discordant samples with GenoType MTBDRplus and MGIT 960 phenotypic drug susceptibility testing. The genomic DNA of the discordant results was extracted and sent to the Inqaba Biotech Pty (Ltd) for the rpoB gene sequencing.

**Results**
Of the 50 samples in which the rpoB gene sequencing was done, rpoB gene sequencing yielded 41/50 (82.0%) interpretable results. Of these, 39/41 (95.1%) patients were co-infected with HIV. There were more females 21/41 (51.2%) than males 20/41 (48.8%). Of the 41 resulted samples, 2/41 (4.8%) yielded no rpoB mutation. The most prevalent mutation was Ser531Leu 12/40 (29.2%) samples, followed by His526Leu 7/41 (17.0%) samples; His526Asp 6/41 (14.6%) samples; Leu511Pro 5/41 (12.1%) samples; His526Try 5/41 (9.7%) samples, and both Asp516Val as well as Asp218His in 2/41 (4.8%) samples.

**Conclusion**
It is important to detect rpoB gene mutations to accurately and promptly diagnose drug-resistant strains to limit the delays in patient management. This study shows that DNA sequencing is the best suited for evaluating suspected drug-resistant Mycobacterium tuberculosis isolates with discordant results for phenotypic susceptibility and rapid molecular testing.

**ID: 8379**

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**ABSTRACT TITLE:** VERIFICATION OF THE HAIN MTBDRSL AND BD MGIT PHENOTYPIC DRUG SUSCEPTIBILITY TESTING (DST) FOR TUBERCULOSIS AT THE TSHWANE ACADEMIC LABORATORY (TAD)

**Introduction**
Drug resistant TB (DR-TB) is associated with higher morbidity and mortality than drug sensitive TB. Rapid detection of these strains is important for appropriate treatment as well as prevention of further transmission of these strains in the community. A tertiary academic microbiology laboratory such as TAD should be able to perform DST on these strains routinely. The aim of this study was to verify the MTBDRsl assay (LPA, Hain Lifescience, Nehren, Germany) and the MGIT (Becton, Dickinson and Company, USA) phenotypic DST assay.
Methods
Our laboratory received a proficiency panel of twenty TB strains with varying drug susceptibility to the second line agents from the TB Reference Laboratory at the National Institute of Communicable Diseases (NICD, Sandringham, Johannesburg, South Africa). These strains were tested using the MTBDRsl assay (Version 2) and the MGIT phenotypic susceptibility testing using five drugs with concentrations of: amikacin (1 µg/ml), capreomycin (2.5 µg/ml), kanamycin (2.5 µg/ml), moxifloxacin (0.5 µg/ml and 2.0 µg/ml) and ofloxacin (2.0 µg/ml)

Results
The TAD laboratory received a score of 95% for DST using the MTBDRsl assay. Nineteen strains were available to compare MTBDRsl and the BD MGIT phenotypic DST assay. There was one discrepant result for fluoroquinolone DST, that tested resistant with the MTBDRsl assay and sensitive with the phenotypic DST. This did not change on repeat testing. For aminoglycoside DST, there were no discrepancies observed between the two methods.

Conclusion
The TAD laboratory was found to be proficient in the use of the MTBDRsl assay. When the MTBDRsl assay was compared to the BD MGIT phenotypic susceptibility result, it had a sensitivity and specificity of 100% for aminoglycoside susceptibility and a sensitivity of 94% and a specificity of 100% for fluoroquinolone susceptibility. The TAD laboratory has subsequently implemented second-line testing for TB as a routine.

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ABSTRACT TITLE: PREVALENCE AND INTENSITY OF SEXUALLY TRANSMITTED DISEASES (SASPID) IN LAGOS METROPOLITAN LAGOS STATE, NIGERIA (A CASE STUDY OF HIGHEST POPULATED COMMERCIAL SEX WORKERS CENTRE)

Introduction
The spread of sexually transmitted diseases is growing at an alarming rate through commercial sex workers. A survey carried out in one of the hotels considered as the highest population for commercial sex workers in Lagos, Nigeria reveals that 70% commercial sex workers were infected with different severities of diseases ranging from HIV, Hepatitis B, N. gonorrhea, Staphylococcus aureus, Candida albicans, etc. 20% accounted for microscopically low pus cell count levels which could not yield significant growth after 48 hours of incubation in the laboratory. However, 10% indicated no record of any form of sexually transmitted diseases.

Case
In addition, 70% of the heavily infected cases of the study suggest that three-quarters accepted treatment and counselling while one-quarter refused to submit themselves for medical treatment and counselling as a result of fear of exposure to the law enforcement agencies.

Three-quarters of the 20% lower cases accepted counselling while 2/5 declined. All the patients with no record of sexually transmitted diseases accepted counselling and do not have any intent in quitting prostitution.

Discussion
The study reveals that the contributory factor to the prevalence and intensity are poverty (41%), unemployment (35%), education (10%), government enforcement (9.5%), refusal of male counterpart to apply contraceptive e.g. condom (4.5%) and are responsible for the prevalence and intensity of sexually transmitted diseases among commercial sex workers.

Finally, recommendations were made on how to address these causes for a lasting solution. Three-hundred-and-fifteen commercial sex workers participated in the study.

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ABSTRACT TITLE: GENETIC DIVERSITY OF MYCOBACTERIUM BOVIS BCG STRAINS; CLINICAL AND VACCINE
Mycobacterium tuberculosis BCG is an attenuated strain of M. bovis used to vaccinate infants against disseminated tuberculosis. Attenuation of the vaccine resulted from mutation of the wildtype M. bovis. While protection from disseminated tuberculosis was shown in infants, complications following vaccination have been reported. Several BCG sub-strains now exist following passaging of the original Pasteur strain for vaccine manufacture. What has not been elucidated is whether these differences have any bearing on the risk for complication with each sub-strain.

The aim of this study was to identify genetic variation between the invasive BCG and non-invasive BCG reference strains.

In this study five clinical (M. bovis BCG Danish) isolates were studied. The M. bovis BCG Moscow vaccine was used as reference. Nitrate reduction test and pyrazinamide drug susceptibility tests were performed to confirm the isolates. The DNA was sequenced using the Illumina Miseq platform. Data was analysed using CLC Genomics workbench7 and Mauve 2.3.1; the sequences were compared with reference genome, M. bovis BCG Copenhagen (Danish).

The biochemical assays confirmed that the clinical and the reference strains were M. bovis. Deletion of the RD1 region, confirmed the isolates as BCG. Deletions, insertions and single nucleotide variation were identified on the target isolates sequences in reference to the M. bovis BCG Copenhagen genome; the target isolates could have mutated.

Variation between the clinical and reference M. bovis BCG genome was confirmed. This could have contributed to the vaccination complications. Further studies are required to identify specific gene mutations that may contribute to dissemination in immunocompromised.

**Conclusion**

Understanding the dynamic nature of chronic HBV infection is essential for management of HBV carriers.

**Methods**

A cross sectional study conducted among 331 patients presenting with jaundice, elevated liver function tests, or previous history of liver disease, attending Princess Marina Hospital, Gaborone, Botswana (February 2015 to July 2016). Serum screened for presence of Hepatitis B surface antigen (HBsAg), core antibody (anti-HBc), surface antibody (anti-HBs), Hepatitis B e antigen (HBeAg) and antibody to Hepatitis B e antigen (anti-HBe) using enzyme linked immunosorbent assay techniques (ELISA). Risk factors assessed using logistic regression techniques.

**Results**

One-hundred-and-eighty-nine (57%) of the 331 participants were females, with median age of 40 years (Q1, Q3: 31–53.5). Serological profiles show: HBsAg positive in 52.57% (95% CI 0.47–0.58); anti-HBc in 56.5% (95% CI 0.51–0.61); HBeAg 8% (95% CI 0.05–0.11) and anti-HBe 34.74% (95% CI 0.22–0.31). Antibodies to HBsAg detected in 115 (34%, 95% CI 0.29–0.40) samples. Using serological profiles, participants were classified into acute infection or with chronic active hepatitis (based on the presence of HBeAg (n = 40, 12.8%); chronic hepatitis B patients (n = 47, 14.2%); presence of immunity due to natural infection (n = 53; 16%); probably hepatitis B infection with anti-HBs below level of detection (n = 4; 1.21%). Anti-HBsAg alone indicating prior vaccination or infection (n = 46, 13.9%); isolated anti-HBc detected in 22 (6.65%) probably due to resolved infection, low level chronic infection or false positive implying they are susceptible. Fifty-two (15.7%) patients were negative for all the serological markers tested. Mean alanine aminotransferase (ALT), aspartate aminotransferase (AST), serum bilirubin, alkaline phosphatase values were 230.8 (SD 410.58), 311.98 (SD 511.53), 83.01 (SD 7.47), and 287.55 (SD 395) respectively. All health professionals in the study were HBsAg positive.

**Conclusion**

Half of the patients with jaundice, history of jaundice or increased aminotransferase levels have hepatitis B infection. Screening for HBV infection and early management is advised. Health professionals need to be vaccinated compulsorily and anti-HBs titre evaluated.
ID: 8561

**Category:** Infectious Diseases (IDSSA)

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**ABSTRACT TITLE:** Mycobacterium Tuberculosis Heparin Binding Haemagglutinin Adhesin (HBHA) and Curli Pili (MTP) are Essential for In Vitro Growth, But Not Viability and Biofilm Production

**Introduction**

*M. tuberculosis* (M.tuberculosis) gene knockout mutants generated by phage-mediated genetic approaches allow for the extensive study of specific encoded proteins and their role in persistence and virulence. Mycobacteria display multiple proteins with primary adhesin function that interact with host cells. The major adhesin is the heparin-binding haemagglutinin adhesin (HBHA), that facilitates dissemination of the pathogen. *M.tuberculosis* also produces pili proteins (MTP) during pathogenesis. This study proposed to establish the contribution of HBHA and MTP, in the growth, viability and biofilm production of *M.tuberculosis*.

**Methods**

A double mutant strain deficient in the adhesin molecules encoded by genes Rv3312A (mtp) and Rv0475 (hbhA) was generated by phage-mediated specialised transduction, followed by complementation of both the double and single mutants with episomal plasmids expressing the deleted genes. The phenotypic contribution of these molecules was assessed in vitro by Optical Density (OD 600) readings, Colony Forming Unit (CFU) counts, resazurin microplate assay (REMA) and crystal violet biofilm quantification.

**Results**

Growth rates varied among the wildtype, single and double mutant strains. However, no significant differences in the viability were observed among these strains. Biofilm production was similar between wildtype and mutant strains, alluding to possible compensatory gene regulation mechanisms in the absence of functional hbhA and mtp transcripts.

**Conclusion**

These findings demonstrate the importance of the two major adhesins in functioning together to facilitate the growth of *M. tuberculosis*. This is suggestive of their potential as biomarkers for drug or vaccine development.

ID: 8678

**Category:** Sexually Transmitted Diseases (STDSSA)

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**ABSTRACT TITLE:** The Role and Impact of Sexually Transmitted Infections (STIS) Surveillance in South Africa (2004–2016) – Where Are We Now?

**Introduction**

Since 2004 in South Africa, several successful attempts for comprehensive STI Surveillance were initiated: a) National microbiological surveillance in all provinces between 2004–2014 (syndromic STI aetiologies and antimicrobial resistance to *Neisseria gonorrhoeae*); b) Cross-sectional sentinel surveillance syndrome aetiologies and HPV genotypes between 2014–2015 in 36 healthcare facilities across South Africa and presently on GERMS-SA platform with incorporates STI aetiologies, antimicrobial resistance and HPV genotyping. We are reporting the impact this surveillance had on the control and management of STIs in South Africa, the progress that has been made, the new approach and challenges that this programme is facing and suggestions how it can be improved.

**Methods**

Sentinel sites for surveillance was undertaken yearly or biennially across the country in all nine provinces. Selection was based on the total volume of STI patient seen yearly in a particular clinic and presently on number of MUS cases (≥ 25 per month). All patients 18 years and above complaining of one of the three syndromes were recruited with informed consent. Genital and blood samples were collected in addition to screening of 100 asymptomatic women aged 18–20 for HPV genotyping.

**Results**

The positive impact on STI control and management illustrated by the change and introduction of new STIs management guidelines in 2008 and recently 2015 in support of STI syndromic management. The information gathered prompted the NDOH to rethink about a new approach to syphilis and the initiation of a new comprehensive STI strategy for South Africa.

**Conclusion**

The impact, challenges, shortcomings and variables identified that hinder progress on STI surveillance will be discussed in detail in this presentation, as well as the detailed new approach incorporated into GERMS-SA.

ID: 8327

**Category:** Clinical Microbiology (SASCM)

**Permission:** Yes

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TRAK HELPER: SOFTWARE TO AUTOMATE MANUAL ENTRY OF MICROBIOLOGY TEST RESULTS ON THE LABORATORY INFORMATION SYSTEM

Introduction

Despite increasing laboratory automation, many routine clinical microbiology test results are generated manually. Manual entry of results in the laboratory information system (LIS) can be laborious and time-consuming, involving many user interface steps for a single result. At the National Health Laboratory Service microbiology laboratory at Groote Schuur Hospital, we developed and implemented Trak Helper software to “overlay” TrakCare Lab LIS (InterSystems, Cambridge, MA). Trak Helper is written in AutoHotkey, a free open-source macro scripting language for Microsoft Windows. Trak Helper consists of > 30 applications and shortcuts that automate manual LIS tasks by emulating user interface steps (mouse clicks, keyboard entries). We evaluated the benefit of Trak Helper in reducing manual steps and hands-on time on a subset of test results.

Methods

At our laboratory, approximately 80 negative (no growth) CSF culture results are entered manually per day. We compared the number of steps and time taken to enter a negative CSF culture result manually versus using Trak Helper. Two technologists were timed entering all negative CSF culture results over six to 11-day periods per method.

Results

Trak Helper reduced the number of steps required to enter a negative CSF culture result from twelve or six steps (depending on the age of the sample) to a single step. For Technologist 1, time per sample was reduced from a median of 39 seconds (IQR 38–41) to 23 seconds (21–29) (41% improvement, p = 0.002). Improvement for technologist 2 was 33% (p = 0.028). Median hands-on time per day was reduced from 54 minutes to 31 minutes for Technologist 1, and from 45 minutes to 30 minutes for Technologist 2.

Conclusion

A simple software application reduced hands-on time for negative CSF culture result entry on the LIS by up to 41%, saving the laboratory up to 15 person-days/year of skilled technologist time. Additional savings could be gained by further software development and wider implementation.

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ABSTRACT TITLE: CLINICAL CHARACTERISTICS ASSOCIATED WITH OUTCOME IN PATIENTS WITH COMMUNITY- VERSUS HOSPITAL-ASSOCIATED METHICILLIN-RESISTANT STAPHYLOCOCCUS AUREUS BLOODSTREAM INFECTION IN FIVE SOUTH AFRICAN HOSPITALS

Introduction

Methicillin-resistant Staphylococcus aureus (MRSA) is a major cause of healthcare-associated infections. We compared clinical characteristics and outcome of patients with community-associated (CA) MRSA and healthcare-associated (HA) MRSA infections.

Methods

We analysed active, laboratory-based surveillance data for S. aureus bloodstream infection (BSI), 2013 through 2015, at five sentinel hospitals. A case of MRSA BSI was defined as any patient with S. aureus, non-susceptible to oxacillin or cefoxitin, cultured from blood within a 21-day period. CA-MRSA was defined as MRSA isolated within 48 hours of hospital admission and no admission within the previous year. A reference laboratory confirmed isolate identification and antimicrobial susceptibility. We used multivariable logistic regression to evaluate factors associated with CA-MRSA versus HA-MRSA, and compare in-hospital outcome between the two groups.

Results

Among 557 patients with MRSA BSI, 44 cases (7.9%) had CA-MRSA and 513 (92.1%) had HA-MRSA; 60% were male (335/555). The median age was three years (IQR: 0–42), with no difference between the two groups (p = 0.816). Thirty-nine per cent of CA-MRSA and 47% of HA-MRSA cases (239/512) were aged < 1 year. HIV-infected patients were three times more likely to have CA-MRSA than HA-MRSA, after adjustment for age and sex (aOR: 3.3; 95% CI: 1.1–9.3). Fever (body temperature ≥ 37.5 °C) at diagnosis was associated with HA-MRSA (aOR: 0.2; 95% CI: 0.05–0.7). Crude in-hospital mortality was 41.2%. There was no association between...
MRSA category and outcome. Increasing age (age 50–59 aOR: 3.14; 95% CI: 1.61–6.13, age ≥ 60 aOR: 5.15; 95% CI: 2.50–10.60, compared to age < 1 year), cardiac arrest (aOR: 23.03; 95% CI: 2.88–183.93), mechanical ventilation (aOR: 1.89; 95% CI: 1.21–2.95) and prior MRSA infection (aOR: 3.44; 95% CI: 1.38–8.53) were associated with mortality among all cases.

Conclusion
HIV infection and fever could predict CA-MRSA and HA-MRSA, respectively. There was no outcome difference between CA-MRSA and HA-MRSA cases.

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ABSTRACT TITLE: RPOB MUTATIONS CAUSING DISCORDANT RIFAMPICIN SUSCEPTIBILITY IN MYCOBACTERIUM TUBERCULOSIS IN KWAZULU-NATAL, SOUTH AFRICA

Background
The rapid diagnosis of drug resistant tuberculosis is limited by the discrepancy in drug susceptibility results between molecular and genotypic assays. Our aim was to determine the prevalence of phenotypically susceptible rpoB mutations as well as to characterise them so as to define optimal ways of diagnosis.

Methods
Discordant clinical isolates showing rifampicin resistance on GenoType MTBDRplus while susceptible on 1% agar proportion method using Middlebrook 7H10 agar were collected from KwaZulu-Natal provincial TB laboratory between May and December 2014. These were tested using agar dilution (AD) in Middlebrook 7H10 and automated Bactec Mycobacteria Growth Indicator Tube (MGIT) 960 MICs. Additionally, subsets of stored discordant as well as 40 fully susceptible isolates were included. Sequencing was performed on all discordant isolates.

Results
Discordant isolates constituted 5% (67) of 1324 rifampicin resistant isolates detected by MTBDRplus during the study period. Out of 90 discordant isolates (including 23 stored isolates); sequencing confirmed the presence of rpoB mutations in all isolates. The most frequent mutation was Gln513Pro at 24 (26%), followed by 17 (19%) Asp516Val and 13 (14%) Asp516Tyr. The majority of MICs were between 0.5 and 2 µg/ml. (84% for AD and 74% for MGIT). Isoniazid resistance was found in 34 (37%) isolates and it was associated with higher rifampicin MICs on the AD. Among the 40 fully susceptible isolates (sensitive to isoniazid and rifampicin -wild type rpoB gene) 38 (95%) had a rifampicin MIC of ≤ 0.25µg/ml.

Conclusions
Given the high burden of disease in our setting, the isolates with discordant genotypic and phenotypic susceptibility results are not uncommon. Their association with MICs around critical concentration plus isoniazid resistance raises concerns regarding clinical implications as these may represent pre-MDR TB phenomenon. The frequent occurrence of an uncommon mutation, 513Pro warrants further investigation in order to define its public health implications.

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ABSTRACT TITLE: INVESTIGATION OF A POSSIBLE RALSTONIA INSIDIOSA OUTBREAK IN A REGIONAL HOSPITAL IN THE WESTERN CAPE

Introduction
Ralstonia insidiosa is a Gram-negative bacillus that has rarely been associated with infections in humans. We report a pseudo-outbreak of this organism in a regional hospital in the Western Cape.

Methods
We performed an outbreak investigation following a clustering of 18 isolates on blood cultures received over a two-month period. The investigation included laboratory diagnosis and environmental sampling. Blood culture isolates were identified by Vitek® 2 automated identification system and confirmation performed on two isolates by mass spectrometry. Environmental samples were confirmed on Vitek ® 2 and mass spectrometry.
Results

Eighteen blood culture isolates were identified as Cupriavidus pauculus by the Vitek 2 platform. All index blood cultures were taken from casualty or paediatric wards and had similar antibiograms. No patients had symptoms suggestive of C. pauculus infection and there was no epidemiological link between patients. The mass spectrometry platform identified the two patient samples as R. insidiosa. We subsequently identified R. insidiosa during initial environmental sampling from diluted chlorhexidine skin cleaning solution. We confirmed the source to be the chlorhexidine solution from a storage container in pharmacy. A private laboratory confirmed R. insidiosa from this sample. No relevant organisms were found in any other solutions, blood culture bottles or tap water.

Conclusion

Incorrect diluting and storage practices of chlorhexidine lead to the pseudo-outbreak of R. insidiosa.Ralstonia spp and C. pauculus are two closely related Gram-negative bacteria that are almost indistinguishable with automated lab identification systems. Identification of these organisms in the microbiology laboratory should alert clinicians to a potential pseudo-outbreak.

ID: 8438

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ABSTRACT TITLE: WATER SUPPLY AND SANITATION CONDITIONS IN RURAL SOUTHERN MOZAMBIQUE AND ITS ASSOCIATION WITH MORBIDITY INDICATORS DURING 2012-2015

Introduction

Water and sanitation (WASH) are major health determinants, responsible for 5.7% of worldwide disease burden. However, the debate about the effect of water quality and sanitation in morbidity such as diarrhoea is still ongoing. The aim of this study is to describe access to improved WASH infrastructure at a community and household level, as defined by the Joint Monitoring Programme, during 2012-2015 in the Manhiça Health Research Centre (CISM) study area and evaluate its association with morbidity indicators.

Methods

We conducted a retrospective cohort study. We included all children under 15 living in the study area during the period 2012-2015 (N = 61,900). Children were followed up until they moved from the study area, turned 15 or until 2015. WASH household data was obtained from the CISM demographic surveillance system in the Manhiça district. Clinical data was obtained from CISM round-the-clock morbidity surveillance system covering outpatient and hospital admissions at the Manhiça District Hospital and rural health posts. A negative binomial regression model using Wald test was performed to assess the minimum community-based incidence rates for every morbi-mortality indicator. WASH community herd protection for morbidity indicators was assessed spatially using QGIS.

Results

Water access conditions enhanced in 2015 (85% children living in improved conditions) compared to 2012 (73%). However, 75% of children remained in unimproved sanitation conditions during this period. Spatial distribution of unimproved water and sanitation facilities showed to be clustered. Access to unimproved sanitation and water facilities were associated to higher rates of malaria parasitemia, which adjusted incidence rate increase was doubled. Other studied morbidity indicators such as diarrhoea, malnutrition and dehydration did show weaker evidence of association between the use of unimproved water and sanitation facilities and its adjusted incidence rate. We did not observed community herd protection by owing improved water and sanitation facilities.

Conclusion

Achieved results are useful to inform sector-related decision-making processes and ultimately improve access to safe drinking water and sanitation in rural southern Mozambique.

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ABSTRACT TITLE: A POINT-PREVALENCE STUDY OF ADULT IN-PATIENTS TO DETERMINE COLONISATION WITH CARBAPENEMASE-PRODUCING ENTEROBACTERIACEAE IN TYGERBERG HOSPITAL, WESTERN CAPE, SOUTH AFRICA

Introduction

The prevalence of carbapenemase-producing Enterobacteriaceae (CPE) has been increasing around the world in recent years, but data regarding the prevalence and clinical significance of CPE in Africa is not well documented. To determine the current prevalence of CPE colonisation a point-prevalence study was performed at TBH’s adult in-patient wards. The need for routine screening and risk factors associated with colonisation were also investigated.

Methods

A total of 439 adult in-patients from 46 different wards in TBH were screened with rectal swabs during a three-month period. Risk factors...
were recorded for each patient. Patient samples were tested for carbapenem-resistant Enterobacteriaceae using a MacConkey agar plate with an ertapenem disc and the Vitek® 2 system. Preliminary positive samples were tested for carbapenemase-production with a multiplex PCR containing primers for five associated genes.

Results
Six hundred adult in-patients were approached for participation in this study and 161 (25.33%) withheld consent. Of the 439 patient samples collected, 12 (2.7%) screened positive for carbapenem-resistance and of these, one (0.22%) sample was positive for carbapenemase-production. This patient was colonised with a *Klebsiella pneumoniae* organism harbouring a NDM-1-type gene and had no commonly reported risk factors associated with CPE colonisation.

Conclusion
It seems that the routine screening of adult patients for CPE-colonisation during admission to TBH is unwarranted. Risk factors associated with CPE-colonisation in this specific setting require further clarification.

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ABSTRACT TITLE: ENGINEERING NEWCASTLE DISEASE VIRUS AS VACCINE DELIVERY SYSTEM FOR ROTAVIRUS VP7 AND NSP4

Introduction
Rotavirus (RV) is the leading cause of severe dehydrating diarrhoea in young children. Due to the zoonotic nature of RV infections, the young of various animals, including livestock, are also affected, resulting in devastating economic losses. Recent reports have described animal-human reassortant strains, such as bovine-human reassortants. These interspecies transmitted strains contribute to the diversity of rotavirus strains circulating in humans and emphasise the need for a One Health approach in RV control. Infection occurs in calves 2-4 weeks of age indicating the protective role of maternal antibodies in the colostrum immediately postpartum. Vaccination remains the only existing prophylactic measure against infection and aims to increase the level of anti-RV antibodies in the colostrum. Newcastle disease virus is an attractive vaccine delivery vector for mammals because of its natural host-range restriction in non-avian species. It is also antigenically distinct from common animal and human pathogens and would, therefore, not be affected by a pre-existing immunity against NDV.

Methods
Recombinant NDVs were constructed to express the outer-capsid glycoprotein (VP7) and the enterotoxin non-structural protein (NSP4) of RV. The recombinant viruses were rescued in BSR-T7 cells and passaged in SPF embryonated chicken eggs. RNA was extracted from allantoic fluid and sequenced using the Illumina MiSeq platform for verification of successful gene insertion. Protein expression was verified using immunoperoxidase monolayer assay (IPMA) and western blot analysis.

Results
Recombinant NDVs containing open reading frames encoding rotavirus VP7 and NSP4 were constructed and successfully rescued from BSR-T7 cells. The viability of recombinant NDVs were confirmed using the hemagglutination assay, virus titrations and sequencing. Expression of rotavirus proteins was subsequently confirmed from BSR-T7 and MDBK cells.

Conclusion
This is the first step towards a possible vaccine vector for RV infection in bovine for a One Health approach to RV control.

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ABSTRACT TITLE: WHOLE-GENOME SEQUENCING AND MICROBIOME ANALYSIS TO IDENTIFY AND CHARACTERISE SHIGA TOXIN-PRODUCING ESCHERICHIA COLI FROM THE STOOL OF A PATIENT PRESENTING WITH HAEMOLYTIC URAEMIC SYNDROME

Introduction
Microbial whole-genome sequencing (WGS) holds great promise for enhancing diagnostic and public health microbiology. WGS analysis of bacterial cultures is well established, but when applied directly on clinical samples is more challenging. We describe the use of WGS and microbiome analysis applied directly on a culture-negative stool sample, from a patient presenting with haemolytic uraemic syndrome (HUS), to successfully identify and characterise Shiga toxin-producing Escherichia coli (STEC).

Methods
Total DNA was extracted from stool samples using the MagNA Pure LC Total Nucleic Acid Isolation Kit. DNA was further processed using the NEBNext Microbiome DNA enrichment Kit. Enriched microbial DNA samples were subjected to WGS using Illumina MiSeq next generation sequencing technology. Raw sequencing data were analysed against microbial databases using SURPI (sequence-based ultra-rapid pathogen identification), a computational bioinformatics pipeline for pathogen identification utilising complex metagenomic sequencing data. Raw sequencing data were further analyzed using CLC Genomics Workbench Software; trimmed reads were assembled de novo. Assembled data were analysed using bioinformatics pipelines available at the Center for Genomic Epidemiology (CGE) (http://cge.cbs.dtu.dk/services/).

Results
SURPI analysis against bacterial databases returned an overwhelming and convincing match to E. coli; additionally, SURPI analysis against viral databases found matches to phages which house stx2 genes (markers for STEC). Analysis of assembled data at the CGE identified STEC with the following characteristics: serotype O26:H11; presence of eae, stx2a and stx2b virulence genes; multilocus sequence typing (MLST) sequence type 21.

Conclusion
WGS and microbiome analysis was able to convincingly identify and characterise STEC from a culture-negative stool sample. If implemented in real-time, WGS applied directly on clinical specimens can drastically reduce diagnostic times and rapidly provide clinically relevant and actionable public health information. This has significant patient management and huge public health benefits, especially for enhancing turn-around times for the diagnosis and characterisation of outbreak-prone pathogens including STEC.

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ABSTRACT TITLE: PHARMACOKINETICS OF RIFAMPIN AND ISONIAZID DURING PREGNANCY AND POSTPARTUM IN SOUTH AFRICAN WOMEN

Introduction
Physiological changes during pregnancy may alter drug pharmacokinetics. Trimester differences in rifampin (RMP) and isoniazid (INH) exposure have not been described. We explored the effects of pregnancy gestation on RMP and INH pharmacokinetics in tuberculosis-infected women.

Methods
P1026s is a prospective study of antiretroviral and anti-tuberculosis pharmacokinetics in HIV-infected and uninfected pregnant women. Intensive 24-hour pharmacokinetic profiles of RMP and INH were performed during the 2nd trimester (2T), 3rd trimester (3T), and 2-8 weeks postpartum (PP). Daily anti-tuberculosis tablets were given according to WHO-recommended guidelines. RMP and INH plasma concentrations were measured using High Performance Liquid Chromatography(HPLC); detection limits being 0.117 µg/ml and 0.098 µg/ml respectively. The pharmacokinetic parameters were characterised using noncompartmental analysis and compared to published non-pregnant South African adult data.

Results
Preliminary pharmacokinetic data are available for ten South African participants; seven African, two mixed descent, and one Indian. Eight women were HIV-infected (seven on efavirenz and one on lopinavir/ritonavir). Median age at 3T was 31 years (range 21-40) and median weight at 3T was 58.6 kg (range 49-99). Median gestational age at delivery was 38 weeks (range 36-41). RMP and INH pharmacokinetic data were available in 5, 8 and three women in 2T, 3T, and PP. Compared to a non-pregnant South African adult cohort (45% male, 10% HIV-infected not receiving antiretrovirals, McIlleron et al. 2006), RMP exposure was similar or higher in 2T and 3T. INH exposure was below the 25th percentile across all stages of pregnancy. Small sample size and unavailable comparator raw dataset prohibited formal statistical testing.

Conclusion
RMP concentrations in pregnancy compared well to non-pregnant concentrations. INH exposure was reduced throughout pregnancy.
If confirmed with larger sample size, an increased dose of INH may be needed during pregnancy.

**ID: 8356**

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**ABSTRACT TITLE:** URINARY TRACT INFECTIONS IN PREGNANT WOMEN IN KZN: BACTERIAL AETIOLOGY AND SUSCEPTIBILITY

**Introduction**

Urinary tract infection (UTI) is one of the commonest infections during pregnancy. Untreated, it can cause significant maternal and perinatal morbidity and mortality. Challenges while treating UTIs in pregnancy include foetal protection and resistance development of uropathogens. Currently, the Essential Drug List (EDL) recommends amoxicillin-clavulanic-acid to treat cystitis and ceftriaxone/ gentamicin for pyelonephritis. Nitrofurantoin is not included in the recommendation. Cefuroxime has been removed from the EDL. Aim: To determine common pathogens causing UTI in pregnancy and their antibiotic susceptibility patterns.

**Methods**

A retrospective analysis of laboratory data for positive urine specimens from obstetric departments of six KZN hospitals during 2012-2016 was performed. Identification and susceptibility testing was performed using the Vitek2 system. Antibiotics tested were amoxicillin, cephalexin, ceftriaxone, nitrofurantoin, amoxicillin-clavulanic-acid, ciprofloxacin, gentamicin, and cotrimoxazole. Results were interpreted according to Clinical and Laboratory Standards Institute (CLSI) guidelines.

**Results**

From 5 855 urine specimens, the commonest isolate was E coli, 2 984 (51.2%), K pneumoniae, 718 (12.2%), C albicans, 377 (6.4%), Candida species, 292 (3.2%), Group B Streptococci(GBS), 225 (3.8%), E faecalis, 218 (3.7%), and 17.7% various others. E coli displayed 71.9% resistance to ampicillin, cotrimoxazole (65.2%), cephalothin (23.7%) and ciprofloxacin (12.3%). Resistance remained low to cefuroxime 4.1%, ceftriaxone 1.8%, amoxicillin-clavulanic-acid 5.3%, gentamicin 5.5% and nitrofurantoin 2.8%. K pneumoniae was similar. GBS displayed 99.5% penicillin susceptibility. E faecalis displayed 92% susceptibility to ampicillin, amoxicillin-clavulanic-acid, and nitrofurantoin.

**Conclusion**

E coli is unsurprisingly the most common cause of UTI in pregnancy (KZN), with > 94% susceptibility to cefuroxime, ceftriaxone, amoxicillin-clavulanic-acid, nitrofurantoin and gentamicin. GBS remains prevalent, and susceptible to penicillin. E faecalis remains susceptible to ampicillin and nitrofurantoin. Candida featured prominently in this analysis, and its significance needs further studying. To spare ceftriaxone, nitrofurantoin could be an alternative in treating cystitis and cefuroxime for pyelonephritis. As drug resistance is evolving, routine surveillance is necessary to provide updated information on recommended antibiotic use.

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**ABSTRACT TITLE:** VIRULENCE PROFILES OF ENTEROHAEMORRHAGIC ESCHERICHIA COLI O157:H7 ISOLATES FROM CLINICAL SPECIMENS AND ENVIRONMENTAL SAMPLES IN GAUTENG REGION, SOUTH AFRICA

**Introduction**

Enterohaemorrhagic Escherichia coli (EHEC) O157:H7 or Shiga-toxins producing E. coli is an important gastrointestinal pathogen causing bloody diarrhoea and more severe diseases such as haemolytic uremic syndrome, an important cause of acute kidney failure in children. The reservoirs for EHEC O157:H7 are healthy ruminant animals. Humans acquire EHEC O157:H7 by direct contact with animal or via the ingestion of contaminated food or water. In South Africa, limited information is available on characterisation of EHEC O157:H7. The aim of this study was to determine the virulence...
profile of EHEC O157:H7 isolates from clinical specimens and environmental water samples in the Gauteng region of South Africa.

Methods
Clinical stool specimens and environmental water samples were collected from October 2016 to June 2017. Samples were cultured on selective chromogenic media and presumptive EHEC O157:H7 were confirmed by using a latex agglutination test. Isolates were characterised using multiplex PCR assays to screen for the uidA, rfbEO157 and hlyC7 genes that encode for β-glucuronidase, somatic-O and flagellar-H antigens, respectively and for the virulence genes: Shiga-toxins 1 and 2 (Stx-1 & Stx-2), enterohaemolysin (hlyA) and E. coli attaching and effacing lesion (eae).

Results
A total of 213 samples were analysed of which 10.7% (23/213) were identified as EHEC O157:H7 by M-PCR assays. Run-off water represented 86.9% (20/23) of isolates and stool specimens represented 13% (3/23). The Stx-1, Stx-2, hlyA and eae genes were present in 7.6% (2/23), 15.3% (4/23), 3.8% (1/23) and 11.5% (3/23) of isolates, respectively. Virulence genes were not detected in 47.8% (11/23) of isolates.

Conclusion
This study reported the presence of EHEC O157:H7 in stool specimens and environmental water samples, showing the under reporting and miss diagnosis of 13% of stool specimens due to the diagnostic screening test used. Environmental surveillance plays an important role in reducing the potential health risk to consumer of untreated water. The inclusion of molecular testing for EHEC O157:H7 of stool specimen can improve the detection of this important pathogen in this clinical setting.

ABSTRACT TITLE: RECONSIDERING CLOSTRIDIUM DIFFICILE TESTING IN SA – A CRITICAL LOOK AT THE NEED FOR A TWO-STEP ALGORITHM!

Introduction
Current Clostridium difficile diagnostic assays are inadequate as stand-alone tests. Molecular assays show greater sensitivity compared to enzyme immune assays (EIAs) in detecting toxigenic strains however, lack the ability to discriminate between active disease (toxin production) and colonisation.

Methods
A retrospective analysis was performed using specimens submitted for C. difficile testing during the period 5 May 2016 to 30 April 2017 from the private health sector in Gauteng. Results from samples tested with both Xpert C. difficile (Cepheid) and the C Diff Quick Check Complete (Alere) were included. Sensitivity, specificity, positive predictive value (PPV) and negative predictive value (NPV) of the glutamate dehydrogenase (GDH) antigen was determined, using PCR as the reference. Toxin detection (TcdA/B) was expressed as a percentage of PCR positives. Association between sample consistency (solid, semisolid and liquid) was determined.

Results
A total of 165 samples were included for analysis of which 113 (68.5%) were negative by PCR testing. Toxin was detected in 14 (30%) of the PCR positives. Diagnostic performance of the GDH antigen demonstrated: sensitivity (84%), specificity (91%), PPV (78%) and NPV (93%). The majority (53%) of the samples were semi-formed and no association between the consistency and toxin production or carriage could be determined.

Conclusion
Toxin positivity correlates best with clinical outcome and assists in targeting patients for treatment. Up to 70% of patients may have received unnecessary antimicrobial therapy. The sensitivity of the GDH antigen to detect and inability to exclude disease limits its use as a screening test compared to PCR, seven (4%) of patients would have been missed in the study group with EIA-based testing alone. Our findings support an algorithmic approach to diagnosis of C. difficile associated diarrhea. We recommend a PCR based assay with subsequent toxin detection of positives to distinguish between active disease and colonisation.

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ABSTRACT TITLE: COMPARISON OF DIAGNOSTIC TESTS FOR CLOSTRIDIUM DIFFICILE IN A PRIVATE SECTOR LABORATORY IN GAUTENG, SOUTH AFRICA

Introduction
The diagnosis of Clostridium difficile infection is a frequently debated topic due to the wide range of diagnostic strategies. The objective of this study was to compare diagnostic tests for C. difficile performed in a private laboratory in Gauteng.

Methods
A convenience sample of 100 consecutive liquid PCR positive stool specimens (BD MAX™ Cdiff) were selected and further tested for GDH and toxin A and B (CerTest Clostridium difficile, BIOTEC S.L.). All were also cultured using selective agar (chromID® C. difficile agar, Biomerieux). Culture positive isolates were confirmed to be toxigenic by use of PCR (Roche LightMix® Kit Clostridium difficile, TibMolBiol) which also detected the 18-bp sequence deletions in the tcdC gene associated with hypervirulence.

Results
Of the 100 PCR positive stools, 95 were culture positive and toxigenic. Of these 90 were GDH positive. 36 were toxin positive of which the majority, 72% (26/36), were positive for Toxin A and B. 2 strains showed the 18-bp deletion in the tcdC gene, indicative of a hyper-virulent strain.

Conclusion
Most PCR positive stools selected were culture positive for toxigenic strains. The detection of toxin in only 36 of 95 C. difficile culture positive stools is a higher rate of discrepant results than reported in most studies. The absence of C. difficile toxins in EIA tests should not rule out the possibility of CDI. As toxin detection is dependent on the level of toxin production at sample collection and the sensitivity of the test, this magnitude of discrepancy may also be due in part to the kit used in this study. The use of PCR remains important.

ABSTRACT TITLE: DIVERSITY OF THE GUT, VAGINAL AND ORAL MICROBIOME AMONG PREGNANT WOMEN IN SOUTH AFRICA WITH PRE-ECLAMPSIA

Introduction
Pre-eclampsia is a multisystem disorder better known as a condition of high blood pressure and proteinuria associated with pregnancy. It is considered a leading cause of maternal as well as fetal/neonatal morbidity and mortality. Pre-eclampsia has a complex aetiology of which the exact cause remains unknown; however, its association with exaggerated systemic inflammation suggests microbial infection and a bacterial source to play a possible role.

Methods
Oral, vaginal and rectal samples were collected from ten primiparous pregnant women in both the normotensive and pre-eclampsia group. Targeted 16S rRNA next generation sequencing (NGS) was performed using an Illumina MiSeq platform. Analysis of the microbial community was done using Quantitative Insights into Microbial Ecology (QIIME) and statistical analysis was inferred using R.

Results
Alpha (within sample) diversity analysis showed the gut and oral microbiome to be more diverse than the vaginal microbiome. Significant differences in alpha diversity of the gut and oral microbiome between treatment groups was not confirmed. However, the vaginal microbiome of women with pre-eclampsia had a significant increase in evenness as observed with the Shannon Index (Kruskal Wallis chi-squared test: P = 0.0472). The evaluation of beta (between sample) diversity by principal coordinate analysis (PCoA) and the UniFrac distances revealed significant community differences of the vaginal microbiome in pre-eclampsia compared to normotensive pregnant women (Adonis: P = 0.05). Lactobacillus iners was identified as the predominant species in the vaginal microbiome, however, a lower abundance of this species was detected in the pre-eclampsia group. Although the gut and oral microbiome did not reveal significant separation of these microbial communities in the pre-eclampsia group compared to normotensive pregnant women, a change in the relative abundance of several taxa was observed.

Conclusion
To our knowledge, this is the first study to investigate the diversity of the gut, oral and vaginal microbiome among pregnant women.
in South Africa with pre-eclampsia. Our results indicate an altered vaginal microbiome in pregnant women diagnosed with pre-eclampsia.

**ID: 8681**

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**ABSTRACT TITLE:** EVALUATION OF PSEUDOMONAS AERUGINOSA ANTIMICROBIAL SUSCEPTIBILITY DATA OBTAINED FROM THE NHLS, LABORATORY INFORMATION SYSTEM WITH THE REFERENCE LABORATORY AT NICD, 2014 TO 2015, SOUTH AFRICA

**Introduction**

Broth microdilution has been regarded as the reference method for antimicrobial susceptibility testing (AST) of Pseudomonas aeruginosa. However, the accuracy of automated AST systems, disk diffusion and Etest have become a concern. We aimed to compare AST results among clinical isolates of P. aeruginosa over a two-year period using data from a laboratory information system and the national reference laboratory.

**Methods**

We analysed data from patients admitted to 12 public sector sentinel hospitals from 2014 to 2015. Cases were identified through data extracts from the National Health Laboratory Service (NHLS), Corporate Data Warehouse (CDW) and the laboratory-based antimicrobial surveillance programme at the Centre for Healthcare-Associated Infections, Antimicrobial Resistance and Mycoses (CHARM). A case was defined as any patient with P. aeruginosa isolated from a blood culture specimen. A positive result obtained after 21 days of the first blood culture result was regarded as a new case. We compared categorical data (susceptible or non-susceptible including intermediate and resistant) for each method based on Clinical and Laboratory Standards Institute guidelines against six antimicrobial agents (cefepime, ceftazidime, colistin, imipenem, meropenem and piperacillin/tazobactam). Data from the NHLS, CDW contained AST data from the Vitek 2, MicroScan, disk diffusion or Etest and surveillance data from CHARM contained AST data for MicroScan and/or Sensititre.

**Results**

We assessed P. aeruginosa AST data for 1067 routine isolates and 571 surveillance isolates. Cefepime susceptibility was similar between NHLS (73% [754/1038]) by NHLS and the reference laboratory (73% [414/571]), meropenem susceptibility was observed in 72% (751/1046) by NHLS versus 69% (392/571) by the reference laboratory and piperacillin/tazobactam susceptibility was observed in 69% (706/1028) by NHLS versus 73% (414/571) by the reference laboratory.

**Conclusion**

Overall, we observed acceptable categorical agreements for AST from both routine NHLS laboratories and CHARM. We recommend that NHLS AST data for P. aeruginosa isolates can be used for reporting purposes.

**ID: 8265**

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**ABSTRACT TITLE:** PREVALENCE AND PROFILE OF TUBERCULOSIS IN LIVER DISEASES: ZILCH BUT EXCEPTIONS

**Introduction**

Tuberculosis has a complex relationship with liver diseases, as the diseases may affect liver in disseminated cases with military disease, can be reactivated in immunosuppressive conditions or could affect the liver with toxic antitubercular therapy.

**Methods**

This study is the analysis of liver disease patients admitted to our tertiary care centre from January 2011 to December 2016. Tuberculosis was diagnosed based on (i) histological evidence of caseating granulomas (ii) smear AFB positivity (iii) growth of Mycobacteria on MGIT culture (BD Microbiology Systems, Cockeysville, MD) or (iv)positive qPCR for Mycobacterium tuberculosis (MTB qPCR) (Cobas TaqMan MTB assay). Mycobacteria other than tuberculosis (MOTT) were identified by positive MPT64 ICT assay (SD Bioline) or MGIT culture positive and MTB qPCR negative samples.

**Results**

- Total no of samples: 816
- MGIT and or qPCR positive samples: 118
• Pulmonary (sputum, ET secretions, BAL) 31 (26.2%)
• Extra-pulmonary TB (ascitic fluid, pus, body fluids, lymph node aspirates, tissue, urine)
• 87 (73.7%) Male 78/118 (66.1%) Female 40/118
• Median age. 44.5 years

Conclusion
• Our study reiterates these findings showing predominance of smear negative, extra-pulmonary TB and MOTT in our patients.
• MOTT accounting for 11% cases, should be ruled out in all cases as treatment varies with this group reducing hepatotoxicity by undue ATT exposure. MPT 64 assay is a rapid and easy method for detection of MOTT.
• A combination of MGIT culture and TB PCR has additive advantage over either test alone, with cumulative sensitivity of 100%.

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ABSTRACT TITLE: THE EPIDEMIOLOGY OF CANDIDA AURIS IN SOUTH AFRICA, 2012-2016

Background
We sought to describe the epidemiology and genetic relatedness of Candida auris, an emerging multidrug-resistant invasive fungal pathogen associated with high mortality, in South Africa.

Methods
Culture-confirmed cases of C. auris infection/colonization were reported 2012-2016 by private- and public-sector laboratories across South Africa. We defined a case as a patient with culture of C. auris (or C. haemulonii if the diagnostic laboratory used a non-confirmatory method) from any specimen. For an isolate subset with confirmed molecular identification, we performed antifungal susceptibility testing, sequencing of the FKS1 and FKS2 genes (if echinocandin MIC ≥1 mg/L) and multi-locus sequence typing (MLST).

Results
1695 cases of C. auris were reported, 1406 (88%) from Gauteng province. The number of cases increased from 4 in 2012-2013 to 380 in 2015-2016. Ninety-three per cent (1545/1659) of patients were admitted to private-sector hospitals. The median age was 59 years (IQR: 42-72) and 62% (1029/1652) were male. Overall 739 (47%) isolates were cultured from sterile site specimens (blood [344, 22%], cerebrospinal fluid [2, 0.1%], fluid [56, 4%], tissue [49, 3%]), central venous catheter tips [288, 18%] and 840 (53%) were cultured from sites of potential colonization (urine [622, 39%], respiratory tract [173, 11%], skin/mucosal swabs [45, 3%]). Of 85 isolates confirmed as C. auris, the MIC 50 (mg/L) and MIC 90 (mg/L) for fluconazole, voriconazole, micafungin and amphotericin B was 128, 256; 0.12, 1; 0.06, 2 and 0.5, 1 respectively. No known FKS mutations were detected. MLST analysis grouped isolates into two clusters comprising of 83 and 2 isolates respectively.

Conclusion
There was a large increase in reported cases of C. auris since its first isolation in South Africa in 2012. C. auris strains were mostly fluconazole non-susceptible and highly related by MLST though the possibility of nosocomial transmission should be explored using more discriminatory whole genome sequencing.

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ABSTRACT TITLE: PREVALENCE AND SUSCEPTIBILITY PROFILE OF ACINETOBACTER BAUMANII ISOLATED AT HIGH CARE WARDS AT DR GEORGE MUKHARI ACADEMIC HOSPITAL IN 2015

Introduction

Antimicrobial resistance is a general public health concern, threatening the effective management and Infection control of infections. Globally, Acinetobacter baumanii has been reported as the major pathogen isolated in hospitals with high resistance to major antimicrobial agents and thus termed Multidrug resistant organisms. The organism results in high morbidity and mortality because of limited treatment options. The aim of the study was to determine the prevalence and antimicrobial susceptibility profile of A. baumanii isolated from blood culture specimens taken at selected high care wards of Doctor George Mukhari Academic Hospital in 2015.

Methods

This was a retrospective descriptive study. National Health Laboratory Services (NHLS) laboratory information system (TrakCare) was used to obtain data on A. baumanii isolates isolated from blood culture specimens in 2015 using Microsoft Excel. Only blood culture results from high care wards (6, 9, 20, 24, 39 and ICU) were filtered, repeats excluded and data on the susceptibility trends to select antimicrobial agents was analysed.

Results

A total of 630 blood culture results were retrieved during the study period. Sixty-five (10.3%) of the isolates were identified as A. baumanii. A. baumanii was more prevalent in ICU at 44.6% (29/65). The organisms showed resistance to majority of the antimicrobials tested. They exhibited resistance to aminoglycosides; with Amikacin at 29%, and gentamicin at 78%. Most of the isolates also exhibited resistance to carbapenems: Imipenem and meropenem, at 88% and 90% respectively. Fortunately, all the isolates were susceptible to colistin.

Conclusion

High level of resistance to carbapenems is worrisome since these are the initial treatment options for patients admitted in high care areas. It also poses a threat to effective management and infection prevention and control measures in our setting.

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ABSTRACT TITLE: MORPHOLOGICAL DIVERSITY OF PANTON-VALENTINE LEUKOCIDINE POSITIVE STAPHYLOCOCCUS AUREUS BACTERIOPHAGES IDENTIFIED IN SOUTH AFRICA AND NIGERIA

Introduction

Panton-Valentine leukocidin (PVL) is a bacteriophage-encoded bi-component, pore-forming leukotoxin, causing leukotoxin lysis. The presence of the PVL gene in Staphylococcus aureus has been associated with severe necrotising pneumonia and necrotising fasciitis. Ten bacteriophage types belonging to three morphologically distinct head-groups have been reported to aid in the horizontal gene transfer (HGT) of PVL genes. The aim of this study was to identify the morphological diversity of PVL positive S. aureus bacteriophages from South Africa and Nigeria.

Methods

A total of 70 previously stored isolates consisting of MRSA and MSSA isolates from South Africa and Nigeria were collected from the culture bank in the Department of Medical Microbiology, University of Pretoria/National Health Laboratory Services (NHLS). The presence of the genus (16S rRNA), species specific (nuc), methicillin-resistance (mecA) and the PVL genes were confirmed using Multiplex-PCR (M-PCR) assays. Multiplex-PCR assays were performed to identify the morphological diversity of bacteriophage types within the S. aureus isolates.

Results

The species confirmation M-PCR assay correlates with previous analysis performed on these isolates. The results indicated the icosahedral head-group II [72.6% (53/70)] was the most prevalent head-group, followed by the elongated head-group [70% (49/70)] and icosahedral head-group I [28.5% (20/70)]. All three bacteriophage head-group morphologies were detected in 5.7% (4/70) MSSA and 2.8% (2/70) MRSA isolates from South Africa and only one MRSA isolate from Nigeria.

Conclusion

While the elongated head-group has been reported as the most prevalent in Italy, an equal distribution of bacteriophage morphology has been reported in China. This is in contrast with the results found in this study. The high prevalence of the icosahedral head-group II bacteriophages carrying the PVL genes, suggests geographic variation according to phage-lineage types and that these bacteriophages may be easily transferred to S. aureus isolates in clinical settings of the study population under investigation.

ID: 8517
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ABSTRACT TITLE: EVALUATION OF GENOTYPE® MTBDRSL (VER 2.0) AGAINST SECOND-LINE DRUG SUSCEPTIBILITY TESTING AND IMPACT OF GYRA MUTATIONS ON FLUOROQUINOLONE SUSCEPTIBILITIES

Introduction
The emergence of Mycobacterium tuberculosis resistant strains is threatening the control of multidrug resistant tuberculosis (MDR-TB) worldwide. To curb the spread, fluoroquinolones (FLQ) and second line injectable drugs (SLIDs) are used to treat MDR-TB. A recently endorsed, GenoType® MTBDRsl is used for rapid detection of resistance towards FLQ and SLIDs and guide treatment options for patients while awaiting phenotypic drug susceptibility testing (DST). The aim of the study was to evaluate Genotype® MTBDRsl assay against DST and correlate the phenotypic results with mutations.

Methods
We collected MDR-TB isolates (n = 54) from the National Health Laboratory Service, Pretoria. All isolates were sub-cultured, and DST was performed on the MGIT 960 system. The critical concentrations (CC) used were 2.0µg/ml for Ofloxacin (OFX), 0.5 and 2.0µg/ml for Moxifloxacin (MOX), 1.0 µg/ml for amikacin (AMK), 2.5 µg/ml for kanamycin (KAN) and capreomycin (CAP) drugs. GenoType® MTBDRsl assay was performed according to manufacturer (Hain LifeScience, Germany) and compared to DST.

Results
Sensitivity of the GenoType® MTBDRsl assay was 100% for OFX, MOX (0.5µg/ml) and MOX (2.0 µg/ml) with specificity of 100%, 91.84% and 88.24% respectively. Sensitivity and specificity for KAN, AMK and CAP were 75% and 100%, 75% and 100% and 50% and 98% respectively. Mutations of A90V resulted mostly in low level MOX resistance (<0.5 µg/ml) as compared to D94G with high level resistance (> 0.5 µg/ml). Mutations of A1401G (n = 2) were detected in AMK/KAN/CAP resistant cases, while G1484T (n = 1) was in AMK/KAN case.

Conclusion
GenoType® MTBDRsl assay is a useful tool and has excellent sensitivity for FLQ resistance, but moderate to high specificity for MOX and OFX respectively. Rapid detection of different mutations can guide clinicians in prescribing appropriate drugs while awaiting DST. More studies are needed, to improve the accuracy SLID rapid detection.
**ID: 8431**

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**ABSTRACT TITLE:** CHANGES IN SUSCEPTIBILITY PATTERN TO OXACILLIN IN STAPHYLOCOCCUS AUREUS ISOLATES FROM PATIENTS WITH BLOODSTREAM INFECTIONS IN SOUTH AFRICA OVER A SEVEN-YEAR PERIOD

**Introduction**

This report aimed to describe antimicrobial susceptibility and distribution of Staphylococcus aureus bacteremia (SAB) isolates among hospitalised patients in South Africa over a seven-year period. We also aimed to determine the molecular characteristics of methicillin-resistant S. aureus (MRSA).

**Methods**

The study population included patients of all ages accessing public-sector healthcare. All isolates from patients with SAB were submitted for confirmatory identification and antimicrobial susceptibility testing for the Laboratory-based Antimicrobial Resistance Surveillance (LARS) programme periodically. Susceptibility testing was performed on the MicroScan Walkaway system (Siemens Healthcare Diagnostics, USA) using the Positive MIC Panel Type 33. The LightCycler 480 II instrument (Roche Applied Science, Germany) was used for the real-time PCR amplification of the methicillin resistance determinant, mecA.

**Results**

A total of 5358 patients with SAB were identified from June 2010 to June 2017. Among 5322 of these cases with a viable isolate, confirmed molecular identification and full susceptibility profile, 1967 isolates resistant to oxacillin and confirmed as MRSA were identified. The most common SCCmec type was SCCmec type III (n=860, 43%) followed by type IV (n=609, 30%). Spa-typing of the 1149 oxacillin-resistant isolates revealed the five most common spa-types were t037 (n=556, 30%), t125 (n=214, 11%), t045 (n=95, 5%), t012 (n=58, 3%) and t064 (n=46, 2%). MLST showed the most common ST to be ST612 (CC8) followed by ST239 of clonal complex (CC8).

**Conclusion**

We found a decrease in MRSA over the seven-year period. SCCmec type III predominated; however, the presence of SCCmec type IV has become evident, indicating that this type is emerging in hospitals.

**ID: 8445**

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**ABSTRACT TITLE:** SPECIFIC DETECTION OF LABORATORY MISIDENTIFIED KLEBSIELLA VARIICOLA BY SANGER SEQUENCING FROM KNEE EFFUSION

**Introduction**

K. variicola was described as a new bacterial species in 2004, based on the phylogenetic analysis of six housekeeping (rpoB, gyrA, nifH, infB, phoE and mdh) genes. It is found mostly from plant and insects and only bloodstream infections. To date, the correct identification of Klebsiella species has not been easily achieved in microbiological laboratories because several species of this genus share similar biochemical profiles. We report a case of the female patient from Easter Cape with left knee effusion from whom K. variicola was misidentified as K. pneumoniae by automated method.

**Case**

A 44-year-old female patient from ECP with left knee effusion was admitted to the St. Elizabeth Hospital in Lusikisiki. Sterile fluid specimen, gram-negative bacilli. On overnight incubation, lactose fermenting colonies resembling Klebsiella species were isolated, which were identified as K. pneumoniae by B & C AutoSCAN using NID2 panel with 96% probability. MIC was determined by B&D Type 37 panel, which demonstrated sensitive to most antibiotics except ampicillin and ESBL -ve. DNA of the isolate was sequenced by the Sanger method at the Inqaba BioTech Lab, using an Applied Biosystems 3500xL Genetic Analyser. MLST typing was done which identified this isolate as K. variicola (matching with K. variicola DSM 15968 NCBI) which was previously identified as K. pneumoniae by phenotypic method.

**Discussion**

K. variicola, a bacterium closely genetically related to K. pneumoniae, is commonly misidentified as K. pneumoniae by biochemical tests. It is mostly isolated from environmental sources such as plants...
and insects (termites) and animals (gorilla). Misidentification of Klebsiella species by automated systems has been reported by Berry et al. S. K. varicola seems to have lower antibiotic resistance as compared to K. pneumoniae but bloodstream infections have higher mortality rate. No additional virulence factors were found for increased deaths by Maatallah et al. Our isolate was ESBL negative and sensitive to almost all antibiotics except ampicillin, but we found presence of blaCTX-M.

ID: 8454

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ABSTRACT TITLE: A TWO-YEAR STUDY OF THE INCIDENCE OF NOCARDIA

Introduction
Nocardia is a gram-positive organism found predominately in soil and the oral microflora of humans. Nocardia has been linked to a wide range of human disease. Taking into account the high level of immune compromised individuals in South Africa due to HIV infection, it would be expected to see a high level of these infections.

This study aimed to investigate the incidence of Nocardia with respect to patient age, HIV status, type of referring ward, species identified and antibiotic susceptibility in state hospitals in Gauteng over a 2-year period.

Methods
Samples sent to the Infection Control Services Laboratory, NHLS, for bacterial identification and antibiotic susceptibility results for the period 01/04/2015 to 30/07/2017 were included in the study. The genus and species of the isolates were identified through 16S rRNA sequencing. Antibiotic susceptibility testing was performed by the manual broth dilution method.

Results
36 isolates were sequenced during this period. The main Nocardia species identified were N.arthritidis and N.exalbida. 12 different species of Nocardia were identified in the study. The average age at presentation was 37.3 years. The main referral wards were the surgical and medical wards. The main type of specimen was fluid at presentation was 37.3 years. The main referral wards were the surgical and medical wards. The main type of specimen was fluid

ID: 8284

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ABSTRACT TITLE: EARLY-ONSET NEONATAL STREPTOCOCCUS GALLOLYTICUS SUBSPECIES PASTEURIANUS MENINGITIS AND SEPTICEMIA

Introduction
Streptococcus gallolyticus subspecies pasteurianus is a highly unusual pathogen in neonates. We describe a case of this organism causing early-onset meningitis and septicemia.

Case
A HIV-exposed premature infant presented on day 1 of life with respiratory distress and shock; complicated by jejunal atresia repaired on day 3. Intravenous ampicillin and gentamycin were initiated after blood cultures and a septic screen were performed. Streptococcus gallolyticus subsp. pasteurianus was isolated from a blood culture. Cerebrospinal fluid (CSF) collected on day 3 of life confirmed a diagnosis of meningitis. Bacteria were not cultured from the CSF; however, a Polymerase Chain Reaction (PCR) performed directly on the CSF demonstrated the presence of Streptococcus gallolyticus subsp. pasteurianus. The patient was treated in the neonatal ICU with 14 days of ampicillin. No neurological complications at discharge were noted.

Discussion
Streptococcus gallolyticus subsp. pasteurianus has replaced Streptococcus bovis II/2; part of group D nonenterococcal streptococci.

Bacteremia is the most common clinical manifestation of early-onset S. bovis group infection. Neonates with early-onset S. bovis group bacteremia generally present with acute onset of respiratory distress and sepsis within the first 5 days of life. In contrast, late-onset S. bovis group infection generally presents with urinary sepsis or meningitis.
Streptococcus gallolyticus subsp. pasteurianus is capable of causing fulminant neonatal sepsis or meningitis that is clinically indistinguishable from that caused by group B streptococcus. Brain abscesses, delayed subdural effusions and brain haemorrhages have been described as complications of this infection and therefore long-term neurological follow-up of these infants are recommended.

**ID: 8300**

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**ABSTRACT TITLE:** PHENOTYPIC AND GENOTYPIC CORRELATION OF CARBAPENEMASE-PRODUCING ENTEROBACTERIACEAE AND RECOMMENDATIONS FOR THE ROUTINE SCREENING OF CARBAPENEM-RESISTANT ENTEROBACTERIACEAE

**Introduction**

The emergence and transmission of CRE is concerning in the clinical and public health arena. Reliable and accurate detection for patient management and infection prevention and control purposes is required. Routinely, phenotypic methods are utilised for identification of CRE.

**Methods**

Phenotypic profiles of 2678 suspected-CPE isolates generated by the automated MicroScan®-Walkaway-system using CLSI guidelines were correlated with carbapenemase production as identified molecularly.

**Results**

Klebsiella pneumoniae accounted for 63% of all isolates (n = 1685) followed by Enterobacter cloacae (n = 361, 14%). Carbapenemases accounted for 75% of isolates (blaOXA-48 (n = 978, 37%), blaNDM (n = 904, 34%), blaVIM (n = 108, 4%), blaIMP (n = 35, 1.3%), blaGES (n = 24, 0.9%), blaKPC (n = 18, 0.7 %)). A substantial number of isolates expressing a carbapenemase/s were susceptible to third- and fourth-generation cephalosporins and carbapenemases demonstrating that confirmed carbapenemase-producing isolates are not presenting as possible carriers of carbapenemases using routine diagnostic methods. Similar results were obtained when CLSI and EUCAST clinical breakpoints were applied. This was improved using EUCAST epidemiological cut-off (ECOFF) values. The recommended ECOFF value of > 0.12 mg/mL for meropenem, reduced the number of missed carbapenemase-producing isolates. Exact quantification could not be performed as dilutions lower than the lowest MIC dilution is required for analysis.

**Conclusion**

While ECOFF values are appropriate for infection prevention and control purposes, clinical breakpoints are suitable for patient management. Considering this, and other published data investigating appropriate methods for carbapenemase screening, routine laboratories should use a combination of the EUCAST meropenem screening cut-off of < 25 mm or MIC > 0.12 mg/ML and ertapenem (or meropenem) APBA and EDTA combination disk tests plus temocillin disk diffusion on MH-CLX agar.

**ID: 8485**

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**ABSTRACT TITLE:** PREDOMINANT METHICILLIN-RESISTANT STAPHYLOCOCCUS AUREUS SPA-TYPES CIRCULATING AT FIVE SENTINEL SITES ACROSS SOUTH AFRICA

**Introduction**

Staphylococcal protein A (spa)-typing is a valuable tool to track the spread of Staphylococcus aureus. Spa-typing exploits S. aureus’ clonal nature and therefore isolates with the same spa-type are related. The aim of this study was to define the predominant methicillin-resistant S. aureus (MRSA) spa-types circulating at five sentinel sites across South Africa.

**Methods**

Staphylococcus aureus bloodstream infection isolates were submitted to the national reference laboratory as part of GERMS surveillance from 2010-2015. Methicillin-resistance was defined as the PCR detection of the mecA gene and non-susceptibility to oxacillin and/or cefoxitin. Molecular testing included spa-typing, Staphylococcal Chromosome Cassette (SCC) mec typing and multilocus sequencing typing (MLST) on selected isolates.

**Results**

A total of 3167 isolates were submitted and 34% (1087/3167) were methicillin-resistant, of which 74% underwent spa-typing. Spa-type t037 predominated in three Gauteng hospitals [Hospital A = 73% (203/278); Hospital B = 47% (52/110) and Hospital C = 50% (56/113)] and in a Western Cape hospital [Hospital E = 27.3% (44/161)]. Spa-type t1257 (27%, 37/139) predominated in another Western Cape hospital. The majority of t037 cases in the Western Cape occurred in...
a burn unit (59%, 26/44) among adult males (85%, 22/26). Most t037 cases in Gauteng hospitals were associated with paediatric patients residing in a specific ward within the respective hospital (Hospital A = 28% (57/203) in a premature ward; Hospital B = 60% (31/52) in a paediatric ICU and Hospital C = 25% (14/56) in a paediatric surgery ward). Most t037 isolates in all hospitals harboured SCCmec type III (93%, 330/355) and three isolates belonged to MLST clonal complex (CC) 8, sequence type (ST) 239.

Conclusion
The predominant spa-types in South Africa are t037 and t1257. Preliminary results show that isolates are possibly related and that further investigation with pulsed-field gel electrophoresis is warranted to assist in infection prevention and control efforts.

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ABSTRACT TITLE: AN INVESTIGATION OF STREPTOCOCCUS PNEUMONIAE COLONISING THE NASOPHARYNX OF CHILDREN ATTENDING THE PAEDIATRIC CLINIC AT DR. GEORGE MUKHARI ACADEMIC HOSPITAL

Introduction
Polysaccharide Conjugate Vaccine (PCV) was introduced in 2009 in the South African childhood immunisation programme. Since then, there has been a dramatic decrease in the incidence of pneumococcal diseases in both vaccinated children and non-vaccinated individuals of all ages. However, increased infections caused by non-vaccine serotypes have been reported. The reduction in vaccine serotypes may however open a niche in the nasopharynx allowing for increases in the acquisition and prevalence of pneumococcal non-vaccine serotype colonization and subsequent disease. Following the South African PCV introduction, the impact of PCV on nasopharyngeal colonization has not been adequately assessed. Therefore, the aim of the study was to investigate the prevalence of S. pneumoniae colonising the nasopharynx of children aged 2 months-14yrs attending the paediatric clinic at Dr. George Mukhari Academic Hospital.

Methods
Collected nasopharyngeal swabs were cultured. Isolates were identified phenotypically and were serotyped using conventional multiplex PCR targeting 29 of the most invasive serotypes and those included in the PCV.

Results
Thirteen-percent (47/350) of the isolates were positive for S. pneumoniae. Forty two percent (42%) of the isolates were serotyped using PCR. Identified serotypes were 6A/B (35%), 1(25%), 7C (25), 8(5%), 31(5%) and 20(5%).

Conclusion
Non-vaccine serotypes detected in this study are an indication of serotype replacement, with the decline of vaccine serotypes giving way for non-vaccine serotypes to prevail. Overall, PCV has shown alteration in colonising serotypes.

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ABSTRACT TITLE: THE USE OF A URINARY ANTIGEN TEST FOR THE RAPID IDENTIFICATION OF S. PNEUMONIAE BACTERAEMIA

Introduction
Novel detection methods for invasive pneumococcal disease (IPD) include PCR and antigen tests. The ‘gold standard’ for diagnosis of IPD is culture from sterile sites. Prior antibiotic therapy, autolysin production and the fastidious nature of pneumococci can yield culture negative results. The aim of this study was to evaluate the use of a novel urine antigen test directly on blood cultures, for the rapid identification of S. pneumoniae.

Methods
The BinaxNOW Streptococcus pneumoniae urinary antigen test was compared to routine culture with optochin, and lytA PCR, on blood cultures with Gram positive cocci in chains or pairs, and where the morphology was indiscriminate.
Results

Preliminary results of 46 blood cultures tested indicated that 5 were non-viable on culture. When compared to culture, the sensitivity (7/7) and specificity (34/34) of the antigen test was 100%. The 5 non-viable samples were however, positive on both PCR and the antigen test.

When comparing the antigen test to PCR, the sensitivity was 75% (12/16), and specificity was 100% (30/30). Of the 16 lytA positive samples, the antigen test reported 4 as negative. These isolates were other Gram-positive cocci on culture, and produced high CT values on PCR. When excluded from analysis, the sensitivity and specificity of the test increased to 100%.

Conclusion

The use of the urine antigen test on blood culture, in conjunction with Gram stain, gave a rapid and accurate presumptive diagnosis of IPD. This would allow for targeted antimicrobial therapy sooner, and promote antibiotic stewardship. It can also be used as a cost-effective alternative to molecular testing in cases of culture negative S. pneumoniae bacteraemia.

ID: 8514

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ABSTRACT TITLE: MOLECULAR CHARACTERISATION AND ANTIMICROBIAL SUSCEPTIBILITY TESTING OF GROUP A STREPTOCOCCUS ISOLATES FROM INVASIVE AND NON-INVASIVE INFECTIONS AT PHEDISONG 4, SOSHANGUVE 3, AND DR GEORGE MUKHARI ACADEMIC HOSPITAL

Introduction

Group A streptococcus (GAS) is responsible for a wide range of invasive and non-invasive infections. Pharyngitis due to GAS may cause complications such as acute rheumatic fever/ rheumatic heart disease which have high mortality and morbidity rates in developing countries. Antimicrobial susceptibility testing and molecular characterisation of GAS isolates will yield information required for treatment using antibiotics and preventative vaccine development

Aim

The study aims to determine the emm genotypes and antimicrobial susceptibility profiles of GAS isolates circulating in the North-West area of Pretoria.

Methods

This is a quantitative cross-sectional study describing the prevalent emm genotypes circulating in the region. Throat swabs from patients presenting with pharyngitis from the clinics were collected for GAS isolation. GAS isolates were also collected from the National Health Laboratory Services (NHLS). Quantitative antimicrobial susceptibility testing was done for Penicillin, Clindamycin, Erythromycin and disk susceptibility testing for Cotrimoxazole. Genotyping of GAS strains will be done using PCR, sequencing and pulsed field gel electrophoresis.

Results

A total of 24 samples were collected, from which, nine were throat swabs and 17 were GAS clinical isolates from the NHLS. From the nine collected swabs, two were identified as GAS. MICs for the Penicillin, Clindamycin, Erythromycin ranged from 0.012-0.08 ug/ml, 0.016-0.125 ug/ml, 0.016-0.064 ug/ml respectively. Zone sizes for cotrimoxazole were between 20-35mm which were also susceptible.

The MICs for the antimicrobials tested were all at the lower ranges of susceptibility breakpoints.

Conclusion

GAS isolates still remain susceptible to penicillin, despite reports of macrolide resistance; all of our isolates were susceptible and were inducible Clindamycin resistance negative. Molecular characterisation of the emm types is ongoing.

ID: 8240

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ABSTRACT TITLE: CULTURE CONFIRMED BACTERIAL INFECTIONS IN HIV-INFECTED AND UNINFECTED SEVERELY MALNOURISHED CHILDREN ADMITTED TO KING EDWARD VIII HOSPITAL, DURBAN

Introduction

Malnutrition results in an alteration in both innate and adaptive host defence mechanisms, resulting in an increased susceptibility to infections. This study seeks to describe and compare culture confirmed bacterial infections in HIV negative severely malnourished children admitted to the same setting.

Methods

Specimens (blood, cerebrospinal fluid, urine and sputum) obtained from 101 children aged between six and sixty months admitted with severe acute malnutrition (SAM) in King Edward Hospital Durban South Africa between 1 January 2015 and 31 December 2015 were retrospectively identified. Positive bacterial cultures obtained within two days of admission and between two to thirty
days of admissions were classified as admission or hospital acquired infections respectively.

Results

101 patients were legible for the study of which 53% were HIV Unexposed.73% of the total 250 cultures obtained were during admission and 44% of all the cultures obtained during admission were from blood. Gram Negative organisms are the predominantly cultured organisms in both admission and hospital acquired cultures. Escherichia Coli (Ecoli) contributes 26% of all the positive cultures on admission.

Conclusion

Gram Negative organisms remain an area of concern in both HIV Positive and HIV Negative patients with Severe Acute Malnutrition with resistant organisms more prevalent in HIV positive patients. Improving infection control practices are of vital importance in these vulnerable children to reduce the morbidity associated with hospital acquired infections.

ID: 8481

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ABSTRACT TITLE: A RETROSPECTIVE REVIEW OF LISTERIA MONOCYTOGENES INFECTION AT TYGERBERG HOSPITAL FROM 2006 TO 2016: IS EMPIRIC AMPICILLIN STILL INDICATED PAST THE EARLY NEONATAL PERIOD?

Introduction

Ampicillin is added empirically to the treatment of suspected sepsis or meningitis in infants < 3 months of age to cover Listeria Monocytogenes (LM). In view of the limited LM cases seen in other countries, the limited South African data available and the recent Ampicillin shortage at Tygerberg Hospital (TBH), we aimed to describe the positive cultures of LM at TBH to rationalize our Ampicillin usage.

Methods

This was a retrospective descriptive study of all patients with a blood or cerebral spinal fluid (CSF) culture yielding LM processed at the TBH laboratory, including Eastern Metro, Winelands and Overberg over the 11-year period 01/01/2006 – 31/12/2016. All positive cultures of children < 13 years of age were included in the analysis.

Results

There was a total of 26 positive cultures for LM and 23/26 (88%) < 3 month of age, all of which were < 1month old. 13/23 (56.5%) infants were managed at TBH. 6/13 (46%) presented on the day of delivery. 12/13 (92%) were admitted to the neonatal intensive care unit (NICU) and 8/13 (62%) died. Babies born and managed at our referral hospitals were more likely to have CSF taken ((90% vs 31% (p = 0.019)), a higher platelet count (239x10^9/L vs 107x10^9/L (p = 0.004)), lower CRP (64 mg/L vs 137 mg/L (p = 0.01)) and a lower mortality rate (0% vs 62% (p = 0.002)) than infants managed at TBH. The calculated incidence of LM at TBH was 0.04/1000 live births, and 2.3/1000 NICU admissions.

Conclusion

In concordance with other countries, incidence of neonatal LM infection at TBH is low. However, infants present with severe disease and a high mortality rate. Given that no cases of neonatal LM presented >10 days of age, it would be safe to limit empiric Ampicillin prescription to infants < 1 month old. This is in keeping with international guidelines from developed countries.

ID: 8495

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ABSTRACT TITLE: EXPRESSION OF ROTAVIRUS VP6 OPEN READING FRAME IN VARIOUS YEASTS AS A MORE AFFORDABLE ROTAVIRUS VACCINE CANDIDATE

Introduction

Rotavirus infection is one of the six leading causes of death among children under five years of age. Developing countries in Sub-Saharan Africa has a high disease burden. Currently, there are two live-attenuated vaccines, Rotarix™ (GlaxoSmithKline) and RotaTeq® (Merck), recommended for global use. The efficacy of these vaccines in developing countries is much lower compared to developed countries. Due to the high cost and global demand of these vaccines, local rotavirus vaccine production is desirable. Antibodies against the immunodominant, VP6, are known to neutralise rotavirus intracellularly. Yeast as recombinant protein production vehicle, is cost-effective and scalable. A unique yeast expression system that allows for the potential recombinant protein expression in any yeast, was previously developed at the University of the Free State (UFS).
**Methods**

The VP6 open reading frame (ORF) of a South African rotavirus strain was optimised for expression in Arxula adeninivorans, Pichia pastoris/angusta and Kluyveromyces lactis. The VP6 ORF was cloned into the yeast expression vector (pKM177_VP6). The UNESCO-MIRCEN yeast culture collection at UFS was targeted for recombinant protein production. Colony PCR was used to verify the integration of the VP6 ORF in each of the yeast tested. The expression of the VP6 was confirmed by western blot analysis.

**Results**

More than 45 yeast colonies representing five different yeast strains (Saccharomyces cerevisiae, P. pastoris, P. angusta, K. lactis and A. adeninivorans) expressing VP6 have been identified. Interestingly, results indicated that integration of the A. adeninivorans optimised VP6 ORF was the most effective.

**Conclusion**

The study provides a platform to select the best VP6-producing yeast strains. Such yeast strains will be used to scale up production for immune response studies in animals. This approach ensure that a high-yield yeast strain can be obtained to reach our long-term objective to develop a more affordable rotavirus vaccine.

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**ID: 8513**

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**ABSTRACT TITLE:** WILL THE REAL PATHOGEN PLEASE STAND UP? A CASE REPORT OF EXUDATIVE PHARYNGITIS CAUSED BY CORYNEBACTERIUM PSEUDODIPHtheriticum

**Introduction**

Corynebacterium pseudodiphtheriticum is a respiratory tract commensal and emerging pathogen that is closely related to C. diphtheriae. We report a case of exudative pharyngitis from the Western Cape, ostensibly caused by C. pseudodiphtheriticum, in the aftermath of the C. diphtheriae outbreak in South Africa in 2015.

**Case**

A 14-month old male presented in shock with fever, features of upper airway obstruction, and an adherent, white pharyngeal membrane. Neck swelling, cranial nerve palsies and muscle weakness were absent. The patient’s vaccination series was incomplete, and he was newly diagnosed as HIV-infected.

The child was resuscitated and commenced on penicillin G, followed by cefotaxime when susceptibility was confirmed, to complete a 14-day course. Diphtheria antitoxin was not obtainable. Clinical response was protracted but steady.

Black colonies were isolated on tellurite-containing media from a throat swab, and were identified using two methods as C. pseudodiphtheriticum. Toxin testing by molecular and phenotypic methods were both negative. Antimicrobial susceptibility testing revealed susceptibility to penicillin, cefotaxime and azithromycin.

Healthcare and family contacts were screened. Two family members cultured C. pseudodiphtheriticum on throat swabs. Catch-up vaccinations and azithromycin prophylaxis were administered where appropriate.

**Discussion**

C. pseudodiphtheriticum has been aetiologically linked to a variety of diseases ranging from necrotising tracheitis to urinary tract infection. Respiratory tract disease is the commonest manifestation and has been reported in the presence of underlying patient risk factors, such as immunocompromise, structural lung disease, and chronic medical conditions, and following invasive respiratory procedures, including endotracheal intubation.

Only three cases of exudative pharyngitis caused by this organism have been reported, in a 32-year old male and a 4- and a 6-year old female.

C. pseudodiphtheriticum is usually susceptible to penicillin. Susceptibility to the macrolides and cefotaxime is variable; laboratory testing is advised. Management of this case required a multidisciplinary approach, including the infection control team.

This organism should be considered in the differential diagnosis of exudative pharyngitis, and further studies should attempt to define its association in this context.

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**ID: 8523**

**Category:** Paediatric Infectious Diseases (SASPID)

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Replacement has occurred.

Isolates, however the assay is less useful in cases where serotype identities not included in the assay with one isolate non-typeable. One isolate gave a discordant result. Two isolates had serotype identities not included in the assay with one isolate non-typeable. Of 104 pneumococcal isolates typed by Quellung, 96 isolates were assigned a serotype; four isolates could not be typed by Quellung. Of the 84 carriage isolates typed by the real-time PCR to serotype pneumococcal carriage isolates obtained from healthy children and infants at two primary health care centres in Gauteng between 2014 and 2016.

Methods

Three hundred and twenty-one nasopharyngeal samples were taken at specified ages and immediately processed. The epidemiological and antimicrobial susceptibility data of the samples was analysed. A 114 identified S. pneumoniae isolates were serotyped using Quellung. Validation of the sequential triplex real-time PCR was done using 77 Quellung typed invasive pneumococcal control isolates. To investigate the utility of the assay, 84 carriage isolates were subjected to molecular serotyping.

Results

Nasopharyngeal carriage of S. pneumoniae 114/309 (37%), H.influenzae 56/309 (18%), S. aureus 61/309 (20%), and M. catarrhalis 5/309 (2%) was observed; with MRSA and C. pseudodiptheriticum also isolated. The most prevalent serotypes amongst pneumococcal isolates were 11A, 23B and 15B/C. Of 77 pneumococcal validation isolates, 74 had a serotype covered by the assay. 73 isolates yielded a positive signal, concordant with the Quellung (99%). One isolate gave a discordant result. Two isolates had serotype identities not included in the assay with one isolate non-typeable. Of 104 pneumococcal isolates typed by Quellung, 96 isolates were assigned a serotype; four isolates could not be typed by Quellung. Of the 84 carriage isolates serotyped by the real-time assay, 48 isolates had serotypes covered by the assay. 43/48 (90%) yielded a positive result. Five of these gave discordant results.

Conclusion

The real-time assay was designed to detect 21 serotypes including those found in PCV-13, which were also detected in carriage isolates, however the assay is less useful in cases where serotype replacement has occurred.