A Case of Lower Limb Myiasis in Hong Kong: Case Report and Brief Review of Existing Literature

Single Stitch Mesh Fixation During Laparoscopic Trans-Abdominal Pre-Peritoneal Groin Hernia Repair: A Retrospective Study of 3800 TAPP Repairs

Medical Student Involvement in and Attitudes towards Audit and Research: The MEDical Student Experience of Audit and Research (MED-SEARCH) Survey

Body Mass Index and Pregnancy Outcomes in Expectant Women at Moi Teaching and Referral Hospital, Eldoret Kenya

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The High Ratio of Undiagnosed Cases of Low Back Pain: Implications for Its Management

Abstracts from the 8th International Academic and Research Conference 2018
SURVIVAL OF OSTEOSARCOMA PATIENTS FOLLOWING DIAGNOSIS OF SYNCHRONOUS SKIP METASTASES

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Background and Aims: Skip metastases occur in 2% of high grade osteosarcomas. They are defined as smaller foci of tumour that occur completely separately from the primary lesion within the same bone. We aim to investigate the prognostic implications of synchronous skip metastases in osteosarcoma by presenting the largest single centre series to date.

Methods: A retrospective study was conducted on all 21 osteosarcoma patients with skip metastases treated at our unit between 1983 and 2004. Patients' demographic data was collected, along with information about margin status, percentage of necrosis, response to chemotherapy, and presence of lung metastasis at initial presentation.

Results: The incidence of skip metastases in patients presenting with osteosarcoma is 1.3%. The median age at diagnosis was 14 years. Overall survival was 81% at 12 months, 60% at 24 months, and 31% at 36 months. Nine patients (43%) had lung metastases at diagnosis, and the remaining 12 patients had no other sites of metastasis. Seventeen patients (81%) underwent limb salvage, and four patients (19%) were treated with amputation. No statistical difference in overall survival was demonstrated when comparing presence of lung metastases to those without (p=0.859). No statistical difference was found in overall survival according to age group (<18yrs vs. >18yrs; p=0.126). We found no statistical difference in overall survival according to percentage of chemotherapy-induced bone necrosis (<95% vs. >95% p=0.056).

Conclusion: The presence of skip metastases confers a very poor prognosis as an independent variable, and also confers a worse prognosis than synchronous lung metastasis. Osteosarcoma patients with skip lesions who require surgical amputation carry a poor prognosis post-surgery due to large tumour size and neurovascular invasion.
Background: The DRSP was established to identify early retinopathy and treatment. The screen involves Digital Photographic Assessment (DPA) where each eye is graded (referable or non-referable). Some patients fail DPA due to pathology, such as cataract and corneal scarring, and must undergo Slit Lamp Biomicroscopy (SLB). DR cannot be treated while this pathology persists.

Aims: To assess whether information gained from one eye can be used as an indicator of disease of the other eye which cannot be visualised and graded by slit lamp. Hence, use the gradable eye as reference for making clinical management decisions in the upgradable eye.

Methods: Data was obtained retrospectively from the Leeds DRSP database for patients who failed photographic assessment and had been sent for slit lamp examination between 01/01/2015 and 31/12/2015. During this period, 881 patients failed photographic assessment. One hundred and fifty-three patients with missing data were excluded, leaving a sample of 728. Conditional probability was used to analyse the data.

Results: The majority of patients had the same grade of retinopathy (92%) and maculopathy (99%) bilaterally. In patients with differences in retinopathy grades between eyes, 91% had a difference of one grade only. Conditional probability analysis showed that in 99% of cases, if non-referable retinopathy is found in one eye, it is also present in the other. Analysis also showed a greater propensity for retinopathy in the left eye.

Discussion: This study provides evidence of using the gradable eye as an indicator of disease in the upgradable eye. Results empower clinicians to be more confident about the likelihood of pathology (or no pathology) in the upgradable eye based on the reference eye before pathology has progressed to advanced stages (e.g., vitreous haemorrhage and retinal detachment). There is also evidence to suggest lateralisation of disease to the left, which requires further research.
Abstracts from the 8th International Academic and Research Conference
5th August 2017, University Place, University of Manchester

Winners in Individual Categories
Oral presentations

First Prize
Category: Clinical and Basic Science Research

RAN GTPASE PROMOTES MALIGNANT MELANOMA CELL PROLIFERATION, MIGRATION AND INVASION BY DEREGLATING MET-MEDIATED DOWNSTREAM SIGNALLING

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Background and Aims: Ras-related nuclear protein (Ran) is a GTPase that is essential for important cellular functions, such as nucleocytoplasmic trafficking and mitosis. Recently, it was discovered that Ran is over-expressed in several tumour types compared with normal counterparts, suggesting a role in driving cancer development. However, Ran expression and its significance in cancer cell proliferation, migration and invasion have not been investigated in malignant melanoma. This study aims to examine if Ran GTPase promotes malignant melanoma cell proliferation, migration and invasion by deregulating met-mediated downstream signalling.

Materials and Methods: A375 and G361 malignant melanoma cell lines derived from patients were cultured in T75 flasks. Ran expression was investigated with real-time PCR and SDS-PAGE. Ran expression knockdown and cell transfection was achieved with siRNA technology. Cancer cell proliferation for transfected and non-transfected melanoma cells was assessed with a resazurin-based assay. Cell migration and invasion were assessed with a transwell assay without or with the use of matrix matrigel respectively and with a wound healing assay. Immunofluorescence was used to detect the c-Met receptor. Statistical analysis was carried out using GraphPad Prism.

Results: Ran expression was detected in both cell line models. Ran knockdown with siRNAs resulted in as much as 94% efficiency. Ran knockdown significantly reduced melanoma cell proliferation, migration and invasion, and increased wound closure time. Furthermore, reduction in Ran levels resulted in c-Met down-regulation.

Discussion: Ran is vital for cancer cell survival so targeting Ran could reduce melanoma progression. However, more data is needed from in vivo studies. The findings relate to studies with Ran knockdown in other cancers.

Conclusion: Our findings show for the first time that Ran is a potential target for novel therapeutic strategies for malignant melanoma.
EXPLORING THE RELATIONSHIP BETWEEN TONGUE MUSCLE STRENGTH AND CLINICAL SIGNS AND SYMPTOMS IN AMYOTROPHIC LATERAL SCLEROSIS (ALS)

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Background and Aims: Bulbar dysfunction is a presenting feature in approximately 20% of patients with amyotrophic lateral sclerosis (ALS). However, there are currently no validated biomarkers for the assessment of bulbar dysfunction in the context of clinical trials. Since the tongue has shown to be affected most by weakness compared to other bulbar muscles, tongue strength (TS) has been identified as a potential biomarker for bulbar disease. This study aims to assess the extent to which TS can detect bulbar dysfunction in ALS patients.

Methods: TS measurements (obtained using a quantitative muscle testing system) were recorded in 27 patients with bulbar ALS and 26 healthy age- and sex-matched volunteers. Patients’ and volunteers’ TS measurements were compared using Student’s t-test. The relationship between patients’ TS measurements and ALS functional rating scale (ALSFRS) scores (total and bulbar sub-scores) was assessed using Pearson's correlation. Multiple correlations were made between volunteers, patients without, and patients with tongue wasting (a clinical sign of bulbar dysfunction). These were assessed using the one-way ANOVA test.

Results: TS measurements differed significantly between patients and volunteers (p<0.0001). A significant difference was also shown between TS measurements of patients with and patients without tongue wasting (p<0.0001), and between volunteers and patients without tongue wasting (p<0.0001). TS did not correlate with total ALSFRS scores (r=0.1807, p=0.2044), but correlated significantly with ALSFRS bulbar sub-scores, with r=0.7102 (p<0.0001).

Conclusion: The difference between TS measurements of volunteers and patients without tongue weakness or wasting suggests that TS may have the potential to detect preclinical signs. This could facilitate a more accurate prediction of disease progression and greatly impact future clinical decision-making. Its ability to detect changes overtime would need to be examined before it can be considered a valid tool for the assessment of bulbar dysfunction in ALS.
THE PROGNOSTIC VALUE OF PSOAS MUSCLE THICKNESS MEASURED ON COMPUTERISED TOMOGRAPHY SCANS IN ALCOHOLIC LIVER DISEASE
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Background: The United Kingdom Model for End-Stage Liver Disease (UKELD) score is used to predict mortality in patients with alcoholic liver disease, and prioritise the waiting list for liver transplantation. However, the UKELD score has been shown to be unreliable in patients with low UKELD score and refractory ascites. Sarcopenia is an independent predictor of mortality in ALD, and psoas muscle thickness may be a measure of sarcopenia.

Aims: To determine whether psoas muscle thickness measured on CT scan can be used to predict mortality in patients with ALD.

Methods: A retrospective study of abdominal CT images of 54 patients with ALD admitted to a tertiary hepatology unit in a one-year period. We identified the status of each patient four years post-admission: alive (n = 26); dead (n=28). Axial and transverse psoas muscle thickness was measured on CT scans at the level of the umbilicus. Mann Whitney U Test (α <0.05) was used to test for significant difference in psoas thickness in alive vs. dead patients. Spearman’s rank-order correlation (α <0.05) was used to test for correlation between psoas thickness and UKELD score.

Results: No significant difference in psoas thickness in alive vs. dead patients: transverse thickness (p = 0.29), and axial thickness (p = 0.49). No significant correlation between psoas thickness and UKELD score: transverse thickness (r = 0.05, p= 0.74), and axial thickness (r = -0.27, p= 0.07).

Discussion: Our findings imply that psoas thickness may not be a reliable measure of sarcopenia. However, this study was limited by a small sample size and a lack of complete data set due to the retrospective study design.

Conclusion: Our data set suggests that psoas muscle thickness on CT scan is not a reliable prognostic tool to predict mortality in patients with ALD. A large prospective study is needed.
STAKEHOLDERS’ VIEWS OF SURGEON-SPECIFIC MORTALITY DATA

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Background and Aims: Public release of Surgeon-Specific Mortality Data (SSMD) was mandated by the National Health Service in 2013, with the aim of enhancing transparency, driving quality improvement and facilitating patient choice. Literature evaluating the impact of SSMD publication is conflicting and there is a current paucity of research elucidating its importance to patients. As such, this study’s aims were to evaluate the views of vascular patients regarding current public outcome reporting at both the surgeon- and hospital-level and to investigate patients’ priorities for future outcome reporting.

Methods: A novel questionnaire was designed and trialled with patients to ensure content and construct validity. This was then distributed to patients in vascular outpatient clinics across three sites at Imperial College Healthcare Trust over a six-week period. Patients with confirmed or suspected arterial disease were recruited. Statistical analyses were conducted to evaluate relationships between responses and demographic variables and to ascertain patients’ preference for surgeon- or hospital-level reporting.

Results: Overall, 165 patients participated in the survey (response rate 63%, n=261). Most were unaware of SSMD (80%), with only 10% accessing it pre-operatively. Factors influencing patient awareness of SSMD included education level (p=0.047) and salary (p=0.049). Patients recognised SSMD’s utility in choice of treatment (60%) or surgeon (53%). However, most prioritised the patient-surgeon relationship (90%) and past surgical experiences (71%) when making pre-operative decisions. In line with views expressed by healthcare professionals in the literature, respondents favoured hospital mortality data (66%) over surgeon-level reporting (49%) (p=0.000006), with 63% expressing that patients were likely to misinterpret SSMD.

Conclusion: The aims of public-outcome reporting in informing patient choice are not being met by current methods. Patients express preference for data at the hospital-level and prioritise previous experiences and metrics other than mortality in their pre-operative decision-making. Therefore, policy makers should expand SSMD to include hospital-specific mortality data and consider publishing accessible and contextualised patient-directed outcomes at the hospital-level.
MAINTENANCE IV FLUID DOCUMENTATION AND PRESCRIBING PRACTICE IN ADULT SURGICAL PATIENTS

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Background: IV fluids for maintenance are commonly prescribed in the inpatient setting. Inappropriate prescribing can lead to significant patient morbidity and mortality.

Aims: The study aimed to evaluate the appropriate use of IV fluids in surgical patients.

Methods: We investigated the current compliance of IV fluid prescription and documentation with national standards which included details of patient assessment; documentation of rate, volume and type of IV fluid to be administered; and IV fluid management plan and electrolyte needs over next 24 hours. Prescription of 1 mmol/kg/day of potassium, sodium and chloride, 25–30 ml/kg/day of water, and 50–100 g/day of glucose. These standards were set with an expected compliance of 100%. Data were collected from three surgical wards over a 14-day period. All adult surgical admissions were reviewed for inclusion. Only those patients requiring maintenance fluids were included in analysis. A total of 20 patients were included in the audit. Data were recorded using a self-designed audit collection tool.

Results: Documentation: Details of fluid assessment were documented in 5% of patients. Type, rate and volume of fluid to be administered were documented correctly in 100%. An IV fluid management plan was documented in 20%. Fluid and electrolyte needs over next 24 hours were documented in 10%.

Prescription: Mean potassium received was 18.397% of target, mean sodium received was 511.68% of target, and mean chloride received was 471.034% of target. Mean water received was 136.397% of target. 25% patients received target glucose.

Discussion: This study demonstrated that 100% compliance of standards set was not achieved. IV fluid prescribing and documentation remain poor. An educational campaign is required to increase awareness of NICE IV fluid guidelines and the risks associated with poor prescribing of IV fluids. We aim to make alterations to the Trust drug chart to aid improved documentation and prescribing practice.
AUDIT ON THE DIAGNOSIS, INVESTIGATION AND TREATMENT OF PANCREATIC NEUROENDOCRINE TUMOURS IN HULL HOSPITALS

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**Background:** Incidence of pancreatic neuroendocrine tumours (PNETs) is increasing in the UK. They may be functioning (with symptoms due to hormone excess) or non-functioning. Imaging of PNETs involves CT, MRI and Octreotide scans. Treatments are based on tumour grade and histological diagnosis. The mainstay of PNET treatment is surgery. The European Neuroendocrine Tumour Society (ENETS) guidelines provide a consensus on the diagnosis and treatment of PNETs.

**Aims:** To retrospectively audit patients with a histological diagnosis of PNETs in Hull hospitals between February and October 2015. To review patient characteristics (age and gender). To assess if investigations and management are in line with ENETS guidelines.

**Methods:** A list of patients was obtained from a Hull MDT pathologist. A dedicated neuroendocrine database has been created to collate information on each patient, with data obtained via the clinical system Lorenzo.

**Results:** Preliminary data on 14 PNET patients. There were equal proportions of men and women; the mean age at diagnosis was 66 (range 33-82). The majority of patients (57%) were diagnosed with a Grade 1 PNET (Ki-67 index of <2%). Histological results following curative surgery was the most common form of diagnosis. The average tumour diameter on histological resection was 1.8 cm. Most patients (71%) were diagnosed with localised disease. Octreotide is recommended by ENETS, but was only performed in 14% of patients. 40% of PNETs were found on incidental investigation, and 21% of patients were diagnosed with a functional PNET; the majority of these were glucagonomas. 79% of patients underwent surgery as initial management with curative intent.

**Conclusion:** The preliminary results from this local audit support evidence from other sources that most PNETs are non-functioning and a significant proportion are diagnosed incidentally. More data needs to be collected to sufficiently analyse the performance of this local neuroendocrine centre compared to European guidelines.
AN ATYPICAL PRESENTATION OF INFECTIVE ENDOCARDITIS IN LYMPHANGIECTASIA: AN EDUCATIONAL AND ETHICAL CASE

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Background: Lymphangiectasia is a rare pathological dilatation of the intestinal lymphatics, causing a protein-losing gastroenteropathy.

Case Study: A 74-year-old female, with known small bowel lymphangiectasia requiring total parenteral nutrition through a central venous catheter, presented to her routine nutrition clinic with a three-week history of fever, shivering, weakness, and generalised muscular pain. Blood cultures identified staphylococcus aureus infection, and the patient was advised to attend her local Accident and Emergency Department. On assessment, the patient had a previously undocumented pansystolic murmur. A transthoracic echocardiogram detected a mitral valve rupture, suggesting infective endocarditis. Infective endocarditis was confirmed by a transoesophageal echocardiogram which found 1.7cm vegetation on the anterior mitral valve, causing a prolapse of the anterior leaflet. The patient commenced a course of vancomycin and gentamicin as per Trust microbial guidance, and the indwelling central venous catheter was removed. A new central venous catheter was recommended five to seven days following commencement of antibiotic therapy to meet the patient's nutritional requirements. She had become anxious about "bugs growing all over a new line" and wished to trial an oral diet. This was discussed in a Multi-Disciplinary Team meeting and it was agreed that the patient could attempt an oral diet with monitoring of her metabolic blood panel. The patient's albumin level continuously dropped, requiring human albumin solution replacement which contributed to a pulmonary oedema and severe respiratory failure. This lead to admission to the Intensive Therapy Unit. The patient required ventilatory support and a central venous catheter was inserted to optimise treatment.

Discussion: This raises the ethical issues of autonomy and beneficence in the face of a quickly changing clinical scenario and patient capacity.

Conclusion: The patient was treated for pulmonary oedema, stepped down to the coronary care unit, and transferred to a tertiary centre for mitral valve replacement.
CERVICAL SPINE OSTEOMYELITIS IN AN INTRAVENOUS DRUG USER: AN INDOLENT CONDITION WITH HIGH MORBIDITY

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Background: Cervical spine osteomyelitis (CSO) is a rare condition with an increasing incidence due to the aging population and an increase in immunocompromised states such as in Intravenous Drug Users (IVDU). Diagnosis can be challenging as patients often present subclinically with normal blood tests and radiographs initially. If this condition is missed, it can result in serious complications, such as permanent neurological deficits.

Case Study: We present a case of a 36-year-old male IVDU who presented to the emergency department with neck pain. Examination findings, radiograph and blood test were unremarkable. The patient returned two months later with new symptoms of bilateral hand paraesthesiae, malaise and restricted neck movements. A CT scan revealed bone destruction in the vertebral bodies C4 to C6 due to osteomyelitis. He underwent debridement with spinal fixation and received a six-week course of antibiotics. He made a full recovery with no lasting complications.

Discussion: In patients with CSO, blood tests may be normal with negative blood cultures, especially in the immunocompromised. The imaging of choice in suspected CSO is an MRI which has high sensitivity and specificity for this condition. The mainstay of treatment is a minimum six-week course of antibiotics. Surgery may be indicated in patients with neurological deficits and various approaches are available.

Conclusion: CSO may not be included in the differential diagnosis as it is not a commonly encountered condition and neck pain is usually secondary to a benign condition. However, with the rising incidence, it should be suspected in patients who present with neck pain, especially in IVDU. An early diagnosis will reduce morbidity and mortality.
HERNIATION OF AN INTRAGASTRIC BALLOON

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Background: Intragastric balloons are a surgical method for treating obesity. Obesity is a leading global health problem today. It is becoming more and more common to perform non-invasive and invasive treatments to help combat this growing problem, with over 16,000 cases being operated on in the UK alone between 2011-2013. Studies have shown a greater mortality rate in patients with a higher BMI. Therefore, minimally invasive, non-surgical options for weight loss are gaining popularity as a mechanism to help achieve this preoperative weight loss, one of these being intragastric balloon.

Case Study: We present a case of a 61-year-old female who presented with vomiting and was found to have an abdominal wall hernia through which stomach containing the balloon had migrated. She had coughed on the day and then had noticed continual vomiting, was unable to keep anything down, and had noted that her hernia had become firm. Bowels had not opened since then, but she continued to pass wind and urine. An urgent CT scan of the abdomen and pelvis revealed a large anterior abdominal wall hernia with a defect of approximately 10 x 7.5cm. The pylorus of the stomach containing the 12cm gastric balloon, alongside a loop of small bowel and colon, were found to be within the hernia sac. There was no evidence of obstruction. OGD was performed the evening of admission and the balloon was deflated and removed. The patient tolerated a rapid upgrade of diet the next day and her nasogastric tube was removed. Post-operative recovery was uneventful, and she was discharged later that afternoon.

Discussion and Conclusion: Intragastric balloons are safe invasive procedures used to temporarily induce weight loss prior to further definitive surgery. We present rare complications that are not widely recognised, leading to the thought that further assessment needs to be undertaken when selecting patients suitable for the intervention.
EQUIPPING MEDICAL STUDENTS WITH THE COMMUNICATION SKILLS TO DISCUSS FEMALE GENITAL MUTILATION IN THE CLINICAL SETTING

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Background and Aims: Female Genital Mutilation (FGM) is a practice that has affected an estimated 137,000 women in England and Wales. 20,000 girls in Greater Manchester are at risk of FGM. Despite these figures, skills needed by doctors to handle these cases are not taught. In 2014, UK medical students backed a BMA vote for FGM to be a compulsory component of the undergraduate curriculum. Yet, little has changed. This has led to student initiatives on FGM education, such as a workshop developed by the University of Manchester Obstetrics and Gynaecology Society. This study aimed to investigate the role of this workshop in equipping medical students with the necessary communication skills to discuss FGM in the clinical setting.

Methods: The two-hour workshop involved two lectures and role-play clinical scenarios. A pre- and post-intervention questionnaire was completed for participants self-reported competency levels. This covered knowledge of FGM and the UK law surrounding it, how to discuss FGM with patients, and to whom patients should be referred.

Results: Student participant profile: eight medical students, four midwifery students, eight nursing students, one psychology student, and one undisclosed. All 22 forms reported that a comparable workshop should be mandatory for healthcare students. All 16 medical and nursing students suggested that the workshop had improved their knowledge of FGM. Of medical student participants, 75% reported that the workshop increased their confidence to a level where they could communicate with a patient about FGM, and 50% revealed that they would now know what to do clinically if a patient presented with a type of FGM.

Discussion: FGM communication competency was significantly increased in medical students. We propose that, based on the feedback from this event, it is in the best interests of both the patients and the students to receive formal teaching on the communication and management of FGM. The intervention is inexpensive to recreate and beneficial particularly for medical students in areas with high FGM statistics. We are therefore keen to encourage this based on our initial model.
Winners in Individual Categories
Poster presentations

First Prize

POST-OPERATIVE TRANSFUSION PRACTICES IN HIP FRACTURE PATIENTS

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Background and Aims: Every day, 4000 UK hospital beds are occupied by patients with hip fractures. Severe anaemia is a common post-operative complication, and many patients require blood transfusion. We completed an initial audit which looked at blood transfusion practices in hip fracture patients within our Trust. We found that the Trust was mostly non-compliant with NICE guidelines. We introduced a strategy of “Transfuse and Check”, alongside additional quality improvement actions. Following implementation, we re-audited practice. The overall aim was to implement the NICE guidelines on transfusion (NICE CG 24) published in November 2015, and the NICE Quality Standards for Trusts published in April 2017.

Methods: A retrospective cohort study of fractured neck of femur patients over a 12-month period from Warrington Hospital (UK Hip Fracture Database). Post-operative red blood cell transfusions were identified using electronic patient records, Sunquest ICE and hospital transfusion records. Pre- and post-haemoglobin Hb checks, units transfused, and Hb checks after each unit were recorded.

Results: Two hundred and ninety-two fractured neck of femur patients over a 12-month period. Forty-six (15.7%) of these patients were transfused. Twenty-seven (56%) patients were transfused a single unit only. A total of 60.4% of the patients received an Hb check after one unit transfused. On re-audit, 61% fewer RBC units were transfused and there was a 30.4% improvement in post-transfusion Hb targets. A 40.4% increase in single unit transfusion was revealed. An average saving of £112 per patient requiring transfusion was noted.

Discussion: Since the introduction of the “Transfuse and Check” protocol to encourage restrictive transfusion practice, there has been a significant reduction in the number of post-operative red blood cell transfusions in hip fracture patients. This has resulted in reduced risk, reduced cost and a reduced need for red blood cell units not only in our Orthopaedic Department, but also throughout the Trust.

Conclusion: Our “Transfuse and Check” protocol has reduced liberal transfusion of post-operative fractured neck of femur patients.
MEMORY INTERFERENCE IN MICE

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Background and Aims: The question of how memories are forgotten is equally as important to consider as how memories are formed. It is suggested that interference of new memories during the process of learning can increase forgetting. This is relevant to conditions such as Alzheimer’s disease where there is increased susceptibility to interference. If we can minimise interference, it may be possible to improve memory retention by allowing undisturbed consolidation of information. This study aims to investigate this postulation.

Methods: To explore this question, this study uses an object identification recognition test to compare the memory retention between control mice and J20 mice models of Alzheimer’s disease. This test is a good model for human episodic declarative memory, which deteriorates in Alzheimer’s disease. The mice were either exposed to visual interference immediately after studying the objects or underwent a period of sensory deprivation in a black box. They were then tested on their object memory.

Results: Our results show that, although J20 mice had lower memory retention than control mice (F(1) = 3.502, p = 0.068), neither group benefitted from minimising interference after learning (F < 1, p = 0.928). There was no difference in exploratory behaviour between all groups.

Discussion: These results suggest that minimising visual interference alone is not sufficient in reducing the rate of forgetting. The main limitation of this study was that the age of the mice was not kept constant, which meant that the J20 mice had different stages of Alzheimer’s disease. Nevertheless, this theory should not be dismissed as it has been successful in previous literature. We propose that further research should focus on minimising different types of sensory interference such as auditory, smell and tactile.
TO WHAT EXTENT ARE PRENATAL ANDROGENS INVOLVED IN THE DEVELOPMENT OF MALE HOMOSEXUALITY IN HUMANS?

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**Background and Aims:** Male sexual orientation has recently received significant publicity due to the worldwide advancement of LGBT+ rights. However, scientific research has not determined its biological basis. Reduced prenatal exposure to androgens is a potential causal factor in male homosexuality, with animal models demonstrating same-sex mate preference when exposure is altered during prenatal or early postnatal development. This review aims to investigate the extent to which prenatal androgens are involved in the development of male homosexuality in humans.

**Review Description:** As similar studies in humans are impossible, indirect physical and cognitive measures of androgen exposure must be used to examine prenatal exposure to androgens. Some, but not all, studies suggest that physical measures affected by prenatal androgens, including the index-to-ring finger ratio, growth indices, and facial structure, are more 'feminine' in gay men. Gay men also exhibit significant childhood gender non-conformity and a 'feminised' anatomical and functional brain pattern in sexual arousal, as well as domains such as language, visuospatial skills and hemispheric relationships, the changes in which are similar to the changes observed with altered prenatal androgen exposure.

**Discussion:** Although these features are useful indicators of prenatal androgen exposure, they are also confounded by other factors throughout the person's life and fail to provide information on when sexual orientation is determined during gestation. Factors such as genetic influences or maternal immunisation may also underlie the observed differences, and require further investigation. Research has also been hampered by inconsistencies in the reporting of sexual orientation and the potentially unrepresentative populations of gay men studied, while additional complexities pertaining to gender conformity and sexual role may also influence results.

**Conclusion:** While existing findings suggest that prenatal androgen exposure is involved in the development of male homosexuality in humans, future research should examine the effects of androgens on the brain throughout development and how other factors interact with androgens or act independently to influence male sexual orientation.
CIRCADIAN REGULATION OF REJECTION AFTER KIDNEY TRANSPLANTATION

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Background and Aims: Circadian rhythms comprise of oscillations of the levels of important regulatory molecules such as hormones. Consequently, these rhythms influence a number of physiological processes, including the immune system. Disruption of circadian rhythms is associated with pathology. Kidney transplants are of the few types of operations that can occur at any time over 24 hours. This retrospective observational study aims to explore the impact of circadian rhythms in generating rejection.

Methods: Data from 974 transplants that occurred in one hospital unit between 2004 and 2014 was collected, and time of organ reperfusion and presence of rejection was recorded.

Results: From these records, 89 were those that had inconclusive biopsy results and, for these, hospital records were investigated to find the diagnosis. 43 of these patients were those of whom notes were not found and these patients were assumed to not have rejection. There was a pattern of increased rejection for transplant performed between 16:00 and 20:00. Statistical analysis of this pattern using JTK_CYCLE and CircWave was used to determine involvement of circadian rhythms in rejection and found that there was weak circadian contribution.

Discussion: Whilst a weak association between kidney transplant rejection has been identified, this association is not significant enough to provide support for the higher number of rejected transplants carried out between 16:00 and 20:00. A number of other confounding factors also need to be considered, such as cold ischemia time, surgical complications and adherence to immunosuppressive therapy. Missing medical records also limit the significance of these results.

Conclusion: There is a weak association between kidney transplant carried out in the evening and rejection. This association is not significant enough to suggest that individuals who have transplants in the evening are more likely to have rejection.

PROGNOSIS AND TREATMENT OUTCOMES OF SMALL CELL LUNG CANCER PATIENTS AT PRINCESS NORAH ONCOLOGY CENTRE (PNOC), JEDDAH, SAUDI ARABIA

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Background and Aims: Lung cancer is the second most common cancer and the most common cause of cancer-related death. Small cell lung cancer (SCLC) constitutes 13%-15% and is clinically divided into limited-stage disease (LD-SCLC) and extensive-stage disease (ED-SCLC). The median overall survival rate is poor in both subtypes. This study aims to explore the prognosis and treatment outcomes of SCLC in patients at PNOC, Jeddah.

Methods: In this retrospective cohort study, data were collected from patients' medical records for 50 SCLC patients, aged over 18, diagnosed at or referred to PNOC between January 2000 and August 2016. These patients were identified from 360 lung cancer cases. Overall survival rates were investigated using Kaplan-Meier method. Prognostic factors were analysed using Cox proportional hazards model.

Results: The mean age of diagnosis is 61.66±11.587. 45 patients were males and five were females. 43 patients had ED-SCLC and seven had LD-SCLC. The most common presenting symptoms were cough, dyspnoea and weight loss, respectively. The response rate was three of six patients treated with concurrent chemo radiation, eight of 34 patients treated with first line chemotherapy, and one of 13 patients treated with second line chemotherapy. The mean overall survival rate is 31.945 months (95%CI: 21.455 - 42.436) among LD-SCLC patients while the median overall survival rate is 6.209 months (95%CI: 4.697 - 11.721) among ED-SCLC patients.

Discussion: It has previously been reported that the median survival among LD-SCLC patients is approximately 23 months while the median survival is seven to 12 months among ED-SCLC patients. In this study, the mean overall survival for LD-SCLC patients is approximately 32 months and the median overall survival for ED-SCLC patients is six months. Two studies mentioned that the significant prognostic factors were performance status (p < 0.001) and female gender (p<0.0001). This pilot study revealed that sex and performance status were insignificant prognostic factors.

Conclusion: Further local studies are recommended to better understand the epidemiology, prognosis and treatment outcomes of SCLC.
NUCLEATION OF MICROTUBULE BUNDLES BY TAU DROPLETS IN VITRO

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Background and Aims: Microtubule-associated protein tau is associated with a number of neurodegenerative diseases, including Alzheimer’s disease and familial front temporal dementia, for which there is currently no cure. Familial front temporal dementia is a rare form of dementia caused by autosomal dominant mutations in the MAPT gene, such as the P301S mutation. This work aimed to investigate the biological differences between wild-type and P301S mutant tau to further our understanding of how the mutant form of tau induces the early age of onset of neurodegeneration.

Methods/Results: Wild-type tau has been shown to undergo liquid-liquid phase separation to form tau ‘droplets’, which could sequester tubulin and promote the nucleation of microtubules in vitro. This project showed that P301S mutant tau could undergo the same phenomena. P301S tau formed more liquid droplets under the same tau concentrations and microtubules polymerised out of these droplets at a faster rate compared to wild-type tau. Overall, the results were used to design an in vitro assay to dissect the differences between wild-type and the P301S mutant tau in the nucleation and stabilisation of microtubule bundles. Co focal microscopy was used to visualise liquid-liquid phase separation of both wild-type and P301S mutant tau. When low levels of tubulin were mixed with the solution of tau, live imaging using a confocal microscope captured the sequestration of tubulin into the tau droplets, which promoted the nucleation and polymerisation of microtubules.

Discussion: This study provides the opportunity for future examination of the effects of other tau mutants and post-translational modifications (such as hyperphosphorylation) of tau on microtubule dynamics to further our understanding of tauopathies. This will hopefully lead to the development of novel therapies for neurodegenerative diseases involving tau.

GENE REGULATION ASSOCIATED WITH THE ONSET OF POSTOPERATIVE DELIRIUM

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Background and Aims: Postoperative delirium (PD) is a common and serious condition that affects up to 50% of elderly patients undergoing surgery. Global financial costs of PD have been estimated to be in excess of $340bn. The underlying pathological mechanisms that lead to the development of PD have not been elucidated and their link with dementia is poorly understood. Altered gene expression and epigenetic processes are important in common human diseases. The extent to which these play a role in the onset of delirium is not known. The aim of this study is to investigate gene regulation following major surgery and their link with short-term and long-term outcomes.

Methods: Our study is on a cohort of 30 patients over 65 years old enrolled for colorectal and orthopaedic surgery at the Royal Devon and Exeter NHS Foundation Trust in 2015. Blood samples were collected before surgery, one day postoperatively and seven days postoperatively, and past medical history was collected. DNA from samples was used to quantify DNA methylation. Levels of gene expression in key genes highlighted from previous work on methylation patterns will be quantified using qPCR at the RILD site. Follow-up data from these patients will be collected in 2018, three years postoperatively, to observe longer-term outcomes.

Results: Five patients exhibited hypoactive delirium postoperatively. Multiple differentially methylated regions (DMRs) associated with both surgery and the onset of PD was identified in a previous study. We are currently quantifying the expression of gene transcripts annotated to 10 of those DMRs to confirm they are associated with alterations to gene expression.

Conclusion: We are undertaking a study to explore the hypothesis that epigenetic regulation of gene expression is involved in the pathogenesis of PD. We will be quantifying gene expression using qPCR laboratory methods. Our research will provide insight into the mechanisms linking major surgery to the clinical presentation of PD.
TARGETING P63 UPREGULATION MAY PREVENT DEVELOPMENT OF MAPK-INHIBITOR RESISTANCE IN MELANOMA

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Background: Melanoma progression is often characterized by mutations in the mitogen-activated protein kinase (MAPK) and phosphoinositol-3-kinase (PI3K) pathways. Understanding regulation of these pathways has led to development of novel targeted therapies which show high response rates. However, many patients relapse with ensuing resistant disease. P63, a p53 homologue, carries a poorer prognosis when over expressed. In keratinocytes, degradation could be regulated by two ubiquitin ligases, MDM2 and FBXW7. This project explored expression of p63, MDM2 and FBXW7 in MAPK-inhibitor (MAPKi) sensitive and resistant melanoma cell lines.

Method: Morphology change associated with development of resistance was assessed by actin immunostaining using epifluorescence microscopy (Figure 1). Western blot and immunofluorescence experiments enabled analysis of p63, FBXW7 and MDM2 protein expression. Finally, a novel technique of flow cytometry enabled analysis of cell death in MAPKi-resistant melanoma cells treated with Nutlin-3A.

Results and Discussion: MAPKi resistance in melanoma cells is associated with increased p63 expression (Figure 2). Resistance is furthermore associated with reduced FBXW7 expression (Figure 3) and enrichment of nuclear MDM2 (Figure 4), suggesting a potential nuclear interaction between MDM2 and FBXW7. The resultant inactivation of FBXW7 explains the increased p63 expression upon MAPKi resistance. Furthermore, by treating MAPKi-resistant cells with the MDM2-inhibitor Nutlin-3A, we were able to restore FBXW7 expression and promote p63-degradation, with resulting increased melanoma cell death.

Conclusion: This project identifies FBXW7 and MDM2 as regulators of p63 expression in MAPKi-resistant melanoma and proposes a possible role for Nutlin-3A in treating advanced MAPKi-resistant melanoma.

WHAT NEURODEVELOPMENTAL OUTCOMES ARE BEING MEASURED IN CONGENITAL INFECTIONS INCLUDING ZIKA - A SYSTEMATIC REVIEW OF THE LITERATURE

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Background and Aims: There is an emerging interest in the outcomes of children affected by congenital infections since the outbreak of Zika virus. The virus has been linked to congenital neurodevelopmental disorders in children in low and middle-income country settings, where few tools are adapted and standardised to assess children appropriately. This study aims to investigate the neurodevelopmental outcomes measured in congenital infections, including Zika.

Methods: A systematic review of the literature to identify what outcome measures have been used in children affected by congenital infections (including Zika). We searched PubMed, Scopus, Google Scholar and Cochrane from 1960 to 2017 to identify studies where outcomes were measured secondary to in-utero viral infections at birth. We collated studies to evaluate the coverage of outcomes against the framework of the International Classification of Functioning and Disability (ICF), synthesising an outcome matrix to understand the extent of coverage of domains in the framework.

Results: Fifty-three studies were identified with information relating to outcomes measured in children exposed to viral infections in utero from the preterm period to 19 years of age. The most populated ICF-CY domain measured was structure, which included microcephaly, eye, ear and neurological structures. The functioning domain was also well reported with a wide range of tools used (Bayley Scales of Infant Development (BSID); Teller acuity cards; Auditory Brainstem Responses (ABR); Gross Motor Function Classification; sleep patterns). Few studies reported outcomes in the activities and participation domain and the environmental outcomes of the ICF-CY. A total of 34 out of 53 studies concentrated on microcephaly (58.8% defining microcephaly as OFC ≤2SD, three studies using multiple definitions, and some had no definition). Only 24.6% of studies measured other factors. Almost all of these focused on child development using (BSID) for 0-2 years and cognitive functioning utilising the Wechsler Intelligence Scale for Children (WISC) (6-17 year olds). Other outcomes measured motor functions, vision and hearing but methods of measurement and validity of those measures are vastly under-reported.

Conclusion: The review demonstrated a lack of cohesiveness and consistency in reporting and measuring outcomes. Studies focus more on health structure and functioning rather than on the other relevant social, psychological and participatory parts of the ICF which may be more relevant for families and children.
LACK OF AWARENESS ABOUT PREVALENCE OF DEVELOPMENTAL COORDINATION DISORDER (DCD) IN PRETERM INFANTS IN THE NORTHERN ALBERTA NEONATAL PROGRAMME

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Background and Aims: Developmental Coordination Disorder (DCD) has a prevalence rate of ~ 5% among school-age children, more commonly among preterm male infants born ≤32 weeks gestational age (GA). Data on DCD in Northern Alberta, Canada is lacking. We hoped to elucidate the prevalence of DCD among preterm infants in Northern Alberta and identify its associated risk factors.

Methods: Analytics, Data Integration, Measurement & Reporting (DIMR) services of Alberta Health Services provided a list of preterm infants (<37 weeks GA) born in Northern Alberta from 2000-2008, admitted to Royal Alexandra Hospital, and diagnosed with DCD based on ICD-10 code. Perinatal and neonatal variables, and developmental testing scores, were abstracted from health and school records, after parental consent. Exclusion criteria: cerebral palsy, significant cognitive delay (full-scale IQ≤70), and/or legally blind (VA<20/200). Research Ethics Board of the University of Alberta granted approval.

Results: Of the total number of infants born <37 weeks (11,787) in Northern Alberta during this study period, 150 infants were identified with DCD based on ICD-10 code. Fifty-two infants were ineligible, and 14 were lost to follow up. Although coded by HIM, 68 infants did not fulfill criteria for DCD diagnosis. Sixteen infants fulfilled the DCD diagnosis criteria with average GA 32.1±4.3 weeks (mean ± SD) and average birth weight 1790±732 g (mean ± SD), mostly prevalent in males and ≥ 34 weeks GA.

Discussion: A survey amongst a physician sample recruited from Canada, USA and UK revealed only 23–41% of physicians are familiar with DCD. Our study agreeably revealed a significantly lower number of DCD cases in preterm infants than is reported in the literature.

Conclusion: Awareness of DCD at a regional, educational, and health education management level is significantly lacking. Increasing awareness of the prevalence of DCD, its impact on daily life, and the serious consequences if left unrecognised is essential.

COMPARING P-POSSUM AND ACS-NSQIP SURGICAL RISK CALCULATOR TO BEST PREDICT UNFAVOURABLE OUTCOMES AFTER SURGERY

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Background and Aims: All surgical interventions are associated with risks of morbidity and mortality. The ability to individually risk-stratify patients preoperatively is highly desired. This project aimed to compare two risk calculators, P-POSSUM and the ACS-NSQIP Universal Calculator, at predicting unfavourable outcomes.

Methods: A retrospective analysis of 158 patients who underwent an emergency laparotomy at Salford Royal Hospital between January 2015 and June 2016 was carried out. The medical notes were reviewed in order to establish what risk prediction score would have been given to each patient by the two calculators preoperatively. This was compared with the actual 30-days outcomes and analysed with observed to expected (O: E) ratios and Brier scores.

Results: ACS-NSQIP and P-POSSUM calculators produced O: E ratios of 1.29 and 0.84 respectively for mortality, 0.88 and 0.34 respectively for serious complications, and 2.15 and 1.02 for any complications. ACS-NSQIP produced superior Brier scores for both mortality and serious complications.

Conclusion: The observed results show that the ACS-NSQIP Universal Calculator has an advantage over P-POSSUM when applied to emergency laparotomy procedures. However, it is worth noting the small sample size, limited to emergency laparotomy procedures only, which was studied. Further work should be conducted to perform a more robust comparison of the two models, which would allow increased confidence in the choice of risk calculator.
18F-FDG PET IN THE DIAGNOSIS OF VASCULAR PROSTHETIC GRAFT INFECTION: A DIAGNOSTIC TEST ACCURACY META-ANALYSIS

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Background and Aims: The diagnosis of vascular prosthetic graft infection (VPGI) is a complex process, often requiring an intra-operative peri-graft biopsy to confirm an infection. Controversy exists regarding the use of imaging techniques in the diagnostic process. We aimed to evaluate the diagnostic accuracy of 18-fluorine fluorodeoxyglucose (18F-FDG) positron emission tomography (PET) in VPGI.

Methods: We conducted a systematic search of electronic databases to identify studies assessing the use of 18F-FDG PET in the diagnosis of VPGI. We conducted a meta-analysis using a mixed-effects logistic regression bivariate model.

Results: We identified 12 studies reporting a total of 433 prostheses, out of which 202 were proven to be infected. Analysis of PETs scan was performed using five different methods: graded uptake, focal uptake, maximal standardised uptake value (SUVmax), tissue-to-background ratio (TBR), and dual-time point (DTP). The pooled estimates for sensitivity and specificity for graded uptake were 0.89 (95% CI 0.73 to 0.96) and 0.61 (95% CI 0.48 to 0.74), respectively; they were 0.93 (95% CI 0.83 to 0.97) and 0.78 (95% CI 0.53 to 0.92) for focal uptake; 0.98 (95% CI 0.42 to 0.99) and 0.80 (95% CI 0.70 to 0.88) for SUVmax; 0.57 (95% CI 0.39 to 0.73) and 0.76 (95% CI 0.64 to 0.85) for TBR; and 1.00 (95% CI 0.48 to 1.00) and 0.88 (95% CI 0.68 to 0.97) for DTP.

Conclusion: Our meta-analysis suggests that 18F-FDG PET scan has a high sensitivity in diagnosing VPGI, and its accuracy can be further increased by combining PET with computed tomography (CT).

THE EFFECT OF SOLITARY HOUSING ON MICROGLIAL CELLS - A POSSIBLE EXPLANATION BEHIND THE GLASGOW EFFECT

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Background and Aims: The circadian rhythm, or internal body clock, is a daily cycle of biological activity observed across species and is based around the 24-hour day. It has been shown that disruption of this rhythm by low levels of light at night can lead to negative health consequences such as cardiovascular disease, depression, and early mortality, among others. This study aimed to investigate the effect that Dim Light at Night (dLAN) would have on markers of microglia in the hippocampus of the mouse brain and to assess whether living in isolation has any effect on these levels.

Methods: Adult mice (n=32) were allocated to be singly housed or housed in groups of four and were randomised to a dLAN (12h dim light: 12h bright light) or control (12h dark: 12h light) group. Immunohistochemistry was performed on sections of the hippocampus for microglia using an antibody to Iba1, a marker present on microglia.

Results: There was a significant difference in immunopositivity between the different housing conditions in the hippocampus (p=0.02), with the socially isolated mice having increased Iba1 expression (2.34% vs 1.58%). There was no statistically significant difference between the lighting groups (p=0.18).

Conclusion: These results give evidence to the hypothesis that living in social isolation increases levels of inflammation in the hippocampus, an area which functions to modulate the stress response. Social isolation is postulated to prime the brain through neuroinflammation, which may lead to an increased risk of negative health effects in those who live in urban environments. This may be one of the explanations behind the ‘Glasgow effect’, as Glasgow has one of the highest levels of social isolation of any city in the UK. This phenomenon describes greater health problems in Glasgow even when compared with equally deprived cities around the UK.
A Qualitative Study to Identify the Opportunity for Health Promotion Intervention in the Emergency Department

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**Background:** Emergency department (ED) staffs frequently see patients with potentially modifiable risk factors for acute, chronic or subsequent illness. Health promotion interventions delivered in the ED have been advocated for these patients, but delivery remains suboptimal. This study aimed to compare the perspectives of doctors and nurses in the emergency department (ED) in order to recommend improved strategies and opportunities for health promotion interventions.

**Methods:** A multicentre, qualitative study was conducted in three EDs in Scotland in 2017. ED staffs at one large teaching hospital and two general hospitals were approached during handover meetings. All staff who provided direct patient care were eligible for the study (n=273) and offered a multicomponent survey. Two pilot phases of the survey conducted before deployment.

**Results:** Of the 197 respondents (72% response rate), 79 (40%) were doctors and 118 (60%) were nurses. More doctors (86.1% CI [76.8-92.1]) than nurses (51.7% CI [42.8-60.5]) report offering health promotion interventions; specifically, for alcohol misuse, smoking, drug misuse and sexual health interventions. Time constraints (n=172, 87.3%) and a lack of health promotion infrastructure (n=100, 50.7%) in the ED were found to challenge widespread delivery (groups not mutually exclusive). Staff felt patients whose presentation was directly related to smoking and alcohol/drug misuse or patients with new-found hyperglycaemia should be prioritised for brief interventions.

**Discussion:** This is the first staff perspective study in the UK and shows encouraging rates of health promotion intervention being reported by ED staff, exceeding those previously reported in US and Australian studies. Staff acknowledge the benefit of health promotion, in agreement with other studies, but time constraints and insufficient ED resources are unanimously recognised as barriers to practice.

**Conclusions:** Staffs require additional training in brief intervention techniques and treatment options to enhance their delivery of preventive care in the ED.

AN ANALYSIS OF THE CAUSES OF DEATH IDENTIFIED AT AUTOPSY IN THE OBESE POPULATION

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**Background and Aims:** Obesity is the fifth largest risk factor impacting on global mortality and its incidence is rising. Contribution of obesity to death rates is only measurable if included on death certificates. Obesity causes deaths directly e.g. obesity cardiomyopathy (OCM), and indirectly as a risk factor for coronary heart disease (CHD) and other conditions. In this study, we aim to investigate the reporting of obesity and its inclusion in death certificates in a single centre coronial autopsy service.

**Methods:** Retrospective review of autopsy reports in the Oxford pathology database across a three-year period (2014-2016). Autopsy reports were reviewed for height, weight and BMI, prevalence of obesity and obesity-specific conditions, all-cause mortality, CHD-related mortality, and mean age of death from CHD in different BMI categories.

**Results:** Height and weight were omitted without adequate reason in 14% of reports analysed (n=1,514). Obesity is poorly recognised on death certificates where present (just 5.1% overall). Identification of OCM in the morbidly obese is rising: 6.6% compared to 2.0% in the previous largest study to date. In total, 739 (40% n=1,868) autopsies were performed on obese individuals. Obesity/obesity-specific pathology where included in the death certificates in 0.2% of obese (BMI 30-35), 7.4% of severely obese (BMI 35-40) and 25.7% of morbidly obese (BMI >40) individuals.

**Discussion:** CHD accounted for 26.4% of deaths in morbidly obese individuals and 20.7% of deaths in those of normal BMI. Strikingly, morbidly obese individuals died from CHD on average nine years earlier (mean age of death 68 years) compared to those of ideal BMI, mean age of death 77 years (p=0.000004, 95%CI 5-13); this effect was not accounted for by concurrent presence of diabetes or hypertension.

**Conclusion:** This study links obesity to earlier death from CHD and indicates that obesity is under-recorded on death certificates by pathologists.
THE MANAGEMENT OF ST-ELEVATED MYOCARDIAL INFARCTION IN A MIDDLE-INCOME SOUTH ASIAN COUNTRY: A MULTI-CENTRE STUDY

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Background and Aims: ST-elevated myocardial infarction (STEMI) is a leading cause of death in Sri Lanka and its outcomes are not well documented. The United Kingdom is a developed country whose treatment and management modalities differ significantly. This study aims to characterise the presentation and outcomes of STEMI patients in Sri Lanka, investigate the predictors of adverse outcomes, and compare the treatment pathways between the two countries to draw recommendations.

Methods: A retrospective analysis of 205 patients presenting over three months to a major tertiary care centre in Sri Lanka was performed. Data from the UK originated from the PCI database at the Queen Elizabeth Hospital, Birmingham, UK. Logistic regression was conducted to determine predictors of adverse outcomes.

Results: The rate of in-hospital mortality in Sri Lanka was 10% significantly higher than in the UK. Thrombolysis was the mainstay of treatment in Sri Lanka. The estimate of late presenters was 27%. Sri Lankans were significantly younger than the British, with higher rates of diabetes and hypertension but lower rates of smoking. Country was not a predictive factor of in-hospital mortality among the Asian ethnicity.

Discussion: The high rate of in-hospital mortality of STEMI patients in Sri Lanka, combined with the frequency of late presentation and thrombolysis, highlight the need for a paradigm shift in the treatment, management and education of the population. The mortality of Asians presenting with STEMI was not significantly improved by being in the UK, despite its sophisticated health system. Although further research is needed to conclude, this indicates the need for targeted interventions in high risk groups within the UK, including Asians.

Conclusion: Implementation of primary care, patient education and pre-hospital care are important recommendations to improve outcomes in Sri Lanka. Research regarding the effectiveness of interventions in Asians within the UK is recommended.

IDENTIFYING FACTORS OF MICROPARTICLES MODIFIED WITH ARGinine DERIVATIVES THAT INDUCE PHENOTYPIC SHIFTS IN MACROPHAGES

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Background and Aims: Macrophages are key players in the progression of many diseases, ranging from rheumatoid arthritis to cancer. Drug delivery systems have the potential not only to transport payloads to diseased tissue, but also to influence cell behaviour. In microtumour environments, macrophage phenotypes become inappropriately shifted towards their angiogenic form, promoting the growth of cancerous tissue with the development of new blood vessels. If this behaviour could be controlled to avoid this, or even shift to more inflammatory behaviour, then tumour growth could be hindered. For example, topical microparticle treatment with this effect might be applied in surgical oncology to treat post-excision sites and reduce the chances of any satellite tumour particles from establishing themselves again. This study aims to identify factors of microparticles modified with arginine derivatives that induce phenotypic shifts in macrophages.

Methods: Poly(N-isopropylacrylamide-co-acrylic acid) (pNIPAm-co-AAc) microparticles were modified with 14 different arginine derivatives. These particles were then incubated with interleukin-4 or lipopolysaccharide-stimulated macrophages or naive macrophages (RAW264.7). The phenotypic state of the macrophages was assessed by measuring arginase activity, tumour necrosis factor-α (TNF-α) secretion, and nitrite production.

Results: Partial least-squares analysis revealed material properties and descriptors that shifted the macrophage phenotype for the three cell conditions in this study. Material descriptors relating to secondary bonding were suggested to play a role in shifting phenotypes in all three macrophage culture conditions.

Conclusion: These findings suggest that macrophage responses could be altered through drug delivery vehicles, and these descriptors identified to be involved could be employed to assist in screening further potential candidates.
PREVALENCE OF DRY EYES IN PREGNANT WOMEN IN COMPARISON WITH NON-PREGNANT WOMEN

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Background and Aims: Pregnancy is a physiological state in a woman which alters all the systems in the body. Ocular changes such as dry eye, though mostly reversible, can cause enough discomfort in a patient to actively seek medical help. This study aims to establish whether there is a significant association between the occurrence of dry eyes and the pregnant state by comparing the prevalence of dry eyes in pregnant women with that in non-pregnant women of the same age group.

Methods: It is a cross-sectional, observational, case-control study with 400 participants. The test group consisted of 200 pregnant women whilst the control group comprised of 200 non-pregnant women. These participants were given a questionnaire that included the Ocular Surface Disease Index Score (OSDI score). They were also subjected to Schirmer's test (type 1). The paired t-test was used to test the significance of the hypothesis. Additional parameters have been used to determine the relation of dry eyes with the age, trimester and the parity in pregnant women.

Results: In the test group, 19 women were Schirmer's positive while, in the control group, 16 women were Schirmer's negative. The p value=0.007 was obtained, which is statistically significant. Observed data indicates a higher prevalence in primigravida females, pregnant women in the age group of 25-29 years, and pregnant women in their first trimester.

Conclusion: The study indicates that the relation between the prevalence of dry eyes and the pregnant state is statistically very significant. Hence, the hypothesis that the pregnant state is associated with a higher prevalence of dry eyes should be accepted.
DOES GRADE OF HALLUX RIGIDUS CORRELATE WITH FAILURE RATE FOLLOWING MINIMALLY INVASIVE DORSAL CHEILECTOMY?

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Background and Aims: Hallux rigidus is a degenerative condition of the first metatarsophalangeal (MTP) joint, typically resulting in impingement pain and limitation of movement of the first MTP joint due to a dorsal osteophyte. Following failure of conservative treatment, traditionally a dorsal cheilectomy can be performed for patients in early stages of hallux rigidus. Minimally invasive forefoot surgery is becoming more popular with advances in technology and it is now possible to perform a minimally invasive dorsal cheilectomy (MIDC) of the first metatarsal with a wedge burr. The aim of this study was to determine if grade of hallux rigidus correlates with failure rate following MIDC.

Methods: We retrospectively analysed patients in our centre who had a symptomatic hallux rigidus treated with MIDC following failed conservative treatment between 2011 and 2016. They were graded preoperatively by Coughlin and Shurnas Classification.

Results: There were 89 patients (98 feet; 44 right, 36 left, 9 bilateral) in our series. The average age was 54 (range: 29 – 71) years old, with 25 male and 64 female. The Coughlin grades were as follows: Grade 1, n=33; Grade 2, n=54; Grade 3, n=11. Seven feet went on to have a first MTP joint fusion for ongoing pain (Coughlin Grade 2, n=6; Grade 3, n=1). There was no correlation between grade of hallux rigidus and failure of MIDC (p=0.38, chi square test).

Conclusion: Grade of hallux rigidus does not correlate with failure of MIDC, although there was no failure in Grade 1.

A CLINICAL AUDIT ON PRIMARY POSTPARTUM HAEMORRHAGE IN NHS TAYSIDE

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Background and Aims: Primary postpartum haemorrhage (PPH) is defined as vaginal blood loss of 500ml or greater within 24 hours of delivery. PPH occurs frequently in the UK and is the leading cause of maternal death in developing countries. The objective of this clinical audit is to assess primary PPH risk factors and evaluate current PPH management in NHS Tayside.

Methods: The design is a retrospective audit for 15 weeks. Participants had vaginal delivery with PPH of 500-1500ml. Current practice is compared with a new PPH management checklist developed by MCQIC alongside Scottish Patient Safety Programme. Main outcomes measured are antenatal risk factors, perinatal risk factors and management from the checklist. Data was collected through obstetric notes.

Results: 83 patients were included. The most prevalent antenatal risk factor was anaemia (34.9%). Other common antenatal risk factors were antepartum haemorrhage (30.1%) and previous PPH (21.7%). The largest perinatal risk factor was labour induction (53%). Comparison between expected and achieved standards showed 100% attainment in administering uterotonics, genital tract inspection and blood transfusion. Some of the least achieved standards were uterine massage, emptying bladder and IV fluids.

Discussion: Patients at high risk of PPH require specialised care. The multidisciplinary team needs to assess risk factors regularly and anticipate PPH in these patients. They should ideally deliver in a consultant-led unit. Modifiable risk factors such as anaemia should be addressed before delivery. PPH management was generally well conducted in NHS Tayside, but quick management steps such as uterine massage and bladder emptying were not always documented.

Conclusion: Regular risk assessment and prompt management of PPH is crucial for good prognosis. Introducing risk assessment forms in antenatal and intrapartum care can improve recognition of risk factors. The use of checklists during PPH can help documentation. Implementing the new checklist or a similar tool for PPH is targeted to enhance the current practice.
EVALUATING THE ADHERENCE OF PRIMARY HEALTHCARE CENTRES IN GAZA STRIP TO THE WHO-PEN PROTOCOL 1: A CROSS-SECTIONAL STUDY

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Background and Aims: Cardiovascular disease is common in the general population. It is a multi-factorial disease. Certain risk factors, such as family history, gender, ethnicity and age, cannot be changed. Other risk factors, including high blood pressure, high cholesterol and diabetes, are modifiable. This study evaluates the adherence of primary healthcare centres in Gaza to the WHO-PEN protocol.

Methods: A cross-sectional study involving 200 patients who were already diagnosed with NCDs was conducted at healthcare centres in Gaza. Data was collected retrospectively using a self-designed questionnaire based on the WHO-PEN checklist. Patients’ files were selected randomly.

Results: Based on the analysis of whole cohort (200 cases). The prevalence of DM was 39% and hypertension was 28.5% whereas 32.5% had both in 2017. There were only 17 smokers among patients, representing 8.5% of the sample. Using WHO/ISH - WHO/International Society of Hypertension - risk prediction charts, half of patients had a risk of less than 10%. On the other hand, 10% of patients had a risk of over 40% which means they need to be given statins. However, 1% did not receive statins. In patients under 55 years old, there was over prescription of baby aspirin. Conversely, diuretics were not used as addressed in WHO-PEN protocol. Interestingly, there were four patients of the whole sample on no medications. All patients were counselled on diet, exercise and smoking cessation.

Discussion: There were slight differences between the two centres. In the first centre, there was better prescription of statins. Contrariwise, in the second centre, drug combination was conserved for advanced cases. Both centres had good counselling and follow-up. Eventually, both show good adherence to WHO-PEN protocol in practice as well as documentation.

Conclusion: Results demonstrate high adherence to WHO-PEN protocol in these centres, reflecting a high quality of care and follow-up. However, there were some deficiencies in the risk estimation, which should be documented for better counselling for patients with high risk.

ASSESSMENT OF PREDISPONING RISK FACTORS FOR SURGICAL SITE INFECTION IN PATIENTS ADMITTED TO THE SURGICAL WARDS OF BAGHDAD TEACHING HOSPITAL

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Background and Aims: Surgical site infections are the most common nosocomial infection in surgical patients, and contribute to an increase in postoperative hospital stay and cost of care. Previous studies have identified some of the risk factors, including diabetes, steroid use and obesity. This study aims to assess some of these risk factors.

Methods: A hospital-based study took place in the Baghdad Teaching Hospital. Data collection was by a paper survey with interviewing of both patients and the surgical teams that are directly responsible for them. Cases were patients with surgical site infection admitted and treated in the surgical wards during the period from 1st July to 20th September 2017. Controls were conveniently selected as patients undergoing surgeries at Baghdad Teaching Hospital and followed up for 30 days for exclusion of the development of surgical site infection. For each participant, pre-surgical, post-surgical and demographic parameters were collected. Each parameter was compared between cases and controls for significant difference using IBM SPSS version 22.

Results: Based on the analysis of whole cohort (200 cases). The prevalence of DM was 39% and hypertension was 28.5% whereas 32.5% had both in 2017. There were only 17 smokers among patients, representing 8.5% of the sample. Using WHO/ISH - WHO/International Society of Hypertension - risk prediction charts, half of patients had a risk of less than 10%. On the other hand, 10% of patients had a risk of over 40% which means they need to be given statins. However, 1% did not receive statins. In patients under 55 years old, there was over prescription of baby aspirin. Conversely, diuretics were not used as addressed in WHO-PEN protocol. Interestingly, there were four patients of the whole sample on no medications. All patients were counselled on diet, exercise and smoking cessation.

Discussion: Surgical site infection showed to be of higher occurrence in diabetic and obese patients. Higher care for such patients should therefore be implemented to reduce their risk. Increased care of pre-operative prophylactic measures is recommended as it showed to be of effect in reducing the risk of infection.

Conclusion: Surgical site infection showed to be of higher occurrence in diabetic and obese patients. Higher care for such patients should therefore be implemented to reduce their risk. Increased care of pre-operative prophylactic measures is recommended as it showed to be of effect in reducing the risk of infection.
A PRIMARY AND SECONDARY CARE COLLABORATION TO IMPROVE THE MANAGEMENT OF INTERMITTENT CLAUDICATION

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Background and Aims: First line treatment of patients with intermittent claudication (IC) should be optimal medical therapy (OMT): a combination of cardiovascular risk modification and a supervised exercise programme (SEP). Further investigation of IC patients is only indicated if vascular intervention is being considered. Our aim was to review the value added to patient care by referral of IC patients to a vascular surgeon.

Methods: Both quantitative and qualitative evaluations of IC patients’ care were undertaken by: a review of IC referrals, interviews with the patients, and GP interviews. New IC referrals over a 10-week period were reviewed. Data was extracted regarding OMT, definitive diagnosis and investigation initiation. Qualitatively, 10 patients at clinics and seven GPs via telephone were interviewed. Themes were identified as understanding and management, exercise therapy opinions, and expectations of secondary care.

Results: Thirty-eight referrals were reviewed and showed only 60% were on antiplatelet therapy (APT), 68% were on a statin, whilst 32% were current smokers and 37% included ABPIs. Only two referrals mentioned exercise therapy; this was not SEP but non-specific advice. Interviews showed that patients lacked an understanding of their condition and its best clinical management. GP interviews highlighted that current management could be significantly improved to ensure OMT is delivered at the earliest opportunity.

Conclusion: The majority of IC patients do not need to see a vascular surgeon and should be referred directly to a SEP. Value added care should focus on OMT, smoking cessation and exercise. Subsequently, Primary and Secondary Care together have developed a new IC management pathway, as GPs can now refer directly to the National Exercise Referral Scheme which provides the SEPs. An educational patient leaflet emphasising the importance of SEP has been developed. All GP practices now have received hand-held Dopplers and practice nurses are being trained to measure ABPIs.

UNCONTROLLED HYPERTENSION; FROM DETERMINANTS TO FIRST ENCOUNTER MANAGEMENT AT EMERGENCY UNIT OF BAGHDAD TEACHING HOSPITAL

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Background: Hypertension is uncontrolled when blood pressure reading is more than 140/90 mm Hg. This condition places the individual at a risk of developing hypertensive crisis, whether emergency or urgency.

Aim: To identify the proportion and evaluate the management of uncontrolled hypertension in the emergency department and review some characteristics of uncontrolled hypertensive patients.

Methods: Four hundred and nine cases presented with blood pressure (>140/90 mm Hg), irrespective of past medical history of hypertension, to the emergency unit at Baghdad Teaching Hospital were included in this retrospective cross-sectional study. Data on socio-demographics, chief complaints, past medical history, work-up, management with anti-hypertensive and disposition was collected from patient records in the archives between July and September 2017.

Results: Cases with uncontrolled hypertension formed 6.56% of all admissions to emergency department. Means of systolic and diastolic blood pressure readings were 169.98 and 96.66 mm Hg, respectively. In this study, 45.47% were male and 54.52% were female. Average age was 54.95. One hundred and fifty-one (2.4%) were cases of hypertension crisis among emergency unit attendants. Diabetes mellitus was the single most common past medical condition among all cases. Shortness of breath was the main chief complaint (22%) in hypertension crisis cases; chest pain was the most common presenting complaint in all included cases. Management was not documented in records of 214 (50%). However, in most of the remaining cases, 137 (70%) consisted of: loop diuretics (furosemide), angiotensin-converting enzyme inhibitors in 22 (11%), calcium channel blockers in 17 (8%), and beta blockers in 13 (6%).

Conclusion: One third of patients who attended the emergency department with elevated blood pressure were cases of hypertension crisis. Diabetes mellitus was the major co-morbidity. Local practice in management was not compliant to the guidelines and severity but subjected to the availability of medication.
AN EVALUATION OF LOCAL SMOKING CESSATION THERAPIES PRESCRIBED BY DOCTORS, THEIR KNOWLEDGE AND THEIR PRESCRIPTION TENDENCIES

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Background and Aims: Smoking is a common practice which negatively affects one’s health in increasing the risk of cancers (such as lung and bladder). It is also a major risk factor for cardiovascular disease. Locally, one can find nicotine replacement patches (NRP), lozenges (NRL), gums (NRG) and inhalers (NRI), as well as varenicline and bupropion, as smoking cessation therapies to quit smoking. This audit focused on NRP and varenicline, which are the two commonest therapies used locally. It aims to evaluate the knowledge and prescription tendencies of doctors to local smoking cessation therapies.

Methods: Data was collected in the form of questionnaires, with the questions being related to doctors’ prescription tendencies, their knowledge and their most commonly prescribed medication.

Results: One hundred and ten doctors (54 junior doctors, 32 senior doctors and 24 specialists) participated in the questionnaire. Most of the junior doctors had heard of only NRP (56%) and only 31% had heard of varenicline, whilst the specialists had heard of all the available modes of medications and therapies (75% for NRP and 75% for varenicline). All doctors were more likely to prescribe nicotine replacement patches (68% for junior doctors, 56% for senior doctors and 35% for specialists), but specialists listed varenicline as a close second (28%). Thirty-one percent of junior doctors, 21% of senior doctors and 25% of specialists were unaware of the side effects of NRP, whilst 65% of junior doctors, 34% of senior doctors and 25% of specialists were unaware of the side effects of varenicline.

Discussion: To further evaluate knowledge, the doctors were asked about the correct way of prescribing both NRP and varenicline. For the former, 28% of junior doctors, 57% of senior doctors and 75% of specialists answered correctly whilst, for the latter, only 20% of junior doctors, 53% of senior doctors and 50% of specialists answered correctly.

Conclusion: Nicotine replacement patches remain the treatment of choice amongst doctors locally for stopping smoking.

HEART FAILURE DISCHARGE LETTERS: A QUALITY IMPROVEMENT PROJECT

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Background and Aims: Heart failure (HF) is an increasingly prevalent condition due to the ageing population and improved acute cardiovascular therapies. HF patients have frequent exacerbations and consequent readmissions. The discharge letter summarises the admission details, and is useful for both primary and secondary care physicians, as well as for clinicians, during readmission episodes. The content of such letters is often written with varying quality. This study implements a quality improvement project (QIP) approach using “Plan-Do-Study-Act” (PDSA) cycles with an aim to improve the quality of HF discharge letters.

Methods: Existing letters were retrospectively analysed objectively using an evidence-based ten-point scale (cycle 1). Further cycles included: education of cardiology junior doctors (cycle 2); stratification of data by discharging team to analyse for any discrepancies between specialties (cycle 3); targeted education and posters to general internal medical (GIM) junior doctors (cycle 4); further stratification of data by discharge specialty (cycle 5).

Results: Cycle 1 data demonstrated a mean score of 4.7/10 (n=20), which increased to 6.2/10 (n=44) in cycle 2. In cycle 3, stratification demonstrated a change in discharge summary quality by the cardiology team (cycle 1 mean 4.7/10; cycle 2 mean 6.8/10, n=47) in contrast with those patients discharged by GIM specialties (cycle 1 mean 4.5/10; cycle 2 mean 4.2/10, n=17). Cycle 4 showed an improved overall mean score of 6.3/10 (n=32). Final stratification in cycle 5 demonstrated improvement in both cardiology and GIM specialties (cardiology mean 7.5/10, n=21; GIM mean 5.4/10, n=11).

Discussion: This study demonstrates that a QIP can be used to improve the content of HF discharge summaries with simple interventions and to identify areas of suboptimal practice.

Conclusion: Our simple interventions have substantially improved HF discharge summaries at our Trust. Changes to the electronic discharge letter template would ensure that these improvements have longevity.
THE ROLE OF PLAIN ABDOMINAL X-RAYS IN ACUTE ABDOMEN

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Background and Aims: This audit was carried out to ascertain the value of abdominal X-rays in finding the cause of acute abdomen, clinician's compliance to guidelines, and if further imaging was performed.

Methods: Data was collected retrospectively using RIS and included A&E patients who had an abdominal X-ray in September. Two hundred and twenty-one patients were included. Data including the indication, findings and evidence for further imaging were collected from PACS.

Results: Out of 221 films, 135 met the standard for AXR whilst 86 did not meet the standard. Out of 135 films, 122 were reported as normal and 13 as abnormal. Of the normal AXRs, 29 patients went on to have further imaging, 14 of which were normal CT scans. The remainder were a mix of pathologies. All 13 patients with X-ray abnormalities had CT scans; 10/13 were abnormal.

Discussion: One hundred and twenty-two of 135 patients who met the criteria had normal AXRs. This amounts to 88% sensitivity of AXR. The reason for which further imaging was performed despite this requires further investigation. Determining the grade of the requesting clinician may be valuable. In patients with normal AXR who still had a CT, only 53% were reported as being normal. Abnormalities such as bladder perforation and caecal mass were noted on CT. In the majority of these cases, the indication had been pain. Bypassing an AXR in such serious cases for an immediate CT scan may have been more appropriate.

Conclusion: In conclusion, AXRs are often requested inappropriately. This audit highlights the need for educational posters about indications for AXR being circulated. The use of various imaging modalities should be reiterated at radiology meetings or incorporated into E-learning for doctors.

FACTORS AFFECTING QUALITY OF HISTOLOGICAL SPECIMENS IN LAPAROSCOPIC ANTERIOR RESECTIONS - A RETROSPECTIVE COHORT STUDY

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Background and Aims: Laparoscopic resections for colorectal cancer are common. High quality specimens are determined by a clear resection margin (R0 resection) and adequate numbers of lymph nodes. One large study showed R0 rates of 86.7%. We aimed to explore the factors that could potentiate obtaining poor quality specimens during laparoscopic colorectal cancer surgery.

Methods: We performed a retrospective cohort study over one year. Patients undergoing laparoscopic anterior resection by one consultant surgeon at Wigan were included. Data was extracted from online notes. Gender, tumour, R0/R1, lymph nodes resected, distance from anal verge and pTNM were recorded.

Results: Thirty patients were included. R0 resection rate was 93% and the median number of lymph nodes was 13 (range 2–28). On analysing the two R1 resections, they were low anterior resections of T3 tumours with the mean distance from the anal verge 7.6cm. Mean distance from the anal verge for all other anterior resections was 13.7 (range 6.1–35cm). Both R1 resections were male. The mean size of the tumour for all anterior resections was 3.62, while that of the R1 tumours was 2.45cm. Only one patient who had an anterior resection had less than five lymph nodes isolated. The distance of this tumour from the anal verge was 8.8cm and the size of the tumour was 0.3cm. The patient was male. 33% of patients who had an anterior resection had less than 10 lymph nodes isolated. Their mean distance from the anal verge was 14.7cm and the mean size of the tumour was 3.22cm. 78% were male.

Discussion: Smaller tumour size, shorter distance to the anal verge, and male gender appear to increase the risk of obtaining a poor-quality specimen.

Conclusion: The results appear to show factors affecting specimen quality following laparoscopic colorectal cancer resection are: gender, distance from the anal verge, and tumour size.
A CLOSED LOOP AUDIT OF PRE-OPERATIVE ADMINISTRATION OF ANTIBIOTICS: WHERE SURGICAL OPINION AND GUIDELINES DISAGREE

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Background and Aims: Antimicrobial resistance has risen, driven by inappropriate and over use. The aim of this audit was to reduce elective surgery antibiotic consumption within a DGH to meet the “Antimicrobial Resistance CQUIN 2017/18”.

Methods: Retrospective data collection of antibiotic used, indication, administration time and adherence to local guidelines was undertaken (October 2017). Ten elective patients were randomly selected from each sub-specialty (Breast, Colorectal, Ear, Nose & Throat/Mailllo-Facial, General Surgery/Vascular, Gynaecology, Orthopaedics, Upper Gastro-Intestinal, and Urology). Findings were presented locally, the guidelines were updated, then re-audited prospectively in 2018.

Results: Originally, 52.5% (n=42) patients were given antibiotics. Sixty-four percent (n=27) were inappropriate (not required/incorrect antibiotic). Thirty-three percent (n=15) received Tazocin. On re-audit, 55% (n=44) received antibiotics; 61% (n=27) inappropriately, with no Tazocin prescriptions. Original guidance advised prophylaxis be administered 30-60 minutes before knife to skin (KTS); only 26% (n=11) adhered. After feedback, Trust policy followed national guidelines, suggesting antibiotics be given as close to KTS as possible, preferably within one hour; 100% (n=44) followed the policy. Of 38 (47.5%) patients not given prophylaxis, original guidelines suggested 19 (50%) should receive antimicrobials. Sub-specialty case reviews deemed none necessary, but a guidance was not amended. On re-audit, of 36 (45%) patients without antibiotics, potentially two (1.3%) required antimicrobials.

Discussion: Preserving institutional microbiology knowledge is difficult: doctors frequently rotate hospitals, and accessing information is time consuming and often impractical. Consequently, clinicians either use a “one-size-fits-all” antimicrobial – a broad-spectrum antibiotic - or align with national guidelines. Although local broad-spectrum antibiotic recommendations remained unchanged, on re-audit no Tazocin was administered. However, inappropriate prophylaxis increased. This may reflect increased clinical awareness of inappropriate Tazocin prescribing following audit presentation, with selection of alternative agents.

Conclusion: Despite policy changes, our local guidelines still over-estimate prophylactic antibiotic requirements and do not reflect sub-specialty opinion. Implementation of a single, national guideline could ensure appropriate antibiotic prophylaxis. Regular re-audit and presentation publicises recommendations and encourages adherence.

AN AUDIT ON PREOPERATIVE FASTING TIMES IN PATIENTS ON THE ACUTE TRAUMA LIST

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Background and Aims: Patients are fasted preoperatively in order to prevent aspiration and regurgitation of gastric contents. Guidelines recommend patients fast six hours from food and two hours from fluids preoperatively. However, excessive fasting on emergency lists is common. Prolonged fasting has shown to increase patient discomfort, prolong wound healing, increase hypoglycaemic episodes, cause electrolyte imbalances, and result in longer hospitalisation. Research has shown that giving carbohydrate drinks two hours before procedures is safe and reduces postoperative complications. The aim of this audit is to assess the preoperative fasting times in patients on the trauma list to prevent prolonged fasting.

Methods: A prospective study was conducted in which we looked at patients operated on the surgical trauma list over a two-month period. Patient demographics, co-morbidities, and the operation performed were collected from patient notes. Information on preoperative fasting times were recorded using patient notes and from directly questioning patients.

Results: A total of 104 patients were identified from this study. The average time for which patients were fasted was 14 hours and 28 minutes. The maximum time for which patients were fasted was 26 hours and 46 minutes. Overall, this study showed that 98% of patients were excessively fasted preoperatively. A significant portion of patients had co-morbidities such as diabetes that would be affected by prolonged fasting or were at increased risk of malnutrition.

Discussion: The results from this audit show that the majority of patients on the trauma list are being excessively fasted. This prolonged fasting can have effects on patient morbidity and increase the likelihood of postoperative complications. Better communication between teams and new protocols on fasting patients on the trauma list can help address this. In addition, the use of carbohydrate rich drinks two hours before operations can help reduce excessive fasting and improve patient outcomes.
DOES ADJUVANT SUB-TENON TRIAMCINOLONE (SSTA) AID IN THE MANAGEMENT OF DIABETIC MACULAR OEDEMA?

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Background: DMO is a major cause of visual impairment. There remains a need to improve care.

Aims: Does SSTA injection affect the number of intravitreal anti-VEGF injections needed for treatment of DMO?

Methods: One hundred six eyes from 56 patients from one vitreoretinal surgeon were studied retrospectively between 01/12/2015 and 30/12/2016. Inclusion criteria: adequate media clarity for fundus visualisation, verification of macular oedema by optical coherence tomography (OCT), and absence of coexisting retinal disease. Analysis was done using Mann-Whitney U test with p<0.05 as significant.

Results: Ninety-two eyes received anti-VEGF alone; the mean number of anti-VEGF injections needed was 2.74 in one year. Fourteen eyes received adjuvant SSTA; the number of anti-VEGF injections needed was 1.42. On average, patients receiving adjuvant SSTA required less anti-VEGF injections (p=0.005). For eyes receiving SSTA, baseline mean logMAR VA was 0.82 (Snellen equivalent 20/132) and Corneal macular thickness (CMT) was 464 microns. At 12 months, logMAR VA was 0.64 (Snellen 20/87), and CMT was 294 microns. For eyes that did not receive SSTA, baseline logMAR VA was 0.64 (Snellen 20/87), and CMT was 290 microns. 3/14 patients receiving SSTA developed non-sight threatening rise in IOP requiring anti-glaucoma medications (21%). None required cataract surgery.

Conclusion: Anti-VEGF with adjuvant SSTA appears to reduce the recurrence rate of DMO and, significantly, the need for more anti-VEGF injections by almost half. Side effects are manageable. The benefits clearly outweigh the risks.

THE INITIAL MANAGEMENT OF PATIENTS MEETING THE SIRS CRITERIA AT A LARGE TERTIARY CARE TEACHING HOSPITAL

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Background and Aims: Sepsis is the initiation of the systemic inflammatory response syndrome (SIRS) due to infection. It is a major cause of patient mortality and morbidity. Sepsis may present as changes in vital signs: the SIRS criteria. The UK Sepsis Trust guidelines state that patients fulfilling two of the SIRS criterion should be screened for sepsis and treated if necessary. This service evaluation aimed to retrospectively assess the management of patients meeting the SIRS criteria at a tertiary care hospital.

Methods: From a larger study involving approximately 30,000 patients, 39 were randomly selected. It was assessed whether: patient notes indicated if infection was considered; any relevant investigations for infection were performed; and any antimicrobial treatment was administered. The time at which each of these occurred was also assessed to determine the quality of care received.

Results: Consideration of infection was documented in only 48.7% of subjects. However, 87% of patients received at least one of the following: infection indicated in the notes; an investigation; or antimicrobial treatment. The median time to perform an investigation for infection was longer than the time taken to administer antimicrobials (23.5 and 8.4 hours, respectively). In addition, fewer patients had investigations performed (56.4%) than antimicrobials prescribed (76.9%). In 13%, no investigation was performed or antimicrobial prescribed.

Discussion: In some subjects, sepsis may have simply not been considered. However, the proportion of patients receiving investigations and antimicrobials suggests that infection was often considered but poorly documented, and that antimicrobials were mostly administered empirically. In some cases, infection screening may not have been performed because the patient’s clinical presentation suggested this was unlikely.

Conclusion: It is recommended that hospitals utilise alerting systems, such as NEWS, to identify patients meeting the SIRS criteria. In addition, a sepsis tool should be made available to aid documentation and guide management.
AN AUDIT ASSESSING THE IMPACT OF TIME FROM PRESENTATION TO PYELOPLASTY UPON MAG3 RENOGRAM OUTCOME AND THE BENEFIT OF REPEAT RENOGRAMS

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Background: A MAG3 renogram/scan investigates the drainage of urine from the kidneys into the bladder. It can also be used to check for reflux. Pyeloplasty is the surgical reconstruction of the renal pelvis in order to drain and decompress the kidney. The purpose of the surgery, in almost all cases, is to relieve a uretero-pelvic junction obstruction.

Aims: To compare the outcomes of pyeloplasty in those with split function with those who have a poorly functioning kidney.

Methods: We compared the outcomes of pyeloplasty in those with split function of >30% with those who had a poorly functioning kidney (split function ≤30%). A total of 175 patients were included. The mean length of FU was 573 days. The mean pre-operative split renal function was 39.4%. Thirty-five patients had split function of ≤30%. The mean waiting time from presentation to operation was 207 days. The mean interval between pyeloplasty and first post-operative renogram was 118 days. The second post-operative renogram was performed at a mean interval of 355 days following the first scan.

Results: A total of 89 patients had split function documented in one pre-operative and two post-operative renograms. Seventy-two cases (81%) showed stable or improved function on their first post-operative renogram (58 stable; 14 improved), and the second post-operative renogram showed renal function remained stable or improved in 68 cases (95%). In the remaining four patients, the split function only marginally deteriorated (average of -7%). Of the 17 cases with worsening renal function on the first post-operative renogram, the split function continued to deteriorate in only two cases. Patients who had worsening split function post-operatively had a longer waiting time for their procedure (the majority of those who waited over 200 days had poor functional outcomes). Pain was the most common presenting symptom (80% of cases); 79.2% of these patients had symptoms post-operatively. Those with split function of ≤30% showed higher levels of pain following the procedure (26.7%) compared to those with better function (20.8%).

Discussion: Longer waiting times from presentation to definitive treatment appear to have a significant negative impact on split renal function post-pyeloplasty. The majority of patients received functional and symptomatic improvement post-pyeloplasty, but there was no significant difference in outcome between those with split renal function of ≤30% compared to those with better renal function. Those with favourable results on the first post-operative renogram may not require further imaging.

CHRONIC LYMPHOCYTIC LEUKAEMIA (CLL): AN AUDIT OF COMPLIANCE WITH KEY ASPECTS OF BRITISH SOCIETY FOR HAEMATOLOGY (BSH) GUIDELINES ON DIAGNOSTIC WORK-UP

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Background: Chronic lymphocytic leukaemia (CLL) is a malignant clonal disorder of B lymphocytes with levels greater than 5 x 10^9/L (5 x 10³/microliter) involving the bone marrow and peripheral blood. The incidence of CLL increases with age. Upon diagnosis, British Society for Haematology (BSH) recommends that all patients should undergo specific tests such as full blood count, reticulocyte count, direct antiglobulin test (DAT), immunophenotype, and serum immunoglobulins. In addition to this, screening for hepatitis B and C, TP53 deletion and a baseline CT scan should be performed prior to treatment.

Aims: The aim of the audit is to assess the compliance with key aspects of BSH guidelines on diagnostic work-up for CLL.

Methods: Electronic records for 32 patients diagnosed with CLL between January 2016 and May 2017 in Royal Albert Edward Infirmary (England) were analysed.

Results: The demographics show a higher proportion of males (59% - 19 of 32 patients) compared to females (41% - 13 of 32 patients). Similar studies show a male-to-female ratio of 2:1 which is comparable to our results. The mean age at diagnosis was 72 years. Three patients (9.4%) had haemoglobin levels below 100 g/l of which 90% had a reticulocyte count and direct antiglobulin test done. All 32 patients had an immunophenotype typical of CLL. Eight patients (25%) did not have their serum immunoglobulins checked. Three of 32 patients (9%) received treatment with FCR (Fludarabine, Cyclophosphamide, Rituximab) as first line therapy. Before treatment, all three patients had screening for hepatitis B and C, TP53 deletion and baseline CT scan.

Conclusion: We suggest that patients with low haemoglobin levels should have direct antiglobulin test and reticulocyte count performed. All patients should have their serum immunoglobulins checked. A diagnostic work-up and pre-treatment checklist should be designed to guide specialist teams.
EVALUATION OF DEATH CERTIFICATION PRACTICES AT WEST WALES GENERAL HOSPITAL - AN AREA FOR QUALITY IMPROVEMENT?

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**Background and Aims:** As outlined by the Office for National Statistics' (ONS) Death Certification Advisory Group, the quality of death certification conveys several epidemiological and clinical implications. More specifically, this data is often implicated in population health monitoring and in the design and evaluation of Public Health interventions, to mention but a few. Subsequently, our aim was to analyse current documentation practices and improve the quality of Medical Certificates of Cause of Death (MCCDs) and Death Advice Letters (DALs) at West Wales General Hospital.

**Methods:**

**Patient Selection**
Retrospective analysis of 55 randomised cases between January 2016 and August 2017.

- Inclusion criteria: 1) deceased between the time period; 2) under medical and/or surgical care
- Exclusion criteria: 1) illegible DALs*; 2) case referred to Coroner

*MCCDs are kept within the National Registry. Therefore, DALs were used as a surrogate.

**Analysis of Patient Notes**
Paper and electronic notes were scrutinised.

- Non-adherence to National (ONS) Guidelines was determined by the use of 1) abbreviations; and 2) vague terms
- Non-compliance with recommended hospital practice was determined by the absence of a 'Senior Discussion Stamp'.

**Results:** 45/55 cases eligible based on exclusion criteria. In line with ONS Guidelines, 22/45 (48% surgical vs. 50% medical) entries were non-adherent, with 23/45 (52% surgical vs. 50% medical) recorded as adherent. Of the 22 non-adherent entries, 82% were due to the use of vague terms and 18% as a result of abbreviation use. Interestingly, no entries contained the Senior Discussion Stamp.

**Discussion:** Despite the medico-legal implications of inaccurate patient documentation, the quality of MCCD/DAL completion was poor. These results highlight the need for targeted education amongst medical professionals to improve this process.

**Conclusion:** In light of our findings we have developed an educational resource for implementation within the hospital. In addition, emphasis on the documentation of discussions between Junior and Senior teams is warranted.

ARE PATIENTS SUFFERING FROM CHRONIC KIDNEY DISEASE (CKD) STAGE 3 MONITORED REGULARLY IN PRIMARY CARE AS PER THE NICE GUIDELINES?

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**Background:** Worldwide, Chronic Kidney Disease (CKD) is associated with significant mortality and morbidity. Ranked 14th in the leading causes of death in 2012, it is becoming increasingly challenging to manage. Predicting disease progression is difficult in patients, especially those with co-existing comorbidities. Risks associated with accelerated glomerular filtration rate (GFR) decline include cardiovascular and end-stage renal disease (ESRD), the latter proving to be a substantial burden on the NHS. The National Institute for Health and Care Excellence (NICE) and Kidney Disease: Improving Global Outcomes (KDIGO) have produced guidelines on how to approach, monitor and manage these patients.

**Aims:** To assess whether CKD patients have been monitored as per NICE guidance in primary care. To develop a protocol on managing fluctuating GFR parameters for primary care use.

**Methods:** A cohort of CKD patients registered at primary care practice were investigated over a 12-month period; monitoring of GFR/ Urine albumin to creatinine ratio (ACR) and stage 3 of the disease were the prerequisite. Further discussion revealed the need of a local protocol on approaching those with changing GFRs. Relevant literature was reviewed and national recommendations were studied.

**Results:** Forty-nine patients were identified, 44 of which had a GFR tested but only 17 who had combined GFR and ACR monitored. The algorithm produced consisted of managing patients whose decline in GFR was due to either pharmacological reasons or pathological causes.

**Conclusion:** The results indicate the need for regular monitoring with both a blood test and urine dip to adhere to advice given by national health bodies. The aim of the protocol is to detect initial stages of progressive disease and to refer to a specialist for early intervention.
**THE RATE OF SURGICAL SITE INFECTIONS FOLLOWING SPINAL SURGERY: A CLINICAL AUDIT**

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**Background and Aims:** Post-operative surgical site infections are associated with post-operative morbidity and present a significant burden to the NHS in relation to readmission rates, extended inpatient stay and poor outcomes. Smith et al. established the overall rate of surgical site infections following 108,419 spinal procedures as 2.1% the accepted gold standard. This audit aims to compare the rate of post-operative surgical site infections with this expected rate, and to assess whether there is any correlation between rate of infection and type of procedure.

**Methods:** A retrospective search was conducted of all cervical and lumbar spine surgeries requiring a general anaesthetic between 3rd January 2017 and 20th November 2017, totalling 188 procedures. All patients were nursed on a spinal unit throughout admission. Electronic hospital records were reviewed for any evidence of treatment or readmission for infection within six weeks of surgery.

**Results:** Of 188 cases, four were re-operations and 34 were joint consultant cases. No surgical site infections were identified during admission. However, three were identified following discharge, two of which required readmission. This gave an overall infection rate of 1.06%. There were two systemic infections during initial admission (one UTI and one LRTI). There was no correlation between the type of surgery and the rate of infection.

**Discussion:** Although surgical site infections conformed to standards, introduction of GIRFT ring fencing of orthopaedic elective beds has the potential to further improve this. Educating nursing staff regarding the signs of wound infections and appropriate management would help to identify infections promptly and reduce complications.

**Conclusion:** Post-operative surgical site infection rates conformed to and exceeded the expected standards. This may be related to the management of patients on a solely spinal unit under the care of experienced orthopaedic nurses. There was no increase in infection rate associated with prosthesis insertion, suggesting good operative sterility.

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**LONG-TERM RESULTS OF THE ONCOLOGICAL SAFETY OF THE THERAPEUTIC MAMMOPLASTY: RESULTS IN A SINGLE CENTRE**

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**Background and Aims:** Therapeutic Mammoplasty (TM) is a combination of breast-conserving surgery and oncoplastic techniques. It allows the conservation of the natural shape of the breast whilst removing a larger amount of tissue, thus achieving an oncologically safe excision. There is little data available on the long-term oncological follow-up. The primary aim of this study is to assess local recurrence. The secondary aim is to determine 30-day readmission rate, wound infection rate and mortality rate.

**Methods:** A retrospective database of all patients undergoing TM at a single unit by three oncoplastic breast surgeons was created. Data on clear surgical margins, re-excision rate and loco-regional recurrence, surgical technique, complications, need for reoperation, and post-operative histopathology was recorded.

**Results:** A total of 49 patients were identified. The mean age was 52 (range = 29-74) and the mean BMI was 29.3. Ten percent of patients were current smokers. Two patients underwent neo-adjuvant chemotherapy. The study shows that the incidence of loco-regional recurrence is 6.1% at a mean of five years, ranging up to 84 months. A total of 4.1% had distant recurrence. This was within six to 18 months post-operatively. Overall, the incidence of incomplete incision was 4.1% and 8.1% of patients encountered a complication. Of these, 6.1% (3/49) were re-admitted for infected haematoma or wound dehiscence.

**Conclusion:** Our data shows low rates of required re-excisions and recurrence in TM. These results continue to support the long-term oncological safety of TM.
IMPLANT-BASED BREAST RECONSTRUCTION WITH ARTIA TISSUE MATRIX

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Background and Aims: In 2015, Artia™, a new porcine Acellular Dermal Matrix (ADM), was introduced into our breast unit. As clinical studies of its use in breast reconstruction are lacking, the purpose of this prospective study was to assess outcome data for patients who had Artia™ assisted breast reconstruction. We compared this data to other studies of non-human ADMs in widespread use from the literature.

Methods: All consecutive patients who had Artia™ assisted breast reconstruction between July 2016 and February 2018 were identified. A prospective database was maintained including patient demographics, type of reconstruction, type of implant, oncological data if applicable, complication rates, and adjuvant treatment delays.

Results: Fifty-one patients undergoing 83 implant-based breast reconstructions with Artia™ were included in the study. Of the 83 reconstructions, 62% were performed following risk-reducing mastectomy, 28% following therapeutic mastectomy and 10% for revision procedures. The mean age of the patients at the time of surgery was 42.9 years and the mean BMI was 24.7 kg/m². Over a mean 276-day follow-up period, the complication rate was 10.8%, including six breasts (7.2%) developing seromas, one breast developing a haematoma (1.2%) and two implant losses (2.4%) in a single patient.

Discussion: As there is currently no reported experience with this ADM in the literature, we compared our results with those from studies of other non-human ADMs, including Surgimend™ and Strattice™, which remains the most popular ADM in the UK. The complication rate with Artia matched those obtained from larger studies with the well-established ADMs.

Conclusion: This is one of the first studies demonstrating that Artia™ assisted implant-based breast reconstruction is associated with low and acceptable complication rates. The results are promising and are comparable to our experience using established ADMs, with an implant loss rate of 4.9% in over 500 ADM-assisted implant reconstructions.

A SERVICE EVALUATION OF THE IMPLEMENTATION OF CLOSED LOOP OXYGENATION VENTILATION FOR BABIES UNDERGOING NEONATAL TRANSFER

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Background: Oxygen therapy is an important aspect of neonatal care; there is strong consensus that SpO2 levels should be kept within a tight limit. A recent systematic review showed that compliance of oxygen saturation targets is low in neonatal units and, therefore, the potential need for automated ventilation has been proposed.

Aims: The aim of this research project was to perform a service evaluation of the Predictive Intelligent Control of Oxygenation (PRICO) mode in a NICU transfer setting. PRICO mode is an automated ventilation system and the research was carried out with the Greater Manchester Neonatal transport team.

Methods: A standardised data collection form was produced which included a five-minute log for SpO2. This was taken on transfers and data was collected for 20 transfers over the data collection period. In 40% of the transfers, the SpO2 was kept within the desired range 100% of the time. In the rest of the transfers, the percentage of time within the range set was above 50% except for three cases.

Results: The recent BOOST II study found only half of neonates' oxygen saturations were kept in the desired range. When compared to the literature, this research indicates PRICO works well compared to manual adjustments.

Discussion: This study was purely observation and a RCT would need to be designed with a larger sample size to allow for statistical analysis. As with all new technology, PRICO will require surveillance as it is introduced into routine clinical practice.
THE USE OF SILDENAFIL IN PRETERM INFANTS WITH CHRONIC LUNG DISEASE

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Background and Aims: Chronic lung disease (CLD) is a multisystem disorder that affects many preterm infants. It is a significant cause of mortality, developmental delay and family distress. Sildenafil is a selective pulmonary vasodilator which has been shown to have potential in the treatment of neonates with CLD and associated pulmonary hypertension. This study aimed to investigate the use of Sildenafil in preterm infants with CLD.

Methods: This case-control audit looked at infants with CLD who were prescribed Sildenafil at Royal Preston Hospital between 2009 and 2016. It compared these infants to a larger population with CLD who were not prescribed Sildenafil. Data was collected from patient notes and analysed using chi-squared testing, unpaired t-tests and Fisher’s exact test.

Results: The data showed that the babies who went on to receive Sildenafil (n = 14) had significantly lower birthweights by an average of 174g (p = 0.0008) and were more premature by an average of 1.4 weeks (p = 0.0083) than the control group (n = 144). They were more likely to be discharged on steroids and diuretics. They spent longer dependent on invasive ventilation and non-invasive ventilation. They had a significantly higher CLD-related mortality rate (28.6% compared to 1.4% in the control group) and were less likely to be discharged home. Eight babies’ level of respiratory support decreased after an average of 23 days Sildenafil therapy. There were no statistically significant differences in maternal risk factors or method of delivery between the two groups.

Discussion: This data demonstrates that babies who receive Sildenafil are smaller, more premature, and need more intensive respiratory support. Improvement in over half of the cases who received the drug may inform prescribing practices.

Conclusion: Babies who receive Sildenafil are a unique subgroup of the CLD population, at the severe end of the disease spectrum. It should add to existing literature to consider Sildenafil as a useful agent when treating severe CLD.

SURGICAL MANAGEMENT OF HIGH GRADE GLIOMAS IN THE ELDERLY POPULATION

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Background: As the population ages, the number of individuals diagnosed with gliomas is predicted to increase. Surgical resection of gliomas followed by adjuvant therapy based on evidence available could contribute to increase overall survival in the elderly population.

Aims: A retrospective study to assess the survival rates in elderly patients with high-grade gliomas against the survival and complication rates found in literature. The secondary objective is to identify whether a significant difference in survival exists between the <75 and ≥75 age groups.

Methods: Patients surgically treated for histologically proven gliomas at the Neurosurgical Unit at Salford Royal Foundation Trust from 1st June 2011 to 3rd March 2016 were included in this study and data was obtained from the Neuro-Oncology Database and Electronic Patient Records.

Results: A total of 124 patients older than 65 years with 131 admissions were included. There were 46 males and 78 females. Gross total resection was performed in 25 patients (19%); subtotal resection in 69 patients (53%); partial resection in 12 patients (9%); and biopsy in 25 patients (19%). At last follow-up, seven patients were alive with a median overall survival of 6.4 months. In log-rank analysis, the median survival for <75 was 6.9 months and ≥75 was 5.2 months, with no statistically-significant difference between the two groups. Complication rates amongst subgroups were not different.

Conclusion: In selected patients, surgery can be considered as a safe option. If surgery is accurately selected, patients above the age of 75 can benefit from surgical intervention with acceptable morbidity.
IMPACT OF ADMITTING SPECIALTY ON EMERGENCY LAPAROTOMY MORTALITY

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Background: Surgical patients may be directly admitted to general surgery or non-general surgery specialties, and their operation may be carried out at different time points. We hypothesise that surgeons are more proficient at diagnosing surgical problems, thereby reducing time to theatre and subsequently reducing mortality rate.

Aims: To establish whether patient admission through general surgery vs. non-general surgery affects the mortality of patients undergoing emergency laparotomy.

Methods: Retrospective analysis of patients undergoing emergency laparotomy between January 2009 and December 2014. Emergency theatre logbooks were used to identify those who underwent emergency laparotomies. Admission dates, admitting specialty, discharge dates, date of death, and postcodes were recorded using The Open Patient Administration System (TOPAS). Exclusion criteria included: emergency department admissions; gynaecological patients; and vascular patients. Fisher’s exact test was used to identify statistical significance. End points were 30-day, 90-day, one-year and three-year mortality.

Results: We included 578 laparotomies in total. Mean patient age: 63 for both surgical and non-surgical admissions. A total of 50.3% of patients were male and 49.7% were female. The overall mortality at 30 days, 90 days, one year and three years were 13.1%, 16.8%, 23.5% and 32.4%, respectively. Data comparing surgical vs. non-surgical cohorts is displayed in Table 1. P-values were 0.0279, 0.0472, 0.0088 and 0.0046 for 30 days, 90 days, one year and three years mortality, respectively.

Table 1: 30-day, 90-day, one-year, and three-year mortality in surgical vs. non-surgical cohorts.

<table>
<thead>
<tr>
<th>Mortality</th>
<th>General Surgery</th>
<th>Non-General Surgery</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>30 days</td>
<td>11.6%</td>
<td>19.8%</td>
<td>0.0279</td>
</tr>
<tr>
<td>90 days</td>
<td>15.2%</td>
<td>23.4%</td>
<td>0.0472</td>
</tr>
<tr>
<td>1 year</td>
<td>21.2%</td>
<td>33.3%</td>
<td>0.0088</td>
</tr>
<tr>
<td>3 years</td>
<td>29.6%</td>
<td>44.1%</td>
<td>0.0046</td>
</tr>
</tbody>
</table>

Conclusion: Patients admitted directly to general surgery and underwent emergency laparotomy had a much lower mortality rate than those admitted through non-surgical specialties. However, better recognition of the need for laparotomy by a general surgeon is one component of a multi-varied equation that predicts overall mortality and concluding that it is the only reason would be an over simplification.

POSTNATAL CARE FOR GESTATIONAL DIABETES MELLITUS: AN AUDIT OF A LONDON-BASED GP PRACTICE

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Background and Aims: Gestational Diabetes Mellitus (GDM) puts women at a high risk of developing Diabetes Mellitus II (DMII). DMII poses a significant disease burden to patients, as well as representing a substantial financial burden on the NHS, approximately £12 billion annually. It is important to identify patients at risk of DMII to enable monitoring and early treatment, and to prevent comorbidities. NICE recommends that every woman who has GDM should receive an annual HbA1c as part of their follow-up care. The aim of this study was to evaluate if a particular GP practice complied with the above recommendations.

Methods: An audit was carried out at the local GP practice and the service was compared to the NICE recommendations. A search on the practice database was performed to find all the patients who have had GDM. The study evaluated if they all had their HbA1c checked within the last year.

Results: It was found that 73% of patients with a history of GDM had their HbA1c checked within the last year. The main reasons identified for not meeting the NICE standards were that some patients were not being recalled to have their HbA1c checked. Other patients had been recalled but did not have the blood tests. New patients who moved into the area were also missed as recalls were not entered into the system for them.

Discussion: As a result of the audit, all the patients identified were put into the system for an annual recall. Additionally, each patient was contacted by telephone to explain the importance of annual HbA1c testing and the requests forms were printed for them to collect at the practice. This audit was presented to the doctors working in the practice to increase awareness. The next step is to re-audit the practice in a year’s time to assess if there has been any long-term improvement.
THE USE OF ECG IN THE ASSESSMENT OF CHILDREN PRESENTING WITH CONVULSIVE SEIZURES: A REVIEW OF LOCAL PRACTICE AT THE ROYAL VICTORIA INFIRMIARY, NEWCASTLE-UPON-TYNE

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Background and Aims: The British Paediatric Neurology Association (BPNA) teaching supports Scottish Intercollegiate Guidelines Network (SIGN) recommendations that every child presenting with a febrile convulsive seizure have a 12-lead ECG performed with corrected QT interval (QTc) calculated. We aimed to review our clinical practice regarding adherence to this.

Methods: All children aged 16 or younger attending A&E or the admissions unit at the Royal Victoria Infirmary (RVI) from 01/12/17 to 31/01/18 with admission diagnosis coded as ‘seizure’ were identified (n=101). Electronic and paper notes were reviewed; those with febrile or non-convulsive seizures and long-term epileptic patients for whom first seizure information was not available were excluded.

Results: Twenty-eight patients were included. Twenty-one patients (75%) had an ECG performed, with 81% of these having a normal ECG with normal QTc. Four patients had an ECG but no documentation of interpretation or QTc calculation.

Discussion: Although we demonstrate high concordance with SIGN guidelines, 25% of children presenting with an afebrile convulsive seizure had no ECG in their medical records, and a further four had no documentation of ECG interpretation or QTc calculation. A similar audit in Nottingham in 2014 highlighted only 23% adherence to SIGN guidelines, suggesting greater clarity around clinical guidelines and further education may be required to improve concordance nationally.

Conclusion: This audit has highlighted high adherence to SIGN recommendations at the RVI. However, a significant minority of patients were not managed accordingly. Improving this is of utmost importance as, whilst long QT syndrome (LQTS) is uncommon, reflected by the absence of any prolonged QTc in this audit, children with LQTS are prone to life-threatening ventricular arrhythmias. These can manifest clinically as syncope and seizures, and mortality can be reduced significantly with appropriate treatment.

BLOOD LOSS IN NECK OF FEMUR FRACTURES - IS THERE A ROLE FOR TRANEXAMIC ACID?

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Background: In hip fractures, a majority of in-hospital mortalities have been linked to postoperative blood loss. Studies have also shown that intra-operative blood loss is often significant, and hidden blood loss in hip fracture patients may be as high as 1500cc. Additionally, a study has demonstrated that blood transfusion increases in cost per admission by about £1200.

Aims: To quantify and compare the blood loss in patients with neck of femur fractures with and without administration of tranexamic acid.

Methods: We collected data for 100 consecutive NOF fractures admitted before July 2017. The ongoing re-audit is a data of 50 consecutive patients with NOF fractures who have received tranexamic acid. All statistical tests were performed using SPSS. Frequencies were used to analyse the demographic data; student t-test and Mann Whitney U test was used for continuous variables.

Results: Mean estimated blood loss was 714 ml. Our study showed no significant increase in blood loss in patients on anticoagulants (p-value = 0.3). Blood loss was significantly higher in patients who had hemiarthroplasty (p 0.04), DHS (p 0.5), IMHS (p 0.7) and THR (p 0.2) did not cause significant blood loss. The data is being collected and analysed for a cohort of patients who have received tranexamic acid.

Conclusion: Neck of femur fracture patients lose close to two units of blood. Patients having hemiarthroplasty are susceptible to blood loss. Hence, these patients may benefit from administration of tranexamic acid.
AUDIT ON PELVIC FRACTURES AND TIME TO RECONSTRUCTION

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Background and Aims: Fractures of the pelvic ring occur following high energy transfer, and are significant injuries with associated morbidity and mortality. These injuries require urgent management with a pelvic binder, followed by either early definitive surgery or temporary mechanical stabilisation if this is unable to be performed. According to the new BOAST (British Orthopaedic Association Audit Standards for Trauma) guidelines on pelvic fractures, pelvic ring reconstruction should occur within 72 hours following stabilisation of a patient’s physiological state. This aim of this study was to evaluate the time to reconstruction in patients presenting with pelvic fractures to the Royal Stoke University Hospital.

Methods: From a trauma admissions database, we retrospectively reviewed records of 14 patients with pelvic fractures who required operative management from 01/01/2017 to 31/12/2017. Patients were deemed to have achieved a stable physiological state once they had achieved a stable serum lactate result, or once they were stable enough to undergo trauma computed tomography (CT) scan.

Results: Of the 14 patients who fitted the inclusion criteria, 13 patients (93%) underwent pelvic ring reconstruction within 72 hours of achieving a stable physiological state. Eight of the 14 patients (57%) had undergone their operation within 24 hours of stabilisation, while 10 of the 14 patients (71%) had their operation within 48 hours of stabilisation.

Discussion: A large proportion of patients who presented with pelvic fractures requiring reconstruction to the Royal Stoke University Hospital (Major Trauma Centre) were able to undergo pelvic ring reconstruction within the target time, as all but one (93%) of our patients had met the target time for pelvic ring reconstruction as per the new BOAST pelvic fracture guidelines introduced in January 2018. This was achieved even prior to introduction of the new guidelines.

Conclusion: Ninety-three percent of our patients met the target time as set by current BOAST guidelines. Following dissemination of the results via presentation at a department-wide meeting, and circulation of the new guidelines, we will aim to re-audit this following application of these interventions.

CONSENT FOR COMMON GENERAL SURGICAL PROCEDURES: A RETROSPECTIVE AUDIT

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Background and Aims: Consent is essential to patient safety. It is covered by the Royal College of Surgeons’ ‘Good Surgical Practice’ guidelines, and has gained attention recently following the Montgomery vs. Lanarkshire case. However, consent taken by doctors is anecdotally highly variable. None of the Surgical Colleges or relevant societies offer procedure-specific consenting guidance. EIDO is a private company which supplies evidence-based patient information leaflets to hospitals for consent, accredited by the Royal College of Surgeons. This study examines the procurement of consent for common general surgical procedures in a district general hospital.

Methods: The consent forms for 135 common procedures (50 open inguinal hernia repairs, 39 laparoscopic cholecystectomies, and 46 diagnostic laparoscopies) performed in the preceding six months at a district general hospital were audited. Data was collected on the time between consent and surgery, and whether patients were given a copy of the consent form. Risks were compared with those listed in the EIDO leaflet for each procedure.

Results: For each procedure, common risks were well-represented on consent forms, with more serious risks such as bile leak in laparoscopic cholecystectomy (72%) and damage to the spermatic cord in hernia (60%) less frequently recorded. Consultants listed fewer risks than junior doctors. No single consent form recorded all pertinent risks as recommended by EIDO. In 42% of forms, the patient copy was retained in the case notes and not given to the patient. Sixty-seven percent of consent forms were completed on the day of surgery.

Conclusion: There was a wide variation between individual clinicians in risks mentioned. Although recent guidance states that consent should be obtained prior to the day of surgery, this is not the case for most patients in this cohort. This data was presented locally at the departmental meeting, and a plan was initiated to provide pre-printed consent forms and patient leaflets in the clinic.
TRUST OPPORTUNITIES IN PLASTIC SURGERY TRAINING

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Background and Aims: Core Surgical Trainees (CST) undertake a two-year training programme in the UK which provides training in a range of surgical specialties. The Joint Committee for Surgical Training (JCST) states that all CSTs should attend three consultant-supervised sessions of four hours each week (40% of our total working time) and one clinic (8% of working time). Increasing service provision demands due to rotas gaps has led to training opportunities being compromised, affecting not only training but also moral and job satisfaction. We aim to review and improve this through our audit.

Methods: We carried out a retrospective study of the training opportunities available to the Senior House Officers at Queen Elizabeth Medical Centre's Plastic Surgery Department by documenting service provision and training opportunities from October 2017 to December 2017. Data was collected via weekly rotas and recorded on a spreadsheet. Our findings were presented to the junior doctors' forum to highlight the discrepancies in our training opportunities. A further audit was performed for the following two months using the same database, and the data was analysed.

Results: Our original loop over the initial four-month period revealed that 36.34% of our working time was dedicated to theatre and 36.8% of our working time was allocated to clinic. Following intervention, our theatre time was 43.7% and clinic time was 6.0%.

Conclusion: We have demonstrated that our current training model within this hospital is below the standard set by JCST. However, following discussion at our doctors' forum, we have managed to increase our allocated theatre time to above the recommended standard of 40%. In the current climate, training often takes a back seat as the demand for service increases ever more. We must recognise that training is important for career progression, and we have highlighted that, through increased awareness, the balance can be readdressed.

IMPROVING OUTCOMES OF EMERGENCY COLECTOMY IN PATIENTS WITH ACUTE SEVERE COLITIS: ENHANCED SURGICAL REFERRAL PATHWAY

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Background and Aims: Acute severe colitis is a life-threatening condition which should be recognised early and managed with prompt initiation of intensive treatment. A delay in surgery due to “hopeful-expectations” from medical management may lead to increased post-operative complications. This study aims to develop an enhanced surgical referral pathway which will guide the management of patients with acute severe colitis.

Methods: A retrospective review of patients diagnosed with acute severe colitis between October 2015 and September 2017 was conducted. Patients who underwent emergency subtotal colectomy were identified. Data was cross-referenced between emergency theatre and inpatient databases; duration of medical therapy, time to surgical referral from diagnosis, and post-operative outcomes were analysed. Current surgical referral pathway was audited to evaluate inpatient care against NICE guidance. Patients were classified in two cohorts based on duration of in-hospital medical therapy and time to surgical referral. Univariate analysis was used to assess the impact on post-operative outcomes, and Student's t-test was used to quantify significance.

Results: Twenty-seven patients identified with acute severe colitis; 21 were treated operatively. All received first-line intravenous corticosteroids. Fourteen patients underwent emergency subtotal colectomy. Mean age was 38 (range 20 – 72); mean duration of first-line therapy was 11.5 days (SD 5.1). Time to surgical referral was 7.1 days (SD 5.2); time to surgery (subtotal colectomy) from surgical consultation was 4.6 days (SD 3.5).

Discussion: Medical therapy >10 days was associated with 50% of post-operative complications (OR = 5.33 (p = 0.21)). Average time to surgery was 16 days (SD 3.2); colectomy risk RR = 6 (p = 0.11). Those in the <10 days of medical treatment cohort had reduced time to surgery of 7.1 days (SD 1.77) with no complications. Time to surgery for referrals >7 days from diagnosis was 12 days (SD 2.7). A total of 38% suffered complications (OR = 5.00 (p = 0.238)), increasing risk (RR = 4 (p = 0.1744)). Surgical referral <7 days improved time to surgery to 3.5 days (SD 2.0), with one 24-hour ITU admission. Collectively prolonged medical therapy and delayed referral comprised 88% of all complications, increasing colectomy risk (RR = 8.07 (p = 0.174)). A paired t-test (p < 0.0001) quantified our results as significant.

Conclusion: Duration of in-hospital medical management of acute severe colitis is associated with an increased colectomy risk, with delay in surgical input worsening this. We developed an enhanced pathway structured on NICE guidance and Oxford Criteria to gauge the severity in those with acute colitis who may require colectomy at the time of admission, allowing early surgical involvement to guide therapeutic management in a multidisciplinary approach.
MAGNETIC RESONANCE IMAGING (MRI) OF THE INTERNAL AUDITORY MEATUS (IAM): AN AUDIT OF LOCALLY SUBMITTED REQUESTS AND FINDINGS

Borg D, Gatt AS
Mater Dei Hospital, Malta

Background and Aims: Early detection of acoustic neuroma (AN) translates into timely hearing-preserving surgery and decreased perioperative morbidity. MRI is the gold standard investigation for visualising the cerebellopontine angle (CPA). Being an expensive resource that is in high demand, we sought to identify ways of fine-tuning MR IAM referrals.

Methods: A retrospective analysis of 690 MR IAMs ordered between January and December 2016 at Malta’s acute general teaching hospital, Mater Dei, was undertaken. Follow-up scans, inadequate and cancelled studies were excluded. Unique first-ever MR IAMs totalled 617. The Radiological Information System (RIS) served as our source of data collection. Patient demographics, referring specialties, inputted reason for examination, and imaging findings were recorded.

Results: The commonest symptomatology leading to investigation with an MR IAM was vertigo/dizziness. Eighty-one percent of MR IAM requests were made by otolaryngologists, while the remaining 19% came from all other specialties. Four of 617 (0.65%) MRIs were positive for AN. All four patients had asymmetric sensorineural hearing loss (SNHL). Two of four diagnosed patients were referred from audiovestibular medicine. Asymmetric SNHL was also the commonest non-unanimous complaint among original scans traced back from 2016 follow-up MRIs. Six of 617 (0.97%) patients had significant disease other than AN. Of the scans which were negative for AN/significant pathology, 50 of 607 (8.24%) had incidental findings which we will describe.

Discussion: No globally recognised guidelines for imaging patients with asymmetric SNHL exist. However, various centres in the UK have published regional protocols which describe frequency-specific criteria for targeted screening. Precise request documentation of SNHL asymmetry in decibels (dB) would help ascertain how Maltese patients fit into foreign protocols.

Conclusion: Better patient selection through inter-specialty consultation and prior audiovestibular testing may reduce any inappropriate burden on radiology services and the medical financing system, whilst simultaneously safeguarding patients.

SEPTIC ARTHRITIS MANAGEMENT IN A DISTRICT GENERAL HOSPITAL

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Royal Gwent Hospital, Newport, United Kingdom

Background and Aims: Septic arthritis carries significant morbidity and mortality. Prompt diagnosis and treatment is vital for patient outcome. The BOA recommends joint aspiration prior to commencing antibiotics. NICE advocates initiating the “sepsis six bundle” for all septic patients within one hour of presentation. This study aims to investigate the management of septic arthritis in a large district general hospital.

Methods: We retrospectively reviewed patients with septic arthritis who were treated surgically over a one-year period. Penetrating injuries, chronic infections, medically treated patients and paediatric patients were excluded. Compliance with the BOA guidance and “sepsis six bundle” was recorded. Time from referral to aspiration, administration of antibiotics, and surgery were all recorded.

Results: Twenty-two patients were identified (native n=15; prosthetic n=7). The knee was the most common joint (16). There were four deaths, all in prosthetic joint infections. Antibiotics were given before aspiration in five of 15 native joint and three of seven prosthetic joint infections. Nineteen of 22 patients were clinically septic on presentation, only 10 of 22 patients had blood cultures taken, and none received antibiotics within one hour.

Patients who survived waited, on average, 19 hours for antibiotics and 29 hours for surgery, compared to 54 hours for antibiotics and 170 hours for surgery in those who died.

Discussion: A limited number of studies consider the relationship between time delay in initiating antibiotics and mortality in septic arthritis cases presenting with sepsis. Prosthetic joints may receive longer delays to antibiotics as they require sterile conditions in theatre for joint aspiration prior to commencing antibiotics. This study demonstrated that delays in treatment in both prosthetic and native joints may cause higher mortality, prompting early surgical management for all septic arthritis cases.

Conclusion: We demonstrated universally poor compliance with guidelines. Mortality and outcomes were worse with prosthetic joint infections and delays in treatment. We have instigated measures to improve care, aiming to reduce time to aspiration, antibiotic administration, and surgical washout.
NATIONAL HANDOVER PRACTICES IN 2018: CONCORDANCE WITH GUIDELINES AND THE IMPACT OF THE NEW JUNIOR DOCTOR CONTRACT

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Background and Aims: The new junior doctor contract, implemented in phases from October 2016 to April 2017, forced Trusts to ensure compliance of working rotas with the new contract, and may have indirectly prompted a revision of handover practices as a result of re-designed working patterns. In this national, telephone-survey audit, we aim to assess the concordance of handovers across different hospitals in England with handover guidelines from the BMA and NPSA. We endeavour to compare the outcomes of the survey between data collected before and after contract implementation.

Methods: A 20-point questionnaire was delivered via a telephone conversation to on-call General Surgery FY1 doctors in over 100 hospitals between November 2017 and February 2018. Equivalent data had previously been collected using identical survey methods in early 2016. Permutation tests were employed in order to determine if there had been a significant change in the response to a particular question over time.

Results: The average time (minutes) allocated for handovers increased from 24.10 in 2016 to 30.69 in 2018. Handover time was protected in more cases in 2018 (83% c.f. 65%), it was less likely to be interrupted in 16% of hospitals (37% c.f. 21% in 2018), and it occurred within working hours 11% more than previously recorded (93% c.f. 82%). We also found that ‘time used’ was correlated with satisfaction in both years, and ‘time allocated’ strongly correlated with satisfaction in 2018 (spearman coefficient = 0.45, p = 0.0008).

Discussion: Improvements in handover practices are evident in comparing data collected in 2016 with recent data from 2018, including longer handover time, less interruptions, scheduled handover time within working hours, and improved junior doctor satisfaction. However these improvements cannot be attributed to new contract changes alone, and must be considered in the context of continued handover improvements over a longer period of time.

DOES THE IMPLEMENTATION OF A WARD ROUND PRO-FORMA IN A MEDICAL WARD IMPROVE ADHERENCE TO BEST PRACTICE GUIDELINES?

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Background and Aims: Robust clinical record keeping has been shown to facilitate the delivery of excellent patient care. Documentation of medical ward rounds, in particular, is critical in effectively communicating essential information regarding patient progress and management decisions to members of the Multi-Disciplinary Team. The aim of this study was to evaluate the effectiveness of a ward round pro-forma in a medical ward to ascertain if it adhered to the best practice guidelines.

Methods: We first assessed adherence to best practice guidelines of every ward round note in a medical ward over two weeks. We then designed and introduced a ward round pro-forma based on these guidelines, and subsequently audited those notes that were recorded using the pro-forma. Rates of documentation were examined for twenty variables that were grouped into five categories: 1) patient demographics and background; 2) subjective findings; 3) objective findings; 4) assessment and plan; and 5) doctor details.

Results: Initial spot auditing of 123 ward round notes revealed poor compliance in all categories, with an overall compliance rate of 54.9%. We then implemented our specifically designed pro-forma in 96 ward round notes over the subsequent two weeks. We detected a statistically significant improvement (p < 0.05) in documentation standards with an overall increased compliance of 40.7% and in each category as follows: category one 45.6% category two 36.6% category three 36.3% category four 16.2% and category five 46.9%

Conclusion: Our findings support the more widespread use of a standardised pro-forma to document medical ward round notes in line with best practice guidelines.
ACUTE MASSIVE HAEMOPTYSIS AND RESPIRATORY DISTRESS: A RARE INITIAL PRESENTATION OF METASTATIC TESTICULAR CANCER

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Background: Testicular cancer is the most common malignancy in men aged between 20 and 35 years. Testicular germ cell tumours make up 95% of testicular tumours and are the commonest solid malignancies in young men. Risk factors for testicular cancer include cryptorchidism, family history, infertility, tobacco use and Caucasian. Testicular mass is the commonest initial presentation. However, patients with advanced metastatic disease may present with various other symptoms. We report a rare case whereby the initial presentation of a metastatic testicular cancer was acute massive haemoptysis and marked respiratory distress.

Case Study: A 38-year-old male patient presented to an emergency department with acute massive haemoptysis and marked respiratory distress. After being stabilised by resuscitative measures, a chest X-ray was taken and showed widespread cannonball metastasis. Tumour markers were elevated; HCG 1539 IU/L (0-10IU/L) and AFP 132 kU/L (0.0-8.0 kU/L). CT scans reported multiple well-circumscribed lesions throughout the lungs. Head CT scan showed a solitary 3.5cm metastatic deposit within the left parieto-occipital region. Ultrasound scan confirmed a right testicular mass. Orchidectomy was followed and histological report confirmed mixed germ cell, 90% classical seminoma and 10% embryonal carcinoma. The patient remained disease free after one year post chemotherapy. All serum tumour markers were normal.

Discussion: From a literature review, only 11 cases were reported whereby haemoptysis was the first initial presentation of a metastatic testicular cancer, usually choriocarcinoma. Primary lung cancer is often thought to be the main differential diagnosis. It is, however, essential to differentiate between primary lung tumour and metastatic lung disease as there is a significant difference in management and prognosis.

Conclusion: Physicians need to be cautious when dealing with male patients presenting with haemoptysis as this may be the first presentation of testicular cancer. Early diagnosis and therapy have very good cancer survival rate.

IS THE PREVALENCE OF ANAEMIA IN PATIENTS IN ATRIAL FIBRILLATION HIGHER IN COMPARISON TO PATIENTS IN SINUS RHYTHM, A CASE-CONTROL STUDY

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Background: Atrial fibrillation has a high impact on morbidity and mortality in the population. Management and prevention of atrial fibrillation requires treatment of potential reversible trigger factors. Anaemia has been observed in patients with atrial fibrillation but no data is available as to date on the prevalence of anaemia in atrial fibrillation.

Aims: The main objective is to establish if a relationship exists between anaemia as a risk factor and atrial fibrillation as an outcome factor depending on the prevalence of anaemia. The secondary outcome is a review of other potential contributing risk factors.

Methods: Retrospective, case-controlled study investigating the prevalence of anaemia in patients with atrial fibrillation, assigned as cases, compared with patients in sinus rhythm, assigned as controls. Four hundred and five patients with atrial fibrillation and 399 patients in sinus rhythm who presented to the Emergency Departments at Imperial College Hospitals were identified. They were assessed with an electrocardiogram and had a blood test which included a full blood count sample. Data was analysed with a statistical program, State. The anaemia rate in both groups was compared with logistic regression test and categorical variables were compared using the Chi-square test.

Results: The study found no difference in the prevalence of chronic anaemia between both rhythm groups after adjustment for acute anaemia. However, gender specific differences in risk factor prevalence were found and hypertension was observed to be a contributing factor overall in the development of atrial fibrillation and anaemia.

Conclusion: Anaemia was not found to be an independent risk factor of atrial fibrillation. The overzealous treatment of anaemia bears high risks of transfusion transmitted diseases and reactions, and would not contribute to the prevention of atrial fibrillation.

No treatment of anaemia is required beyond that set out in current guidelines in patients with atrial fibrillation.
ASSESSING STROKE VOLUME VARIATION AS A PREDICTOR OF FLUID RESPONSIVENESS AFTER A FLUID CHALLENGE: A PROSPECTIVE STUDY AND SERVICE EVALUATION

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Background and Aims: A change in stroke volume has for a long time been used as a measure of fluid responsiveness after a fluid challenge. The gold standard measurement for stroke volume is currently pulmonary arterial catheter (PAC) based thermal dilution, but over the years there has been a trend for less invasive monitoring. The Deltex Combi Oesophageal Doppler Machine is one such device that measures cardiac variables including stroke volume variation. This ongoing service analysis aims to look at whether SVV is correlated to stroke volume and can therefore act as a predictor of fluid responsiveness.

Methods: Eleven patients whose treatment required the use of the Deltex Combi ODM and PAC were identified. All variables were recorded continuously, five minutes before a fluid challenge and for fifteen minutes post-challenge. Each recorded variable was then correlated with the recorded stroke volume and with the other recorded cardiac variables. The mean average both pre- and post-fluid challenge was also recorded and the difference was analysed. Finally, the diagnostic odds ratio was recorded for the SVV.

Results: The correlations for each of the variables with stroke volume are: SVV = 0.823, PPV = 0.841, FTc = 0.62. The correlation of SVV vs. PPV = 0.88, PPV vs. FTc = -0.453, and SVV vs. FTc = 0.487. The mean differences pre- and post-challenge are: SV = 6.93%, SVV = 36.57%, PPV = 50.58%, FTc = 7.93%. The diagnostic odds ratio for SVV is 4.

Conclusion: Recording stroke volume with a pulmonary artery catheter as a gold standard measure of fluid responsiveness, a stroke volume variation of >10% showed a strong correlation coefficient of 0.823, a diagnostic odds ratio of 4, and, consequently, could be used as an accurate measure of stroke volume. Flow time corrected has no correlation with stroke volume and cannot be individually used to predict fluid responsiveness.

PERITONITIS SECONDARY TO FOREIGN BODY PERFORATION OF RECTUM IN THE PRESENCE OF ACUTE PARANOID PSYCHOSIS: A CHALLENGING PRESENTATION

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Background: The insertion of rectal foreign body to cause physical injury without suicidal intent is a recognised but poorly understood presentation of Deliberate Self Harm (DSH) in patients with mental health conditions. We report a case of peritonitis secondary to foreign body perforation of rectum in the presence of acute paranoid psychosis.

Case Study: A 43-year-old gentleman suffering from paranoid schizophrenia presented with foreign body insertion into his rectum, resulting in rectosigmoid perforation and peritonitis. No further history was obtained. He had historical presentations of DSH, including two episodes in the months prior. This continued behaviour had resulted in detainment under the Mental Health Act. On examination, the patient was septic and had generalised peritonitis. His inflammatory markers were markedly raised. Digital rectal examination and cross-sectional imaging confirmed the presence of a perforating 20cm x 3cm cylindrical object at the rectosigmoid junction abutting the abdominal wall. He underwent a laparotomy and Hartmann's procedure with retrieval of the foreign object.

Discussion: A comprehensive search of Medline and EMBASE databases confirmed the paucity of evidential study exploring RFB in the context of mental ill-health. Unruh et al. suggested that management of patients who present with RFB insertion should involve exploring the aetiology to prevent future occurrences. Patients with mental health difficulties require a strong support system (e.g. family and friends, community psychiatric nurses, care coordinators, and psychiatrists), willingness to engage with treatment, and abstaining from substance abuse, amongst others.

Conclusion: Attitudes and behaviours of admitting clinicians have historically been blamed for the lack of divulging the intentions underlying these presentations. It is therefore proposed that patients with RFB should ideally have a psychiatric evaluation alongside their surgical management to prevent future occurrences.
HIGH NUMBER OF BED DAYS SAVED WITH OUTPATIENT PARENTERAL ANTIBIOTIC THERAPY (OPAT) IN MANAGING MALIGNANT OTITIS EXTERNA IN A DISTRICT GENERAL HOSPITAL

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Background and Aims: Malignant Otitis Externa (MOE) is a condition associated with high morbidity and mortality, requiring prolonged parenteral antibiotic therapy where duration can extend to months. Our objective is to determine cost savings in treating MOE via the newly established Outpatient Parenteral Antibiotic Therapy (OPAT) service at Ipswich Hospital NHS Trust.

Methods: Retrospective data was collected of all ENT referrals to OPAT services from November 2016 to April 2018.

Results: Of 14 ENT referrals, 10 were accepted for OPAT; of which 9/10 were MOE (7/9 treated for Pseudomonas alone, 1/9 treated for Pseudomonas and methicillin-resistant Staphylococcus aureus, and 1/9 treated for Pseudomonas and anaerobes). All of the MOE patients were elderly; 7/9 were males (mean age: 83.3; range: 73-93) and 2/9 were females (97, 94). Under OPAT, 8/9 MOE patients had successful outcomes (discharge from ENT follow-up, no treatment failures/relapses). However, 1/9 had unexplained death following OPAT to oral switch. The aggregated duration of MOE treatment was 558 days (mean: 62.0; range: 30-130). The aggregated OPAT duration was 448 days (mean: 49.8; range: 14-118) accounting for 80.3% (448/558) of the total treatment days. The remaining 1/10 treated under OPAT had post-mastoidectomy infection (ESBL E. coli).

Discussion: Ipswich Hospital OPAT services enabled nine MOE patients to have, on average, 80.3% of their total treatment from home, with an estimated average saving of £8,711.11 per patient (calculated at £175 per day). Overall, there was a total estimated saving of £78,400 for all patients throughout the 17-month period.

Conclusion: MOE remains a challenging condition to manage with high associated morbidity and mortality, requiring prolonged antimicrobial treatment. OPAT services significantly shortened inpatient hospital stay which resulted in considerable cost savings to the Trust and outcomes comparable to other case series. Patients with MOE can be effectively managed in the community through local OPAT services.

WHAT IS THE PREVALENCE AND CONTROL OF THE COMMON RISK FACTORS FOR TYPE 2 DIABETES MELLITUS (T2DM) IN GIZO HOSPITAL, SOLOMON ISLANDS (SI)?

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Background and Aims: Type 2 Diabetes Mellitus (T2DM) has reached pandemic proportions and is rapidly increasing the Solomon Islands are no exception. The effect and burden of T2DM is profound, posing a serious public health challenge, and action is needed. Research is lacking in this area in the Solomon Islands and this study is the first that aims to explore the prevalence and control of the common risk factors of T2DM. This may help to inform improvements made in both managing and preventing the condition within the community setting.

Methods: World Health Organisation (WHO) STEPS method for non-communicable diseases was used. Six common risk factors were assessed using behavioural, physical and biochemical readings from 32 subjects (53% females; 46% males) in Gizo Hospital throughout May 2017. Results of each risk factor measured per subject were compared to the recommended clinical targets to establish how well T2DM is being controlled.

Results: There was a high prevalence of risk factors, with 78% (n=25) categorised as high-risk from failing to achieve clinical targets for 3-6 risk factors. Smoking was the best controlled risk factor, compared to glycaemic control being the worst (87% and 13% respectively reached the clinical targets).

Discussion: This study provides essential baseline data, but further research and improved lifestyle interventions from both a national and community setting is indicated in order to improve risk factor control and subsequently reduce the burden of T2DM.
FIRST TIME PRESENTATION OF LARYNGEAL HAEMORRHAGE AS A PRESENTING COMPLAINT FOR ACQUIRED HAEMOPHILIA

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Background: This is a case study on a very rare manifestation of acquired factor VIII deficiency. Almost 50% of cases of factor VIII deficiency are idiopathic and challenging to our understanding of its occurrence. Acquired haemophilia has a tendency to cause bleeding in the skin, mucous membranes, soft tissue and muscles without joint involvement. Patients typically present with an acute history of bleeding without being known to be of a high bleeding risk.

Case Study: A 70-year-old gentleman presented with stridor of acute onset after flu-like symptoms for few days prior to admission. He was previously well with no previous stridor, nor did he have any bleeding under his skin, mucous membranes or from his orifices. The stridor was followed by bruising involving the anterior and upper chest, specifically the suprasternal notch. His general physical examination was normal. Naso-endoscopic examination revealed that the stridor was due to an acute haemorrhagic event. An urgent coagulation sample was sent for INR and APTT analysis, and a diagnosis of secondary acquired factor VIII deficiency was confirmed. The patient responded to steroids, mycophenolate, and fresh frozen plasma.

Discussion: The recommendation for similar cases would be to start by measuring the APTT, then conduct mixing studies looking for correction in APTT duration. If correction is achieved, then check factor VIII assay. Recombinant activated factor VII, activated prothrombin complex concentrate, and FEIBA have been shown to be effective in treatment. The second step involves removal of factor VIII inhibitor with prednisolone or cyclophosphamide. Lastly, control the bleed by plasmapheresis or specific immunoadsorption of immunoglobulins.

Conclusion: Acquired haemophilia A is a rare disease associated with severe bleeding complications. Therefore, its prompt recognition is mandatory to initiate an initial treatment. We report a case with bleeding in the glottis which was challenging in terms of diagnosis and management.

THE INNOVATIVE USAGE OF PLATYSMA IN THE MANAGEMENT OF PHARYNGOCUTANEOUS FISTULA FOLLOWING A TOTAL LARYNGECTOMY

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Background: Pharyngocutaneous fistula (PCF) is the most common complication following total laryngectomy and is associated with increased morbidity, delay in adjuvant treatment, prolonged hospitalisation and increased treatment costs. Surgical intervention is indicated following a failed conservative approach and may include primary closure, loco-regional flaps, and free flaps. This study examines the use of platysma flap in the management of PCF post-total laryngectomy.

Aims: Identifying whether platysma flap has a role in reconstructing PCF in patients who have had a total laryngectomy and are not responding to conservative approaches.

Materials and Methods: A retrospective study recruiting consecutive patients who presented (2015-2017) with persistent PCF following total laryngectomy, despite medical management and treatment with the use of platysma flap. There were five patients; all were male with a mean age of 74 years (range 64-82). Post-operative evaluation was carried out with a formal clinical and radiological swallow assessment.

Results: Three patients demonstrated no recurrence of leakage using sole muscle flap. Two patients had post-operative complications using myocutaneous flap. The mean total hospital stay for patients following the operation was 36 days (range 13-85). The mean total days post-op for a successful swallow blue dye test was 10.2 days (range 6-15).

Discussion: The use of sole muscle flap (as opposed to myocutaneous flap) had a similar mean hospital stay (11 days) and mean post-operative successful swallow blue dye test (7.7 days). Patients are less likely to develop post-operative complications if muscle flap is utilised instead of a myocutaneous flap, no prior chemo/radiotherapy, and have no significant co-morbidities.

Conclusion: Platysma muscle flap can serve as a potential surgical intervention for a sub-selected group. Validation via large-scale longitudinal cohorts is needed. It is unlikely, at present, that this novel procedure can replace current conventional surgical intervention methods.
ATRIAL FIBRILLATION IN THE PERIOPERATIVE PERIOD DURING NON-CARDIAC SURGERY

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Background: Episodes of atrial fibrillation (AF) occurring in the perioperative period, especially during non-cardiac surgery, can lead to the destabilisation of haemodynamics, aggravate the patient's condition, and trigger acute coronary syndrome, heart failure and stroke.

Aims: To identify the predictors of paroxysmal atrial fibrillation arising in the perioperative period during non-cardiac surgery, using the preoperative examination data.

Materials and Methods: A total of 1051 patients yet to undergo abdominal, urological and articular surgeries were used for the analysis. These patients had no history of AF. Preoperative examination was performed: clinical examination; electrocardiogram (ECG) at rest; 24-hour ECG monitoring; echocardiography (EchoCG), including the evaluation of myocardial deformation using the Speckle Tracking Echocardiography (STE) method; cardiopulmonary exercise test (CPX); and, in some patients, coronary angiography.

Results: Paroxysmal AF was registered in 61 patients (5.8% of all patients included in the study). Majority (81.9%) of the episodes of AF occurred in the first five days following surgery. In the perioperative period, episodes of AF were regarded as the main cause of 25% of all non-fatal myocardial infarctions (MI), 27% of fatal MIs, and every second episode of acute cerebrovascular accident. Patients with AF significantly more often had a history of diabetes (31.2% vs.13.6%, OR 2.9, r = 0.02) and hypertension (82.0% vs.60.9%, OR 2.9, r= 0.02). AF developed more often following surgery for oncological pathologies of the abdominal cavity. The results of the EchoCG showed that the development of perioperative episodes of AF was associated with a more than 40 ml/m2 increase in the index volume of the left atrium (in 17.0% of patients with AF vs. 7.5% in patients without AF, OR 2.5, r <0.01).

Conclusion: During non-cardiac surgery in patients with diabetes mellitus, hypertension, and an increase in the volume of the left atrium greater than 40 ml/m2, as well as prior to surgery for oncological pathologies of the abdominal cavity, the risk of paroxysmal atrial fibrillation in the perioperative period is significantly higher.

SCREENING FOR ASYMPTOMATIC RENAL DISEASES AMONG SCHOOL CHILDREN IN MAN-SOURA, EGYPT

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Background: In seemingly healthy youngsters, the preliminary detection of renal diseases can help to subdue the progression of occult renal diseases.

Aims: The purpose was to identify asymptomatic renal diseases among school children aged between nine and 18 by urinalysis.

Materials and Methods: A cross-sectional study of 870 apparently healthy students was conducted in 11 teaching institutes in Mansoura using urinalysis for microscopic haematuria, proteinuria and pyuria. Students who had renal or medical illness, menstruation or congenital malformation were excluded. After screening was complete, suspected cases of haematuria, proteinuria and pyuria were referred, for free, to Mansoura Urology and Nephrology Center for further checking and follow-up.

Results: Abnormal urine findings were encountered in 200 students (approximately 23%) of the 870 students screened: the prevalence in males was 35.4% of the total cases screened whilst the prevalence in females was 17.8%. Of the 200 positive cases, haematuria and proteinuria were the most witnessed: haematuria 28.3% proteinuria 20% and pyuria 2% while combined haematuria and proteinuria cases were 4%.

Conclusion: Screening for renal diseases by urine analysis is not only indispensable to identify insidious renal diseases in asymptomatic children but it also proves a pragmatic method to define if further investigation is needed to know the exact aetiology.
ASSESSMENT AND OUTCOMES OF COLORECTAL CANCER PATIENTS WHO DO NOT UNDERGO SURGERY

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Background and Aims: Surgery is the staple curative method of treatment of colorectal cancer. However, not all patients will undergo major operations. More patients are regarded as too frail or too much cancer for operations at Lancashire Teaching Hospital Trusts (LTHT) than the national average. This study aims to determine the outcomes of these non-operative patients.

Methods: All patients discussed in the Colorectal Multi-Disciplinary Team meetings over a two-year period (April 2013 – March 2015) were categorised into the National Bowel Cancer Audit (NBOCA) groups of major resection, too little cancer, too much cancer, too frail for surgery, or others. Patient demographics were then documented and survival time post-diagnosis recorded.

Results: Four hundred and fifty-three patients (mean age 69.9 years (range 23-98 years)) were found to have colorectal cancer and classified into major resection (n=285), too little cancer (n=38), too much cancer (n=81), too frail (n=41), and others (n=8). Frail and too much cancer patients had a higher proportion of patients with more co-morbidities and a higher performance status than patients that had too little cancer. Median survival times for patients who had too much cancer or were too frail were 8.3 months and 10.4 months, respectively.

Discussion: Patients with too much cancer or who are too frail for resection surgeries have a poorer prognosis than those who are fit enough for operations. Pre-operative frailty assessment still has room to be improved, with the introduction of the quantification of frailty using frailty scores and geriatric assessment. Further studies comparing non-operative outcomes and frailty assessment will aid future decision-making.

PULMONARY FIBROSIS IN A 44-YEAR-OLD FEMALE FROM SOUTHERN ITALY AFFECTED BY HERMANSKY-PUDLAK SYNDROME

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Background: Hermansky-Pudlak syndrome, a rare autosomal recessive disorder, is characterised by oculocutaneous albinism, bleeding tendency due to platelet storage pool deficiency, and, in certain individuals, pulmonary fibrosis, granulomatous colitis and/or immunodeficiency. Among the 10 subtypes described, HPS-1, HPS-2 and HPS-4 patients tend to develop pulmonary fibrosis.

Case Study: A 44-year-old non-smoker female presented with chronic cough for 10 years and reported moderate dyspnoea worsening over the years. Physical examination revealed albinism, nystagmus and bilateral crackles upon auscultation. Pulmonary function tests showed moderately severe restriction with a TLC of 56% and a reduced DLCO. The chest-CT revealed bilateral upper lobe fibrosis, as well as traction bronchiectasis consistent with UIP pattern. The transbronchial biopsy showed foci of fibrosis and alveolar septal thickening associated with pulmonary fibrosis. A definitive diagnosis of HPS4 was established by the Endocrinological Unit based on clinical findings and a genetic test which identified a newly reported homozygous mutation.

Follow-up investigations showed worsening PFTs and fibrotic progression with extensive parenchymal involvement of the left upper lobe, plus the presence of bronchiectasis on chest-CT; consistent with pulmonary fibrosis. Oxygen therapy was promptly initiated following desaturation on room air at 88%SpO2.

Discussion: Even though pulmonary fibrosis has been largely associated with affected individuals from north-western Puerto Rico, it can also occur in other individuals, as demonstrated by this case. At present, lung transplantation is the mainstay life-prolonging treatment for HPS-PF; the supplemental oxygen is only supportive. The patient was referred to a transplant centre and evaluated to be ineligible due to significant bleeding diathesis. Influenza and pneumococcal vaccination were advised, as well as the avoidance of pulmonary irritants including tobacco use.

Conclusion: HPS-PF carries a poor prognosis with progression to death within a decade from the onset of symptoms. This highlights the compelling need for a much more effective management strategy.
POLAND SYNDROME - CLINICAL PRESENTATION AND CASE REPORT

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Background: Poland Syndrome is a congenital malformation characterised by the absence or underdevelopment of the pectoralis major, leading to a concave impression in the chest. This condition mostly affects the sternocostal head of the pectoralis major. Another common manifestation of this syndrome is brachysyndactyly on the affected side. The incidence of Poland syndrome varies in different countries, but it is estimated to be present in 1-to-3 per 100,000 individuals.

Case Study: This is a case study of a male neonate who was born in January 2017 with features of Poland syndrome. Other members of his family do not show any features of Poland syndrome. On examination, the right palm was smaller than the left, with the second and the third digits of the right hand fused with hypoplastic nails. The right nipple was deviated downwards, and a deep sacral dimple was observed. The right sternocostal head of the pectoralis major muscle was absent. These features are indicative of Poland syndrome.

Discussion: The infant was scheduled for surgery in March 2018 to split the skin between the middle and index fingers. During the same procedure, the partial web between the third, fourth and fifth fingers was removed. The patient has now gained full function in his right hand. A further operation has been planned to transfer a segment of the latissimus dorsi to fill the space of the absent right pectoralis major. This procedure will be performed when the patient is five to six years of age.

Conclusion: This case explores the diagnosis and prognosis of a patient with Poland syndrome. This report describes past and potential future management plans taken from a biopsychosocial aspect.

TO DEVELOP HAND-HELD NOTES FOR WOMEN WITH PREMATUR E OVARIAN INSUFFICIENCY (POI)

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Background: Women with POI see many healthcare professionals which leads to multiple sets of notes, many of which are either inaccessible or misplaced. Introducing handheld notes would streamline care and increase overall POI patient satisfaction.

Aims: Develop handheld notes that will: collate the patient's medical information; allow for autonomy of care; avoid unnecessary investigations; and assist healthcare professionals in understanding the patient's journey.

Methods: We designed handheld notes based on Guy's and St Thomas' (GSTT) guidelines, and patients' and healthcare providers' feedback. Ten women, who were seen in the POI Clinic at Guy's Hospital, were invited to a focus group and asked to complete two surveys. Each survey contained 16 questions; questions 1-13 measured patient satisfaction with and without handheld notes (maximum score of 65) whilst questions 14-16 asked about ease of using the notes.

Results: In a three-factor model of clinical experience (patient understanding: doctor's knowledge of the patient's journey: ease of service use), we found an improvement in patient satisfaction was enough to raise the global satisfaction score from 2.8 (without notes) to 3.6 (with notes). A t-test showed significant improvement, with a 95% confidence interval. The questions that revealed a significant improvement were related to patient understanding.

Conclusion: Overall, handheld notes have shown initial promise by improving patient perception of POI management, ease of bringing to appointments, and overall patient satisfaction. Limitations include a small sample size and limited time resulting in one data collection. This is an ongoing project with an aim to collate longitudinal evidence with a larger sample size and additional feedback from healthcare professionals.
IMPLICATIONS OF CHANGING OUTPATIENT COLORECTAL PRACTICES ON SURGICAL TRAINING

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Background and Aims: Increased patient and professional demand has led to more index procedures being performed outside the traditional setting of the Outpatient Department. The impact of this on training is undetermined. This study aimed to quantify the reduction in opportunity from a colorectal perspective.

Methods: Design: Prospective, cross-sectional study at a District General Hospital.
Sampling: All consecutive patients seen in the colorectal clinic in December 2017 were recruited.
Procedure: Data was collected with regards to the number of index procedures performed in outpatients and corroborated with clinic letters.

Results: Four hundred and sixty-two patients attended clinic during the study period. A response rate of 72.5% was achieved. Of these, 100 were targeting two-week wait referrals. Only 13 patients had a procedure performed in clinic (proctoscopy = 11; haemorrhoidal injection = 1), equating to 3% of patients. 41% patients, however, were booked for further diagnostic tests to include flexible sigmoidoscopy and colonoscopy. All target two-week wait patients did not have a procedure performed in clinic.

Discussion: A surprisingly low proportion of patients had a procedure performed in clinic, meaning that trainees have reduced exposure to procedures such as proctoscopy in an outpatient setting. The impact on two-week wait patients was particularly interesting, with these patients being more likely to be booked for further diagnostic tests immediately without a clinic procedure.

Conclusion: Increasing volume of two-week wait referrals has led to a substantially increased demand for colorectal clinics. Consequently, procedures that were traditionally performed in clinic are now being escalated straight to the endoscopy suite or operating theatre. As a result, trainees have fewer opportunities to meet their competencies.

PNEUMONIA AS A PRESENTING FEATURE OF FAMILIAL MEDITERRANEAN FEVER (FMF)

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Background: A 51-year-old South African lady of Ashkenazi Jewish descent presented with a four-week history of fever, pleuritic chest pain and shortness of breath. This was on a background of intermittent chest pain and fatigue for 18 months, and intermittent abdominal and calf pains for 30 years. Her past medical history included peptic ulceration, depression and social smoking. Her identical twin sister in Israel had a similar history of abdominal pain. Her travel and drug histories were not significant.

Case Study: She was treated for LLL pneumonia and a small exudative effusion was drained. She continued to have recurrent pyrexia with a persistently raised CRP/ESR and mild LFT derangement. Investigations for Pyrexia of Unknown Origin (PUO) (including atypical infections, CT-TAP and immunology) were unremarkable. She was discharged with doxycycline and rheumatology follow-up, but re-admitted with fever shortly after. Her symptoms resolved with colchicine, and a provisional diagnosis of Familial Mediterranean Fever (FMF) was made. FMF gene testing was negative, suggesting a novel gene mutation.

Discussion: FMF is typically caused by an autosomal recessive abnormality in the MEFV gene, with symptom onset by 20 years. At least 80% of patients experience arthralgia, abdominal pain, constipation, nausea and vomiting, fever, or myalgia. Bloods often show raised CRP/ESR, leucocytosis and raised fibrinogen during flares, which can last 12-72 hours. This patient met both the Tel-Hashomer and Livneh diagnostic criteria for FMF. Colchicine treats symptoms and prevents flares and amyloidosis.

Conclusion: This lady presented a diagnostic conundrum for rheumatology, respiratory, cardiology and haematology. She had a lengthy admission, leading to worsening anxiety. Her symptoms were vague, with little objective evidence of illness. She received multiple invasive investigations, with the risk of complications, high cost and further anxiety. The overall learning points are that FMF can present with pneumonia and should be considered in patients with PUO.
COMPARATIVE STUDY OF THE SURFACE ANATOMY OF THE AXILLARY NERVE AND ITS IMPLICATIONS ON RADIOLOGICAL INTERPRETATION

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**Background and Aims**: Being aware of the position of the axillary nerve (AN), particularly in relation to its immediate surroundings, is crucial from an anatomical standpoint. The purpose of this study was to produce a ratio between two different humeral measurements, and subsequently apply it to predict the position of the AN in X-rays. Thus, the ratio would ultimately serve as a potential diagnostic tool, specifically in cases of proximal non-displaced humeral fractures.

**Methods**: Measurements were taken from a total of 25 cadaveric upper extremities in order to produce a ratio between the superior aspect of the greater tubercle of the humerus to the lateral epicondyle (also known as the humeral length), and the superior aspect of the greater tubercle to the AN. The mean ratio produced was then used to predict the position of the AN on radiological images by measuring the distance of the humeral length and dividing the resulting figure by the mean ratio identified.

**Results**: An average ratio of 5.56cm was created; statistical analysis performed on the cadaveric data found a normal distribution of the ratios (P>0.05). In addition, further statistical analysis was carried out on the cadaveric data to determine whether upper limb side, sex or age had any effect on the ratio; it was found that there was no significant difference within any of the groups (P>0.05), and thus these factors had little, if any, effect on the ratio.

**Conclusion**: The ratio that has been attained in this study may be clinically useful to predict the position of the AN on X-rays of the upper limb. This method may be applied in cases of non-displaced proximal humeral fractures to predict the likelihood of injury to the AN. It also has the potential to reduce the risk of iatrogenic injuries during surgical intervention.

FACTORS INFLUENCING MEDICAL STUDENTS’ CHOICE OF COUNTRY TO PRACTICE

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**Background and Aims**: Shortages of doctors in different specialities can have a strong negative influence on the healthcare system, particularly in low- and middle-income countries. This study aims to explore the reasons for which medical students intend to emigrate rather than work in their country of study.

**Materials and Methods**: A total of 39 International Coordinators (IC) were recruited from an online international medical student forum, after sending out a letter of participation. The IC was instructed to circulate an identical survey to medical students in the penultimate and final years of their respective university. The Chi-squared test was used to analyse results.

**Results**: A total of 628 students (74 from Africa, 14 from Asia, 24 from South America and 514 from Europe) participated. Two-thirds of students stated that they were planning to work where they studied following graduation. Females, those aged 25 and under, with previous degrees, or from an Asian university were more inclined to stay in their country of study. The most common reasons for this decision included job development potential (28%), patriotism (19%), and family and friend influences (14%), with 1% of students expressing that they do not have any other choice. There was a statistically significant difference in the reasons between different genders (p=0.048), age groups (p=0.001), those with or without previous degrees (p=0.001), and the continent of study (p=0.029).

**Discussion**: The large number of students considering moving away from the country of study following graduation is consistent with the trend of increased multiculturalism and globalisation. Analysis of the influential factors among medical students may provide an insight for employers to create pull factors to fulfill job demands.

**Conclusion**: One-third of students are not planning to work in the country from which they obtain their medical degree. There was a significant difference between student decisions and influences based on student demographics.
THE VALUE OF SIMULATION TEACHING DURING INDUCTION FOR FOUNDATION TRAINEES TO TRAUMA AND ORTHOPAEDICS

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Background and Aims: Rotating into a new specialty is often daunting for Foundation Year 1 (FY1) doctors. In order to deliver a clinically-relevant and practical introduction to their clinical attachment, we developed a patient story simulation session as part of their departmental induction. The defined outcomes of the session were to:

- give a basic introduction to approaching trauma.
- give Foundation Trainees the basic knowledge to deal with a common Orthopaedic emergency.
- improve trainees' confidence in patient management.

Methods: A simulated patient pathway was designed and six different simulation stations were created and conducted at induction for the new intake of FY1 trainees. Pre- and post-course questionnaires were completed to quantify the value of teaching and assess the level of confidence.

Results: Overall results showed an increase in the level of confidence across all set learning objectives. Before the session, 100% of trainees were 'Not at all' to 'Neutral' in their confidence level towards: 1) managing trauma calls; 2) managing open fractures; 3) applying a POP cast; 4) intra-operative use of drivers, screws and plates; and 5) reading and presenting radiographs. Post-course feedback showed that 57-100% of trainees in the above-mentioned stations were “Confident” to “Very confident” in their ability to perform the skills taught in a clinical environment.

Discussion: The results show that simulation can be used to improve confidence in patient management for trainees starting in a new specialty.

Conclusion: Simulation can be used at induction for FY1 trainees entering a new specialty to assist with confidence levels and to provide basic skills that will be useful within that specialty. Entering into a new specialty with these skills taught at the beginning of the placement should allow for easier transition and a better overall experience in the specialty during their placement. Further study is ongoing to reassess the value of simulation at the end of trainees’ placements.

THE EFFECTIVENESS OF A BRIEF HEALTH EDUCATION INTERVENTION IN PROMOTING EMOTIONAL WELLBEING IN SCHOOLCHILDREN

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Background and Aims: Mental illness is staggeringly prevalent in adolescents, with 10% of children aged between five and 16 being clinically diagnosed with a mental health condition. Despite NICE Guidelines and Public Health England recommendations, there is currently no national provision of mental health teaching. However, numerous educational interventions have demonstrated success. This study aims to investigate the effectiveness of an educational intervention on the promotion of emotional wellbeing amongst schoolchildren.

Methods: A ‘quasi-experimental’ study with mixed-methods, and wait-list with follow-up design. An educational intervention was delivered to 84 Health and Social Care Class Students (five male; 79 female) in year groups 10 to 13 at an Essex Secondary School. Questionnaires relating to mood and wellbeing (WEMWBS, PANAS) were delivered at various time-points. Baseline wellbeing data was also collected from an additional 19 students.

Results: Schoolchildren had an average level of wellbeing of 41/70, considerably lower than that of the general population (50/70). Between t-tests also demonstrated an immediate large improvement on the Positive Affect Scale (mean 1.59, d=1.04, p<0.001), and a larger effect at significantly reducing Negative Affect (mean 2.33, d=2.16, p<0.001).

Discussion: In keeping with the literature, wellbeing was found to be variable, with males having a higher score. Additionally, feedback questionnaires demonstrated that a short classroom-based intervention affected individuals positively. The study was limited by the timeframe constraints, and was only conducted in one school. The findings of a low level of wellbeing have important implications for students, parents, teachers and the wider society.

Conclusion: Wellbeing in schoolchildren can be considerably low. Successful short-term classroom-based interventions can be conducted to improve the state of mental health of schoolchildren. Future research should be conducted in a wider age range and various schools to validate these findings.
ASSESSING THE USE OF E-LEARNING METHODS TO PROVIDE NUTRITIONAL EDUCATION WITHIN THE UNDERGRADUATE MEDICAL CURRICULUM FOR THIRD YEAR STUDENTS AT BARTS AND THE LONDON MEDICAL SCHOOL: A PILOT AIMED TO ASSESS AND IMPROVE CONFIDENCE AND KNOWLEDGE

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Background and Aims: The relationship between food, nutrition and health is complex. Yet, current medical professionals receive limited teaching in the field of nutrition. It is therefore imperative that the current medical curriculum adapts to change this and that learning outcomes are relevant to what is seen in clinical practice. A UK Undergraduate Curriculum in Nutrition exists within Tomorrow’s Doctors for Graduates. Yet, learning objectives are often dispersed across existing modules, diluting the content of what is recognised as core learning objectives. By equipping ‘Tomorrow’s Doctors’ with a multifaceted learning tool, we aim to improve confidence and knowledge in nutritional education and translate this into clinical practice.

Methods: This pilot study aimed to address this through adopting an e-learning module which targets year three medical students, utilising clinical vignettes in line with the outcomes within the GMC’s brief. By providing clinical scenarios encompassing nutritional components, a key aim was to assess students’ confidence, knowledge and perception of a given topic. On completion, strengths, weaknesses and perceptions would be identified, guiding future development. A nutritional module was devised by a clinical dietitian and GP with a specialist role in medical education. A half-day teaching session was allocated, with relevant pre-reading material and questionnaire. This provided a comparable baseline and post-assessment of confidence and knowledge. The module was subsequently run across three successive groups (120 students/term) with an anticipated total number of 360 students for the academic year 2017-2018.

Results: Term 1 results (n=120 students) show a significant improvement in both knowledge and confidence. Data has also been collected in terms of satisfaction of content, volume and access to the online content. Further knowledge will be tested during the end of year exams.

Discussion: Incorporating this specific nutrition teaching into the undergraduate curriculum at an early clinical phase means knowledge can be built upon both theoretically and practically before qualification, and has the potential to be revisited and studied at further depth later in the course as the “spiral curriculum” permits. Further developments include looking towards a national accreditation of the module with the British Dietetic Association, and research across several medical schools in terms of developing undergraduate curriculum content on nutrition.

CLINICAL MEDICAL STUDENT INTEREST AND KNOWLEDGE OF ANATOMY: A FEEDBACK ANALYSIS OF THE ROYAL COLLEGE OF SURGEONS EDINBURGH ‘WADE’ WORKSHOPS

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Background and Aims: Time spent on undergraduate anatomy teaching has declined since the introduction of modern medical curricula. Clinical medical students, taught via a case-based learning spiral curriculum, were evaluated on their interest and knowledge of anatomy through aggregated feedback from the annual Royal College of Surgeons of Edinburgh (RCSEd) Workshop in Surgical Anatomy in 2016 and 2017, with the aim to improve similar future events.

Methods: A feedback form was used with responses to questions rated on a 4-point scale – 1 (strongly disagree) to 4 (strongly agree). The questions asked if students were able to explain why knowledge of anatomy is important for surgical practice and, secondly, if they felt able to cite examples of anatomical knowledge applied to surgical problems in the head and neck, trunk and limbs. Open-ended feedback from workshop participants was also evaluated.

Results: Students (n = 40) strongly agreed that they were able to explain why anatomy is important in surgical practice; mean = 80% (2016, 84% 2017, 76%). Students were able to cite examples of the application of anatomical knowledge to surgical problems in the head, neck, trunk and limbs; mean = 75% (2016, 74% 2017, 76%). Improvements included suggestions to provide more time for the workshop (n = 17), to give post-workshop handouts (n = 10), and to offer pre-workshop preparatory reading material (n = 5).

Conclusion: The study showed that the RCSEd Wade Surgical Anatomy Workshops are valuable to medical students.
COMPARATIVE STUDY OF THE RADIAL GROOVE IN HUMAN CADavers, DRY BONE, AND X-RAYS

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Background and Aims: The objective of this study was to identify a method that could be used to predict the radial groove’s position on X-rays. This is clinically relevant due to the radial nerve’s close association with the radial groove along the posterior surface of the humeral shaft. When injury occurs to this portion of the arm, such a method may be used to ascertain from an X-ray whether damage to the radial nerve is likely to have occurred. This study aims to compare the radial groove in human cadavers, dry bone specimens, and X-rays.

Materials and Methods: Measurements were taken from 20 cadaveric arms and 20 dry bone specimens. The ratio produced between the measurements of the greater tubercle to the lateral epicondyle, and the greater tubercle to the start of the radial groove, was then used to predict the position of the radial groove on 16 X-rays.

Results: The cadaver cohort’s average humeral length, and average distance between the greater tubercle and the radial groove, was 28.5 cm and 13.2 cm, respectively. The dry bone cohort’s average humeral length, and average distance between the greater tubercle and the radial groove, was 29.1 cm and 13.29 cm, respectively. Statistical analysis performed on the cadaver and dry bone data for humeral length found both to be of a normal distribution (P>0.05). The ratio of both groups was generated. An average of the two ratios (2.19) was then used to predict the location of the radial groove on 16 X-rays.

Conclusion: The ratio that has been obtained in this study may be clinically useful to predict the position of the radial nerve via the radial groove on an X-ray. It also has the potential to reduce the incidence of iatrogenic radial nerve injuries.

THE APPLICATION OF 3D PRINTED, KINAESTHETIC MODELS FOR UNDERGRADUATE MEDICAL ANATOMY TEACHING

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Background and Aims: Technological advances have facilitated the creation of various novel teaching tools that improve the effectiveness of anatomy teaching, including 3D printed models and the creation of kinaesthetic models, where three-dimensional anatomy models can be physically manipulated to illustrate functional anatomy. The aim of this project is to combine these two modalities, create a 3D printed kinaesthetic anatomy model, and then test its efficacy in teaching undergraduate medical students the biomechanics of the rotator cuff in comparison to the currently used plastic models.

Methods: Anonymised CT scans were transferred to ©Cura Software 3.1 for modification and 3D printing. The 3D printed model was then modified to illustrate the biomechanics of movement of the rotator cuff. Twenty-one volunteers were split into two groups of intervention (3D printed model) and control (plastic model), and attended the same teaching tutorial and quiz at the end. The overall scores and qualitative data on the models were then analysed for statistically-significant differences.

Results: A kinaesthetic 3D printed model was successfully produced to illustrate all movements of the rotator cuff accurately. No significant differences were found in the overall score of students, but the 3D printed model was generally found to be more “helpful” and “recommended for future teaching”.

Conclusion: A kinaesthetic, 3D printed model did not increase overall exam performance compared to a plastic model. It was, however, significantly preferred by students and, with some refinement, 3D printed kinaesthetic anatomy models may be used in the future for undergraduate medical teaching.
KNOWLEDGE OF HEALTHY LIFESTYLES AMONGST GULF MEDICAL UNIVERSITY STUDENTS AND HEALTHCARE PROFESSIONALS IN AJMAN, UAE

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Background: Non-communicable diseases (NCD) are now at the forefront of mortality rates; this is especially true in the UAE. Leading a healthy lifestyle has been shown to improve NCD morbidity rates and knowledge of what a healthy lifestyle consists of is pivotal to practicing one.

Aims: To assess the knowledge of healthy lifestyles according to the WHO/CDC guidelines amongst university students and healthcare professionals (HCPs). To determine the sources of knowledge of healthy lifestyles.

Methods: The sample size (670 participants) included students and HCPs. A self-administered questionnaire was used with a cross-sectional study design to assess knowledge of physical activity, diet, sleep and tobacco use.

Results: This study shows that 3.7% and 55.4% of participants were aware of the CDC guidelines of physical activity and sleep, respectively. A total of 0.3% of participants were aware of the USDA guidelines for a healthy diet, and 88% of participants understood tobacco use carried a severe or moderate risk to one's health. The most accurate sources of knowledge for healthy lifestyles were from: friends for physical activity (82.5%), academic institutions for diet (23.9%), academic institutions for sleep (90.7%), and family for tobacco use (98.2%).

Discussion: The majority of participants had good knowledge of healthy lifestyle guidelines, except for diet and physical activity. Research conducted in India, US and Canada supports our findings for physical activity, sleep and tobacco use, respectively. The most common sources of knowledge included media, academic institutions, and family. Similar findings were reported in Europe and Canada, amongst others.

Conclusion: Participants displayed appropriate knowledge regarding the CDC guidelines for sleep, and portrayed awareness of the harmful risks of tobacco use. However, they had incredibly poor knowledge of the CDC guidelines for physical activity and the USDA guidelines for a healthy diet. Academic institutions were the most reliable sources of information for most domains.
CRITERIA FOR THE DIAGNOSTIC OF THE METABOLIC SYNDROME IN THE ADMINISTRATIVE PERSONNEL OF THE UNIVERSITY

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Background and Aims: The Metabolic Syndrome (MetS) is the association of metabolic alterations constituted by central obesity, dyslipidemia, hyperglycemia, and arterial hypertension. This study aims to identify the presence of MetS with the criteria of the International Diabetes Federation (IDF), American Heart Association (AHA), the third report of the National Cholesterol Education Program (ATP III), and the World Health Organization (WHO), in the administrative staff at the Centro Universitario de la Costa.

Methods: Descriptive transversal study, with 76 Mexican participants between 21 and 43 years of age. They took socioeconomic and anthropometric measurements, fasting glucose, arterial tension, and serum testing; the latter represented a risk factor according to the criteria for the diagnostic of MetS: IDF, AHA, ATP III and the WHO.

Results: The obtained data shows that the cases that represented MetS were 6.57% according to the ATP III criteria, 10.52% according to the AHA, 11.84% according to the criteria for the WHO, and 18.42% according to the IDF, the latter being the one with the highest number of affected.

Conclusion: The importance of early diagnosis and the criteria of the IDF are paramount. These factors allow the start of a treatment with a more preventive focus. This will lower the morbimortality associated with this syndrome in the long-term.

RENAL OUTCOME AMONG SAUDI CHILDREN WITH POSTERIOR URETHRAL VALVE: WHEN TO WORRY?

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Background: Posterior urethral valve (PUV) is one of the most common causes of lower urinary tract obstruction in male neonates. The incidence of PUV is <1 in 8000 fetuses. Most fetuses suffer miscarriage, and those continued with the pregnancy will develop a life-threatening obstructive uropathy which requires urgent surgical intervention. Two surgical approaches are used for the management of PUV: endoscopic valve ablation and urinary diversion.

Aims: To evaluate the renal outcome of endoscopic valve ablation versus urinary tract diversion.

Methods: An observational, retrospective cohort study involving all patients who were diagnosed with PUV at King Abulaziz Medical City from 1998 to 2016. The patients were divided into two groups according to the type of intervention: those in Group 1 were offered endoscopic fulguration whilst those in Group 2 were offered urinary diversion. Group 1 comprised of 24 patients (61%) and Group 2 was composed of 15 patients (39%) (either vesicostomy 56% or ureterostomy 44%). The data was collected from the medical records and collated in a data collection sheet.

Results: Thirty-nine patients with PUV were managed, 23% of which were preterm. Chronic kidney disease (CKD) developed in 17 patients (45%), and 18% developed end-stage renal disease (ESRD). CKD developed in 60% of patients with urinary diversion compared to 33% of patients undergoing endoscopic ablation; OR = 3 (95%CI = 0.8 - 11.4, P value = 0.09). Patients who underwent diversion had a higher risk of developing ESRD; OR = 2.55 (95%CI = 0.48 - 13.46).

Conclusion: Endoscopic fulguration has a better renal outcome compared to urinary diversion. PUV is a rare pathology, but our patients had poor renal outcomes in comparison to international studies.
## AWARENESS OF EYE DISEASES CAUSED BY DIABETES AMONG DIABETIC IN-PATIENTS AT KING ABDULAZIZ UNIVERSITY HOSPITAL, JEDDAH, SAUDI ARABIA

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**Background:** Saudi Arabia is the second most common country affected by diabetes mellitus in the Middle East, and is the seventh in the world. Almost 10% of diabetic patients develop visual impairment and blindness due to ocular complications. These could be prevented by increasing awareness.

**Aims:** To assess the awareness of diabetic inpatients about diabetic eye disease at King Abdulaziz University Hospital (KAUH).

**Methods:** A cross-sectional study was conducted in KAUH, Jeddah, Saudi Arabia in 2017 amongst 300 diabetic inpatients in KAUH between 2014 and 2017. The data was collected through a telephone-based interview and analysis was carried out using Chi-square test.

**Results:** The study involved 241 out of 300 patients; 140 (58.1%) were male. Of these patients, 80.9% knew of diabetic eye disease. The study found that educational level and economic status played a key role in the awareness of eye diseases, and 79.7%, 41.9%, and 32.8% of patients were aware that retinopathy, cataract, and glaucoma, respectively, can be caused by diabetes mellitus.

**Conclusion:** Most of those with diabetes were aware of ophthalmologic complications. Despite their awareness of diabetic retinopathy, more than half of them were unaware that it is curable. Hence, more public health education is needed.

## A RARE CASE OF IVEMARK SYNDROME (RIGHT ISOMERISM) WITH AN EXTREMELY UNFAVOURABLE COMBINATION OF HEART DEFECTS

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**Background:** Ivemark syndrome is a rare congenital abnormality typified by heart defects, asplenia and disarrangement of the viscera. The complexity in managing this condition means a high mortality rate in 60% of patients before they reach the first year of their lives, and only 5-10% of affected individuals reach adulthood.

**Case Study:** A three-month-old male Caucasian child was hospitalised in the Paediatric Department due to skin cyanosis, decreased appetite and presence of mucus in stool following antibiotic therapy for acute bronchitis. According to obstetric history, the child was born full-term. Birth weight was 3750g, body length was 53cm, and Apgar rating scale showed 8/8 points. During the third trimester of pregnancy, the mother developed pyelonephritis and anaemia. From the first day of life, the child was hospitalised in the neonatal department with a diagnosis of congenital bilateral pneumonia with cardiovascular syndrome. He was on mechanical ventilation for seven days after birth, and was consulted on the tenth day by cardiology and cardiovascular surgery specialists. He was subsequently treated symptomatically until he died.

**Results:** CT scan of the chest and abdomen revealed: asplenia; full A-V communication (common ventricle); aorta located away from the right ventricle; pulmonary atresia; patent ductus arteriosus; total anomalous pulmonary venous drainage; stomach located on the left; mid-location of liver; non-visualised spleen; kidney without features; large-sized adrenal glands; pancreas located on the right; intestinal malrotation; and small thymus.

**Discussion:** Ivemark syndrome is rare, and cases from the literature are sporadic and multifactorial, indicating varying combinations of presenting anomalies. Hence, evidence to support a specific management approach is lacking. We describe here a case of a three-month-old male child diagnosed with syndrome of heterotaxy.

**Conclusion:** In this case, an extremely unfavourable combination of heart defects made radical correction impossible, eventually leading to the death of child. It is therefore imperative to create an awareness of this rare syndrome in order to motivate prenatal detection of this disease.
AN IATROGENIC TRACHEOESOPHAGEAL FISTULA IN AN ELDERLY PATIENT COMPLICATING A CENTRAL VENOUS CATHETERISATION PROCEDURE: A CASE REPORT

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Background: Adult tracheoesophageal fistula (TEF) is a rare clinical occurrence that results from various causes. The authors report on an elderly patient with an ESRD who is on regular renal haemodialysis and who had an iatrogenic TEF complicating a central venous catheter (CVC) insertion.

Case Study: A 65-year-old Sudanese male with ESRD presented with recurrent cough and choking following meals. The symptoms started soon after he had a CVC inserted at a nephrology centre; he had never experienced these symptoms before prior to the procedure. Barium swallow demonstrated an abnormal flow of contrast into the trachea and main bronchi, with a suspected high tracheoesophageal fistula. A CT of the chest confirmed fistula position, but there were no other abnormal findings in the trachea, main bronchi, lungs or oesophagus. Esophagoscopy and bronchoscopy revealed a large TEF in the cervical oesophagus at 21cm from the incisors. No other luminal pathologies were detected in the oesophagus or bronchi. He underwent surgical repair of the fistula with a vascularised strap flap as it was not amenable to clipping or stenting per bronchoscopist advice. Biopsy was negative for any chronic inflammatory or neoplastic findings. His symptoms resolved with no recurrence over a year.

Discussion: Aetiology of adult TEF involves a variety of neoplastic, inflammatory, traumatic and iatrogenic causes. A detailed history, a thorough physical examination, and findings of serial investigations did not reveal any specific cause of our patient's fistula, apart from the CVC procedure he underwent. The onset of complaints correlated with the CVC insertion as he had never experienced TEF symptoms before. Isolated tracheal and oesophageal injuries are well-known complications of CVC insertion.

Conclusion: Adult TEF is a rare clinical entity. Nevertheless, a careful clinical work-up is needed to establish the aetiology of the fistula in elderly patients. CVC insertion could be a possible cause of an adult TEF.

EXTRACELLULAR MATRIX-DERIVED HYDROGEL FROM DECELLULARISED UTERINE TISSUE: A SYNTHESIS AND ANALYSIS

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Background and Aims: Female infertility is estimated to affect 72.4 million, or 9% of, women of reproductive age. There are multiple causes such as Asherman syndrome, or symptomatic intrauterine adhesions, which results from inappropriate tissue healing after damage to the endometrium's basalis layer (i.e., from surgery, infection or abortion). These adhesions can obstruct sperm passage or embryo implantation, leading to infertility. A modern approach to tissue healing is thus warranted. Extracellular matrix (ECM) derived hydrogel is a novel scaffold that mimics the micro-environment of native tissues. Recently, numerous studies have illustrated its wide-ranging applications in regenerative medicine with one ongoing Phase I Clinical Trial on patients with post-ST elevation myocardial infarction. However, its application in uterine tissues has not been explored or characterised. This study aims to further investigate this.

Methods: Uterine tissues were harvested from female Sprague Dawley rats and decellularised using high hydrostatic pressure of 980 MPa. Histological analysis and dsDNA assay were performed to assess decellularisation. Samples were then lyophilised, digested, and used to form hydrogel as per modified Freytes protocol. Cytotoxicity was examined by culturing rat endometrial stromal cells on the hydrogel for two, four or seven days, followed by live/dead double staining assay.

Results: After seven days, hydrogel surface was almost entirely covered with living cells, with few dead cells detected.

Conclusion: We demonstrated for the first time that ECM-derived hydrogel could be successfully synthesised from decellularised uterine tissues. Its capability to host living cells hints at its potential to facilitate wound healing in the uterus. More research is needed to analyse its characteristics and utilities in clinical medicine.

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Background: Surgical revascularisation of the vertebral artery by transposition to the common carotid artery provides secondary prevention of posterior circulation strokes in patients who have not responded to best medical therapy or endovascular therapy. Surgical revascularisation provides an alternative treatment, however, it is associated with a high complication rate.

Aims: To demonstrate the anatomical relations of the vertebral artery to surrounding structures commonly damaged during surgical revascularisation and hypothesise methods to avoid damage.

Methods: Hemi neck dissections were undertaken with reference to published surgical approaches to demonstrate the relevant anatomy. Findings: The cervical sympathetic trunk was adhered to the vertebral artery and at greatest risk of injury. The thoracic duct, recurrent laryngeal nerve, phrenic nerve, brachial plexus and apex of the lung were encountered in each approach.

Discussion: The study showed that despite an evidence-based approach to the vertebral artery there is still potential to damage structures early in the procedure. Complications can be minimised by following a set of key principles intraoperatively to protect the vertebral artery. Conclusion: The results from the study provide awareness of the anatomical relations of the vertebral artery in a surgical context and potential methods to avoid complications. To continue developing the safest approach to the vertebral artery further dissections are needed to trial the hypothesised methods and directly involve surgeons specialising in the procedure.

THE EFFECTIVENESS OF FASCIA ILLIACA BLOCKS AS PAIN RELIEF IN NECK OF FEMUR (NOF) FRACTURES AT UNIVERSITY HOSPITAL OF NORTH TEES

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Background and Aims: NoF fractures are one of the commonest causes of admission to an Orthopaedic Unit. Fascia iliaca blocks are cheaper than, and have a reduced side effect profile compared to, opioids. This study aims to examine the effectiveness of fascia iliaca blocks as pain relief in NoF fractures.

Methods: Data was collected prospectively between the dates of 22/02/18 and 27/03/18. The inclusion criteria for this clinical audit included skeletally mature patients over the age of 18 years who were admitted via the trauma service at UHNT. Patient notes were analysed from the Emergency Department, with the pain relief given to each patient recorded alongside multiple pain scores (between 0-10): post-block, six hours post-block, and 15 hours post-block. Identical measurements were recorded for patients who did not receive a block. Additionally, age, gender, side effects (likely caused by pain relief), and any alterations in AMTS prior to surgical input were recorded.

Results: A total of 20 patients were identified from the trauma list. Twelve female and eight male patients were recorded in this audit. The patients varied in age, from 66 to 97 years old (mean 81.1). Sixty percent of the cohort were administered a fascia iliaca block. There was a mean reduction of 4.22 in the pain score immediately post-block, compared to an average instantaneous reduction of 2.5 in the pain score for patients who did not receive a block. An average of 0.54 side effects were experienced by patients who had received a block, compared to a mean of two side effects in the group of patients who were treated without a block.

Discussion: We have identified that delivering a series of teaching sessions on the technique used in performing this block to Emergency Department Physicians, Orthopaedic SHOs and Foundation Doctors can act as a route to improving the proportion of NoF patients who receive a block. Additionally, altering the treatment guidelines for the NoF pathway in Accident and Emergency should help further.

Conclusion: The benefits in using fascia iliaca blocks have been confirmed by this audit.
PREDICTING HEART DISEASE USING ARTIFICIAL INTELLIGENCE AND MACHINE LEARNING

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Background and Aims: Internationally, heart disease is a leading cause of death amongst patients. To improve their chances of survival, early and accurate diagnoses of heart disease is essential. In this research project, a computational model based on machine learning algorithms is created to aid medical professionals in optimising the accuracy of heart disease diagnoses.

Methods: The computational model was developed using ensemble machine learning meta-algorithms and networks for heart disease diagnosis. The model was applied to heart disease data sets from 878 patients (53% male; 47% female) in hospitals and medical institutions located in the United States, Hungary, and Switzerland. Data from 50% of the patients was used to train the machine learning model, and the remaining 50% of the patient data was used for testing performance.

Results: The developed machine learning models achieved overall heart disease diagnostic accuracies of 94% based on training data set results and 87% based on testing performance results, exceeding the accuracies of previously published research.

Discussion: The machine learning model was evaluated for accuracy, using F-score, Area under the Curve (AUC), and K-S measures. Its average model F-score was 0.89 and average K-S score was 60%. The ensemble machine learning model prediction results were more accurate compared to those of other data mining approaches for heart disease diagnosis, including those of decision trees (80%) and Bagging algorithm (82%).

Conclusion: Computer-aided heart disease diagnoses based on machine learning algorithms are clinically useful in aiding medical professionals globally to improve patient outcomes. Furthermore, for patients in areas with fewer heart disease specialists, the machine learning model can serve as a cost-effective option for earlier detection and diagnosis of heart disease.

TEXTURE ANALYSIS USE IN ASPECT SCORING OF ACUTE STROKE

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Background and Aims: Acute strokes are one of the major causes of mortality in the UK and are medical emergencies requiring immediate admission to hospital. Upon admission, a brain CT is acquired to check for contraindications before thrombolysis is administered. A 10-point ASPECT score can then be calculated by a senior radiologist using the CT images of patients presenting with middle cerebral artery (MCA) strokes. ASPECTS quantifies ischaemic changes and allows for prediction of outcomes. This study aims to determine if texture analysis can be used to find ischaemic changes that are indistinguishable to the human eye.

Methods: In this retrospective single centre study, the scans of 40 patients were equally divided into groups with either low ASPECTS (7) or high ASPECTS (10). The images were collected from Ninewells PACS network and anonymised using the inbuilt software. Images were analysed by manually placing Regions of Interest (ROI’s) on ASPECT regions at the two basal-ganglionic and supra-ganglionic slices using the DICOM viewer “RadiAnt” and texture analysis software “MaZda”.

Discussion: Limitations of this study included difficulty in placing ROI’s manually due to user learning curve and some problems with the programme “MaZda” recognising the DICOM image format of the CTS.

Conclusion: At this stage, we are currently awaiting results. However, qualitative results can be discussed. There are a number of pros and cons to the use of TA. Pros include that TA can be conducted on CT images but with some major limitations specific to my project, as previously discussed.
ALL CABG PATIENTS WHO HAVE NO CONTRAINDICATIONS: DO THEY GET PERIOPERATIVE BETA BLOCKERS?

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**Background and Aims:** Atrial fibrillation (AF) is the most common heart rhythm disorder. AF has been reported in up to 15 to 40% of patients in the early postoperative period after CABG. Beta blocker administration reduces the incidence of postoperative atrial fibrillation (POAF) from 30–40% to 12–16% after CABG. This study aims to determine if all CABG patients without contraindications receive β-blockers perioperatively.

**Methods:** Retrospective data was collected. Consecutive 50 patients treated with isolated CABG between August 2017 and October 2017 were included. Patients were excluded if they had physician-documented contraindications to β-blocker therapy, or underwent CABG combined with valvular or other cardiac surgery.

**Results:** According to the standard guidelines, all patient undergoing CABG should receive β-blockers in the immediate postoperative period, namely within 24 hours. β-Blockers were continued perioperatively only in 12% of patients. However, in 20% of patients, it was resumed on the first postoperative day. In 48% of patients, β-Blockers were resumed on the second postoperative day.

**Discussion:** According to the ESC Guidelines issued in August 2009, β-Blockers are recommended in patients with known ischaemic heart disease or myocardial ischaemia on preoperative testing (IB). In our audit, those patients who received β-blockers on the day of surgery or the following morning (Continued + Restarted on 1st POD) met the standard guidelines. Thus, according to the data, (12%+20%) 32% of the patients met the standard guidelines. To compare the rate of AF, we divided the patients into two groups: Group A, who followed the guidelines (16 patients), and Group B, who resumed β-Blockers 48 hours onwards (34 patients). In Group A, only one patient developed postoperative AF whereas in Group B, 11 patients developed postoperative AF.

**Conclusion:** β-Blockers significantly reduce the incidence of AF after CABG. Considerable attention must also be focused on understanding and improving β-blocker use during the perioperative period. From this audit, it can be said that standards were met – partially.

PREVALENCE, CLINICAL PATTERN AND DETERMINANTS OF ALLERGIC RHINITIS AMONGST CHILDREN AGED ONE TO SIX IN JEDDAH CITY, KSA

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**Background and Aims:** Allergic rhinitis (AR) is an important global public health problem because of its markedly increased prevalence worldwide. It has a considerable impact on the quality of children’s life. It affects social life, sleep, school and work, and its economic impact is substantial. This study aimed to evaluate the link between allergic rhinitis and its related determinants among children in Jeddah City.

**Methods:** A cross-sectional analytical study was selected. The questionnaire was distributed to 748 children aged between one and six years who were attending two hospitals in the North and South of Jeddah. Data was collected from the mothers or caregivers of healthy children attending with them using: interview questionnaire, anthropometric measurements, and ISAAC core questionnaire on asthma and allergic disorders.

**Results:** This study was conducted on children aged between one and six years (n=748): 357 males (47.7%) and 391 females (52.3%). A total of 62.8% of the children were from the North of Jeddah, while 37.2% were from the South of Jeddah. Saudi children constituted 47.6% of the sample, while 52.4% were non-Saudis. Out of the 748 children studied, AR was found in 128 children (17.0%). Smoking of the father, the North of Jeddah, high social class families, and family history of allergies were more common among children with AR. Children of male gender, born premature, formula-fed, did not consume milk daily or consume a sweet diet were more likely to suffer from AR. Children with AR were more prone to repeated respiratory tract infection and had asthma symptoms. Only 63.3% of ISAAC diagnosed AR were diagnosed by the doctor as having AR.

**Conclusion:** AR is a major health problem among children aged between one and six years. Socio-economic characteristics, environmental factors and diet are important determinants of AR. Obesity and asthma, as well as family history of allergies, are important clinical disorders associated with AR.
POLICY COHERENCE, TRADE LIBERALISATION AND OBESITY: A CASE STUDY OF NEW ZEALAND'S TRADE OBJECTIVES AND DEVELOPMENT COMMITMENTS IN THE SOUTH PACIFIC

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Background and Aims: The small domestic economies and limited self-sufficiency of South Pacific islands has led to dependency on foreign nations for development assistance and importation of essential goods. These islands also face a non-communicable disease (NCD) crisis, containing some of the highest rates of obesity in the world. This study aims to consider the connection between trade liberalisation and obesity in the South Pacific.

Methods: A literature review was conducted to investigate the relationship between trade liberalisation and obesity in the South Pacific. Twenty-six articles were reviewed to investigate the pathways of influence and extent of the relationship in this region, with a focus on the nutritional impact of New Zealand's exports. This was followed by a thematic analysis of New Zealand’s food, trade and development policies in the South Pacific.

Results: Trade liberalisation has facilitated the entry of nutritionally-inferior food products to Pacific nations and reduced subsidies available for domestic production, increasing food-import dependency. Free-trade agreements have also reduced the revenue available to re-invest in health systems over-burdened by the NCD crisis. New Zealand’s policy objectives include a reduction in islanders’ cardiovascular disease and tobacco use and an increase in domestic export revenue through further free-trade agreements.

Discussion: Tensions are evident between New Zealand's objectives to maximise export revenue, particularly from food and beverages, and commitments to reduce NCD incidence. The failure to incorporate health outcomes into economic policies threatens to undermine New Zealand's commitments to policy coherence in the Sustainable Development Goals.

Conclusion: The prioritisation New Zealand affords to its economic objectives underpins its failure to achieve policy coherence in the South Pacific. New Zealand appears reluctant to acknowledge its contribution to the NCD crisis, opposing the development of trade-restrictive health policies. Consideration must be given to the disparities in global economic strength between New Zealand and the island nations, with monitoring systems implemented to guarantee greater future policy coherence.

PATIENT-RELATED OUTCOMES AND EXPERIENCE MEASURES (PROMS/PREMS) RELATING TO ORAL SURGICAL PROCEDURES UNDER IV SEDATION: A SUMMARY OF A LOCAL PROJECT AND A SUGGESTED MOVE TOWARDS A STANDARDISED QUESTIONNAIRE

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Background and Aims: Conscious IV sedation (IVS) is a useful adjunct for many patients undergoing oral surgical or dental procedures. The clinical team should strive to improve the quality, safety and satisfaction of the service provided for patients. Evidence of this is becoming increasingly important in the commissioning of NHS services. To improve a service, we must first evaluate aspects of it in order to find areas of improvement. The patient’s perspective is one major aspect of the quality of a service; patients could receive the best treatment but still be unhappy. Patient Reported Experience and Outcome Measures (PREMS/PROMS) help us to gain a general idea of the patient's perspective. This study aims to explore the PREMS and PROMS relating to oral surgical procedures performed under IVS at a local hospital.

Methods: We studied the PREMS and PROMS of the oral surgery outpatient IVS service at the Royal Devon and Exeter Hospital (RD&E). A questionnaire was formulated which included criteria from two national commissioning documents. One hundred questionnaires were sent by post with pre-paid return envelopes. All patients underwent dental extraction(s) and were over 18 years of age. Forty-three questionnaires were returned and results analysed.

Results: Patient satisfaction was generally very high, on a Likert scale from 5 (excellent) to 1 (very poor), 90% answered excellent while 98% answered 4 or higher. A total of 95% of patients would definitely and 5% would probably recommend the service to a friend or family member. Seven percent of patients needed to seek advice or assistance post-procedure, and 5% required additional surgery.

Conclusion: Overall, patients were happy with the service provided: 98% would rate it 4 or higher out of 5. Waiting times were identified as the commonest reason for dissatisfaction, which could be overcome by increasing clinic sizes. The widespread use of standardised PROMS/PREMS criteria would help evaluate and compare IVS services consistently across providers. The development of a validated questionnaire would be of future interest.
PAEDIATRIC AUTISM IN PRIMARY CARE

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Background: Autism Spectrum Disorder (ASD) is a multifactorial, lifelong neurodevelopmental disorder. Several factors have contributed to its aetiology, including genetics, neurological changes, immune dysregulations and environmental elements. ASD is characterised by pervasive communication and impairments in social interaction, in addition to restricted and repetitive behavioural patterns which usually arise in childhood.

Aims: This project reviews the diagnostic process and management framework of paediatric ASD in primary care. Furthermore, it aims to provide a means for educating parents and carers about the condition.

Methods: National guidelines and international criteria (ICD-10) were used to create a protocol for the identification, referral and management of ASD in general practice. Based on information gathered from literature search, a leaflet was also created to increase public awareness of the condition.

Results: A multi-step protocol was produced for General Practitioner use and guidance. Additionally, a case study involving two autistic siblings was conducted to identify challenges encountered by families and the support received. Finally, a leaflet on the condition was created to improve parents’ and carers’ understanding of ASD and to address their concerns.

Conclusion: Being a lifelong condition, this study has reiterated the necessity of early diagnosis and management of ASD for better future outcomes. Due to the diversity of presentations, referral pathways and service provisions differ. However, a set protocol and multidisciplinary approach to every suspected ASD case is chief to maximise the children’s potential in the long run. Moreover, providing education and support helps families to cope with the diagnosis, as well as the physical and psychological demands of caring for a child with a disability.


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Background and Aims: The human larynx is responsible for manipulating sound waves, altering their pitch within a range of 100 to 50,000 Hz. These alterations are the result of rapid movements of the vocal folds, membranous tissue connecting the anterior and posterior cartilaginous surfaces of the larynx. Responsibility for moving these folds falls to the skeleton of the larynx, made of a number of cartilages. The size and orientation of the cartilages therefore have a dramatic effect on voice. Size in particular is influenced by the hormonal influences exhibited during puberty, creating a great sexual dimorphism. This aim of this study was to evaluate the differences in the anatomy of the larynx between genders.

Methods: This study used dissection to expose the larger cartilages of the larynx, the hyoid bone and the vocal folds, and measures them across 20 different cadavers; 10 male and 10 female.

Results: The results were analysed using one-way and two-way ANOVAs and Kruskal-Wallis to investigate differences, and Pearson’s and Spearman’s tests for correlations.

Discussion: The study showed a significant difference in the larynges between genders, as expected. The epiglottis was similar to the size of the laryngeal outlet both in width and length, confirming its functional role. Other lesser correlations may show further potential links that result from the developmental processes in the growth of the larynx.
VALIDATION OF THE OMRON HBP-1300 IN NORMOTENSIVE AND HYPERTENSIVE PREGNANCIES ACCORDING TO THE BRITISH HYPERTENSION SOCIETY PROTOCOL (1993)

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**Background:** Automated blood pressure devices have become a popular alternative to the ‘gold standard’ manual sphygmomanometer. However, few automated devices have been shown to be accurate in pregnancy and pre-eclampsia. The haemodynamic profile of pregnancy, and the challenges this presents to most automated devices, emphasise the importance of separate validation in these subgroups.

**Aims:** The objective of this study is to validate the Omron HBP-1300 monitor for use in normotensive and hypertensive pregnancies using the British Hypertension Society (BHS) protocol.

**Materials and Methods:** Nine consecutive same-arm blood pressure measurements were taken, alternating between the mercury and the Omron HBP-1300. The final seven measurements were used in the analysis. The device was graded according to the BHS criteria.

**Results:** Fifty-three women were recruited, and 50 were included in the study. The Omron HBP-1300 achieved an A/A grade in both systolic and diastolic measurements in normotensive and hypertensive pregnant women, using the BHS protocol.

**Discussion:** The majority of readings were within 5mmHg. Nonetheless, the maximum over-estimation was 25 mmHg. This may be an anomaly. However, it could have a large impact on patient diagnosis and management. Due to the morbidity associated with hypertensive pregnancies, it is imperative that validation studies continue to evaluate the accuracy of automated devices in pregnancy. This will enable better availability of alternatives to mercury sphygmomanometers, and improve upon available technology.

**Conclusion:** The Omron HBP-1300 has achieved the highest-grade possible using the BHS validation criteria and, therefore, can be recommended for clinical use in both normotensive and hypertensive pregnancies.

A SERVICE EVALUATION OF THE MANAGEMENT OF DEVELOPMENTAL DYSPLASIA OF THE HIP (DDH) AT THE CHILDREN'S HOSPITAL FOR WALES

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**Background:** DDH occurs in 4.9/1000 UK live births and is a leading cause of childhood disability. Early intervention using a Pavlik Harness or hip spica cast can fully correct the abnormality. Late intervention leads to hip instability and arthroplasty.

**Aims:** To evaluate parental perspectives of the screening, diagnosis and management of DDH at the Children’s Hospital for Wales (CHfW), and to define the standard of care achieved.

**Methods:** We designed a questionnaire to collect parental feedback about the service provided at the hospital. We also utilised the CHU9D validated paediatric outcomes score to collect patient well-being data.

**Results:** We gave questionnaires to 105 parents and 91 paediatric patients. In the duration of this project, insufficient responses were received. The project is still ongoing until sufficient responses from parents and patients are received.

**Discussion:** Long-term consequences of missed diagnosis or untreated DDH come at a high cost for society, the individual and the family. Such a condition that is treatable during infancy must be studied in order to reduce the incidence of preventable adulthood disability.

**Conclusion:** Current literature scarcely utilises paediatric patient-reported outcomes or evaluates parental experiences. More data is needed to facilitate future practice.
A PATIENT WITH POLYMYOSITIS

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Background: This case report is a reflection on a seventy-three-year-old lady who I saw in the Rheumatology ward at the Glasgow Royal Infirmary whilst completing a student-selected study module. In the report, I have reflected on the condition with which the patient was diagnosed - polymyositis - and have related the typical clinical presentation of the condition to the specific presentation of this patient.

Case Study: Polymyositis is an idiopathic inflammatory myopathy characterised by striated muscle inflammation causing muscle weakness. The inflammatory myopathies are a group of disorders sharing the pathophysiology of immune-mediated muscle injury, but they each differ histologically. The epidemiology of polymyositis is 6-8/100 000 cases/year, and the female to male ratio is 2:1. Polymyositis affects areas of striated muscle, and complications include dysphagia, respiratory muscle weakness leading to respiratory failure, and cardiovascular involvement. The pathophysiology involves infiltration of striated muscle fibres with inflammatory cells, resulting in necrosis of myocytes and inhibition of normal physiological muscle contraction.

Discussion: The patient in this case report suffered from many disease complications, including opportunistic infections due to treatment regimes, and her admission was a very long and complex one. The risk of cancer in a female with polymyositis is increased 3.4 times. It was discovered during her admission that this patient had a cystic lesion in the pancreas which is still to be investigated, but it would not be surprising if this lesion turned out to be a malignancy due to the high associated risk with polymyositis.

Conclusion: This rare case demonstrates the severe and sudden debilitating impact that polymyositis has on a patient, preventing him or her from being able to stand up unaided, let alone continue to live independently as this particular patient could prior to becoming ill.

BASE OF TONGUE ABSCESS - A RARE BUT POTENTIAL DIFFERENTIAL TO TONSILLITIS: A CASE REPORT

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Background: Tongue abscesses can be classified as either anterior or posterior. Due to the tongue's resistance to infection, abscesses are rare, especially posterior abscesses. Causes of infection of the base of the tongue include lingual tonsillitis, infected remnants of a thyroglossal cyst, and lower molar teeth infections.

Case Study: A 37-year-old lady presented to the GP clinic and Accident and Emergency multiple times and was treated for tonsillitis with antibiotics following a one-month history of worsening throat pain, tongue swelling, dysphagia, and breathlessness with a feeling of constriction in her airway. Worsening of symptoms and raised inflammatory markers required admission, IV antibiotics and ENT input. Flexible nasoendoscopy showed swelling of base of tongue on the right side with narrowed but adequate airway. CT scan of the neck showed a large heterogeneous, multi-cystic collection of 5x4.5x4cm at the base of tongue on the right side, crossing the midline and involving the intrinsic tongue muscle. Incision and drainage, and a tracheotomy, were performed.

Discussion: Tongue defence mechanisms include the mobility of the tongue and presence of saliva allowing for persistent cleansing, the thick keratinized mucosa acting as a barrier, and the tongue's rich blood supply and lymphatic drainage. Airway obstruction may present acutely due to rapidly expanding base of tongue abscess, management of which is particularly challenging and accounted for 40% of all cases (72 out of 184 patients) reported to the 4th UK National Audit Project of Major Airway Complications (NAP 4). Significant recommendations arising from the review of these cases have been instrumental in guiding the assessment, planning, and management strategies of patients with an obstructed airway to optimise their outcome.

Conclusion: Although challenging, it is vital to identify and diagnose tongue base abscesses as the resulting swelling may become life-threatening due to upper airway compromise.
CHRONIC ANAEMIA IN A 30-YEAR-OLD GENTLEMAN: UNUSUAL CASE OF MECKEL’S DIVERTICULUM

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Background: Meckel’s diverticulum is the most common congenital abnormality of the small bowel and, when symptomatic, can present as intussusceptions.

Case Study: A 30-year-old man of oriental origin with no significant past medical history presented to the Emergency Department with a five-hour history of central abdominal pain, generalised abdominal tenderness, and tachycardia on examination. This was his fourth admission in the last five months. Given the multiple presentations of abdominal pain, anaemia with a normal OGD and sigmoidoscopy, a CT scan confirmed a small bowel intussusception secondary to Meckel’s diverticulum. The patient was taken to theatre for a short segment small bowel resection.

Discussion: Intussusception is defined as the invagination of a proximal segment of bowel into the lumen of an adjacent segment, and is described in only 5% of all cases in the adult population. Almost 90% of intussusception cases in adults are secondary to intra-luminal pathologies: neoplasms, lipomas, inflammatory lesions and, rarely in this case, a Meckel’s diverticulum. Whilst a Meckel’s diverticulum causing intussusception is rare, a thorough and appropriate investigation of the gastrointestinal tract is required in young adults with repeated presentations, chronic anaemia and normal endoscopies. Recognising this early and confirming the diagnosis is essential to enhance patient outcomes.

DIABETES STRUCTURED EDUCATION PROGRAMME - EFFECT ON DIABETES CONTROL

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Background and Aims: The National Institute for Health and Care Excellence (NICE) UK recommends structured education programme (SEDP) as an integral part of diabetes management. This project aims to study the effect of patients’ participation in SEDP on diabetes control.

Methods: Specially designed questionnaires were distributed to all diabetes patients ≥18 years old attending the diabetes centre at the Royal Liverpool University Hospital (RLUH) from 13/02/2018 to 01/03/2018. Data on treatment targets and presence of complications were collected from Diabetes Register and hospital database.

Results: Out of 221 intended patients, we have a total of 142 respondents, of whom 55 have attended SEDP. Respondents who attended SEDP were more likely to achieve HbA1c ≤ 58 mmol/mol (Odds Ratio 1.302, 95% CI: 0.643 to 2.636, P=0.464) and cholesterol <5 mmol/L (Odds Ratio 1.294, 95% CI: 0.484 to 3.460, P=0.607), but less likely to achieve target of blood pressure ≤ 140/80 (Odds Ratio 0.687, 95% CI: 0.315 to 1.495, P=0.344). They also had reduced odds of getting cardiovascular complications (Odds Ratio 0.698, 95% CI: 0.301 to 1.623, P=0.404), peripheral vascular complications (Odds Ratio 0.500, 95% CI: 0.284 to 1.623, P=0.315), cerebrovascular complications (Odds Ratio 0.775, 95% CI: 0.222 to 2.706, P=0.689) and nephropathy (Odds Ratio 0.657, 95% CI: 0.284 to 1.519, P=0.325). No significant difference on the odds of having neuropathy (Odds Ratio 1.073, 95% CI: 0.492 to 2.342, P=0.860) and diabetic eye disease (Odds Ratio 1.046, 95% CI: 0.529 to 2.069, P=0.897).

Conclusion: Participation in SEDP could increase the likelihood of achieving treatment targets and reduce the likelihood of developing most long-term diabetes complications.
EVALUATION OF THE EFFECTIVENESS OF ‘FUNDAMENTALS OF MEDICAL SCIENCE FOR NON-MEDICAL PROFESSIONALS’ COURSE

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Background and Aims: Many non-medical professionals, such as medical translators and interpreters, medical journalists, healthcare lawyers, and medical engineers, have limited knowledge of medical science but are required to work closely with medical professionals. This study considers how a course that proffers a focused overview of fundamental concepts and terminologies pertaining to medical science can assist such professionals in their careers.

Description: Entitled ‘Fundamentals of Medical Science for Non-Medical Professionals’, this structured course was held on two occasions (June 2017 and February 2018) at Cardiff University with different cohorts of medical linguists. The interactive discussions provided delegates with a focused overview of fundamental concepts and terminologies within the field of medical science. The course also offered an insight into the basics of medical history taking, reading and interpreting a medical report, construing expert opinions, understanding hospital transfer reports, and comprehending patient discharge documentations.

Results: The feedback (n=15) revealed that the course enabled the delegates to develop a good understanding of medical concepts and terminologies. In response to the question ‘Do you feel more confident in understanding medical terminologies as a result of attending this course?’, 100% selected ‘Most Certainly’ in 2017 whilst 67% answered ‘Most Certainly’, 22% answered ‘Certainly’, and 11% answered ‘Neutral’ in 2018. When asked if they were more confident in reading a medical document as a result of attending the course, 100% responded ‘Most Certainly’ or ‘Certainly’ in 2017 and 90% responded ‘Most Certainly’ or ‘Certainly’ in 2018. One hundred percent of delegates belonging to the 2017 cohort felt ‘Very Confident’ in applying the knowledge gained through the course to the context of medical translational work, whilst 89% answered between ‘Very Confident’ and ‘Confident’ in 2018.

Discussion: The value of specialist and focused courses that offer professionals a greater insight into an area with which they frequently interact is notable. Although the cohort in this study were medical linguists, all non-medical professionals working closely with the healthcare profession will benefit from similar courses. This will ultimately facilitate an efficient and fruitful working partnership between the professions.

A QUALITY IMPROVEMENT PROJECT TO ASSESS AND REDUCE INFECTION RATES IN PATIENTS WITH DISTAL PHALANGEAL FRACTURES TREATED WITH KIRSCHNER WIRE FIXATION

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Background: Recently, concerns were voiced amongst colleagues in the Department of Plastic Surgery at the Queen Victoria Hospital, East Grinstead that cases of infection amongst patients treated with K-wire fixation for fractures of the distal phalanx were increasing. The use of K-wires for fracture fixation is known to increase the likelihood of patients developing deep infections, namely osteomyelitis, and this can have potentially devastating outcomes.

Aims: To determine the incidence of superficial and deep infection amongst patients who had been treated with K-wire fixation at the QVH for fractures of the distal phalanx within the previous 12 months. To use an analysis of the data collected, in conjunction with a systematic review of the literature, to devise a local protocol for the safe and appropriate use of K-wires for fractures of the distal phalanx.

Methods: We retrospectively reviewed the notes of all 38 adult and paediatric patients who underwent this procedure at the QVH during the period of December 2016 to December 2017.

Results: The overall rate of infection after K-wire fixation in this patient group was 18.4%(n=7). The proportion of patients who developed osteomyelitis was 10.5%(n=4).

Discussion: We have made a protocol available in theatres and in trauma clinic that contains a number of recommendations and points for surgeons to consider before using K-wires to treat fractures of the distal phalanx. These recommendations pertain to the timing of fixation, mechanism of injury and fracture location.

Conclusion: We are currently unsatisfied with the rate of infection in this patient group. Thus, we have implemented an intervention designed to redress this problem that we hope will have demonstrable results when we prospectively re-investigate in 12 months.
A SYSTEMATIC REVIEW AND META-ANALYSIS OF THE USE OF TISSUE-DERIVED THROMBIN SEALANT IN DRAINLESS PAROTIDECTOMY

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Background: Drains are traditionally placed after parotid surgery, with the intention of reducing post-operative seroma and haematoma formation. This usually denotes an inpatient bed until the drain is removed, depending on an acceptable low output. With the availability of tissue-derived thrombin sealants, wound closure in facial plastics and head and neck procedures can now be facilitated without the use of a drain in the dead space.

Aims: To assess whether tissue sealant improves wound-related outcomes in parotidectomy and allows for a drainless procedure.

Methods: A systematic literature review was performed using a standardised published methodology and custom database search strategy. Methodological assessment was performed using the Quality Assessment of Diagnostic Accuracy Studies (QUADAS-2) tool, and a fixed-effect meta-analysis of the combined complications was performed.

Results: Eleven studies were identified relating to parotidectomies using tissue sealants. Three of them have shown statistically significant decrease in drain output. Tissue sealant use enabled a shorter in-hospital stay (0-2 days) compared to the conventional method (1-5 days), thus lowering the overall cost per case. There was also a tendency for tissue sealants to reduce complication rates, including haematomas and seromas (Odds Ratio 6.25 [1.13, 34.45], 95% CI, I² = 47%, P = 0.06).

Conclusion: The use of drains post-parotidectomy is superseded by tissue sealant due to the shorter admission time and the lower risk of post-operative complications.

IS THERE A NEW KIND OF DOCTOR? AN INTERVIEW-BASED STUDY OF FORMER BBT TRAINEES AND THEIR SUPERVISORS

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Background and Aims: As a result of an ageing population, more patients are presenting with chronic, multi-system diseases, increasing the complexity of care. The Shape of Training Review recommended changes to the training of doctors to meet these healthcare needs, including training doctors with more generalist skills. Broad-based training (BBT) was an approach designed to achieve this. Following Foundation Year 2, trainees spend six months in four specialities: General Practice, General Medicine, Paediatrics and Psychiatry.

This longitudinal study aims to consider whether BBT has developed a new kind of doctor. We explore how well BBT prepares trainees for their current role, how BBT impacts on clinical practice, whether trainees adopt a holistic approach to care, and what lessons can be learned for the future.

Methods: Following ethical approval, 14 BBT trainees and three clinical supervisors were interviewed by telephone. Recordings were then transcribed and thematically analysed using NVivo.

Results and Discussion: Exposure to broad specialities appeared to increase preparedness to meet changing demographics. Greater awareness of NHS facilities and resources improved decision-making and had a positive influence on the management of complex patients. Exposure to BBT specialities improved understanding of the patient journey and specialty integration. This led trainees to think in different ways and to adopt a more holistic approach to care. Despite difficulty disentangling the specific effects of BBT from other clinical experience, participants identified lessons for the future, including calls for a programme with similar principles to run in the future.

Conclusion: Changing patient needs had led to an agreement that postgraduate training needs to change. BBT represents one approach. Although BBT appears to develop generalist skills and holistic care, it is not possible, without further research, to conclude that BBT produces a new kind of doctor. However, our results show the benefits of a broader style of training and identify important lessons for the future of medical education.
THE COMMUNICATION AND CLINICAL ACTION OF BLOOD CULTURE GRAM STAIN RESULTS IN THE HOSPITAL SETTING

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Background: Acutely unwell hospital patients may have microbiological blood cultures performed to guide antimicrobial therapy. Gram staining is carried out in the laboratory. The results are communicated to the responsible ward clinician.

Aims: To assess the communication of blood culture Gram stain results during normal working hours (NWH) and out of hours (OOH). To ascertain whether the responsible clinician acts on the results received from the consultant or from the laboratory. To assess whether this action is clearly documented in the patient’s medical notes.

Methods: Multi-departmental data was collected prospectively in January 2018. Thirty-three patient records were analysed over a 10-day period. Databases were accessed to view blood culture Gram stain results, microbiology consultant and laboratory communication, drugs prescribed, and on-call bleep documentation. Multiple visits were made to the wards on which patients were based to view notes. A collection pro-forma was created. No exclusion criteria.

Results: Five patients passed away before antimicrobial clinical action could take place. There was one outpatient case. Thus, of the 27 remaining: Doctors visited the wards and acted on the clinical information received from the microbiology consultant (NWH) or laboratory (OOH) in 41% of cases. Written documentation of this was available in 37% of cases.

Discussion: Appropriate clinical responses to blood culture Gram stain results are beneficial for patient safety and antimicrobial stewardship. There are differences in the pattern of results collected from OOH cases versus NWH cases; staffing numbers and demands may be causative. Escalation wards, as opposed to permanent wards, also pose difficulties. Education amongst clinicians is paramount. Wider discussion is encouraged and better practice may be influenced.

Conclusion: The prompt clinical action and documentation of blood culture results is suboptimal at present; adherence rate is 41% and 37%, respectively. This requires improvement. Re-audit shall take place in late 2018.

IS THERE AN INCREASE IN FRACTURE NECK OF FEMUR PATIENTS ADMITTED FROM WITHIN THE HOSPITAL YEAR ON YEAR OVER A THREE-YEAR PERIOD AT A LEVEL 1 TRAUMA CENTRE?

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Background: Between 4% and 7% of hip fractures occur in hospital. Mortality is higher in those who sustain hip fracture in hospital than those who sustain a fracture in the community. With an ageing population and multiple co-morbidities, there is an increased length of stay in hospital for elderly patients. These patients can suffer from acute delirium, often on the back of memory problems, and can be susceptible to falls.

Aims: To assess the variation and trend of the number of fracture neck of femur (NoF) patients admitted from within the hospital due to falls on the wards at a Level 1 Trauma Centre.

Methods: We used our NoF database that holds prospectively-collected data on all the NoF admissions. We then used Clinical Portal to look through the patient notes for information on admission dates, type of surgery, discharge dates, and AMTS. We extracted demographic data, and 30-day and long-term mortality. Age, gender, American Society of Anaesthesiologists (ASA) grade, and time to surgery were also recorded.

Results: During the study period of 2015 to 2017, 1533 patients were admitted with fracture NoF. Of these, 58 (3.78%) were sustained while in hospital. There were 23 in-hospital fractures in 2015, 21 in 2016, and 14 in 2017. More females (30/58 (52%)) sustained a fracture in hospital than males, and the median age was 80 years. The most common procedure performed was DHS (23/58 (40%)), followed by a hemiarthroplasty (17/58 (29%)). Overall mortality was 19% (11/58) and 30-day mortality was 16% (9/58).

Conclusion: Our in-hospital fracture NoF patients is less (3.78%) than those reported in the literature. We have seen a slight reduction in the number of patients admitted from in-hospital falls over the three-year period. However, as these patients are the sickest, it therefore is unsurprising to see slightly higher mortality rate compared to those NoF patients admitted from outside the hospital.
AN AUDIT OF PATIENTS REFERRED WITH CATARACT WHO PROCEED TO SURGERY

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Background: Cataract surgery is the most common surgical procedure carried out on the NHS. Literature shows poor levels of quality in cataract referrals, resulting in many patients not being operated on.

Aims: To evaluate cataract referrals made to a large tertiary hospital and to establish the number of referrals that resulted in an operation, as well as the reasons for those that did not. In addition, to see if there was any difference between over- and under-75s when being listed for operations.

Methods: Clinic letters were reviewed for all patients seen within a four-week period in 2016 to assess information regarding their surgery, or the reason that they did not undergo surgery.

Results: Seventy patients were seen, with a mean age of 73 and a female: male ratio of 2.3:1. A total of 99% (n=69) of patients were correctly diagnosed, with 63% (n=44) listed for surgery. The most common reason for not being listed was that the patient was asymptomatic, accounting for 42% (n=11) of cases. Fifty-three percent (n=37) were under-75, of which 70% (n=26) were listed for an operation. Forty-three percent (n=33) were over-75, of which 55% (n=18) were listed for an operation.

Discussion: The epidemiological data retrieved from this cohort is similar to national figures, despite the relatively small scale of this study. Increasing age decreases the likelihood of having a cataract operation, which may be explained by the fact that more elderly patients are more likely to have routine eye checks and therefore be referred while being asymptomatic.

Conclusion: This study found that almost all patients had a correct diagnosis of cataract. The most common reason for not listing a patient for an operation was that they were asymptomatic. This was more common in the over-75 cohort. More care should be taken when referring patients for cataract assessment to reduce unnecessary numbers of asymptomatic patients, resulting in a more efficient service.

PNEUMOTHORAX AS A COMPLICATION OF PERCUTANEOUS LUNG BIOPSIES: TECHNICAL APPROACHES TO IMPROVE PATIENT SAFETY

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Background and Aims: A QIP introduced in our Radiology Department focused on the incidence and outcome of pneumothorax as a complication of CT-guided percutaneous lung biopsy. This aim of this study was to evaluate the effectiveness of handover in surgical patients.

Methods: This retrospective clinical audit studied a cohort of patients who underwent CT-guided lung biopsy within a six-month period between March and September 2017. Insignia image information system reports and Electronic Patient Records (EPR) were examined, searching for immediate (identified on CT scan) and delayed post-procedure pneumothoraxes (PTX) seen on chest radiograph during the follow-up period.

Results: A total of 27 patients were studied; 17 females (63%) and 10 males (37%). The mean age was 68 (range 18 - 83). In terms of lesion type, 4.7% were reported as a mass, 22.2% as multiple nodules, and 33.3% as a solitary nodule. The immediate PTX seen on CT scan accounted for 30% of all lung biopsies and 3.7% required chest drain within the surveillance period. The main biopsy technique was coaxial with 18G or 20G needle, and the number of passes varied from one to four.

Discussion: A national survey about current practice and complication rates of percutaneous lung biopsies in the UK gathered a total of 1860 auditable data from 39 centres. Among other complications, this survey showed that pneumothorax accounted for 20.5% of biopsies and that those requiring chest drain was 3.1%. This QIP demonstrated that the incidence of PTX and chest drains in our centre surpassed the national survey by 9.5% and 0.6%, respectively. Although cutting needle with coaxial technique increases diagnostic yield, it is also associated with an increased complication rate.

Conclusion: Following the first results of this QIP, prophylactic techniques, for example self-expanding hydrogel plug into the pleural space, and therapeutic post-procedure strategies such as Heimlich valves have been introduced in our department as they have the potential to reduce complications and hospital admissions.
THE PERCEPTIONS OF HEALTHCARE PROFESSIONALS ON BARRIERS TO THE UPTAKE OF STRUCTURED DIABETES EDUCATION PROGRAMME

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Background and Aims: Diabetes is a chronic and progressive disease that causes significant levels of morbidity. Despite NICE's recommendation to offer all patients with diabetes education within a year of their diagnosis, the number of patients taking part in structured education remains poor. This study aimed to assess perceptions of healthcare professionals (HCPs) relating to factors which influence the promotion of diabetes education.

Methods: A structured questionnaire with 10 questions was used to gather responses from HCPs within the diabetes team at a tertiary centre to assess barriers limiting uptake of diabetes structured education. Retrospective data on patient referrals to diabetes education between 2015 and 2017 was audited against the standard outlined in NICE guideline (NG17) and the National Diabetes Audit.

Results: Of the 305 patients offered structured education between 2015 and 2017, nearly half had been diabetic for more than 10 years, with only one-tenth receiving a diagnosis within a year. Of those offered education, almost 90% attended the programme. Fifty-seven HCPs completed the questionnaire, with 80% being aware of the education programme and 54% knowing how to refer patients. Only half of the HCPs encouraged their patients to attend education.

Discussion: The poor referral rates are likely to be the result of time limitations, perceived low-utility of education, inadequate understanding of the referral process, and service limitations. It would be appropriate to prioritise those patients with the greatest clinical requirement for education, such as those patients with erroneous HbA1c values, morbid obesity, poor compliance with hypoglycaemic agents, poor engagement with the diabetes team, and those patients who face the greatest risk of diabetic complications.

Conclusion: The tertiary centre was not meeting all diabetic patients being offered structured education. A key area for improvement includes developing a package of education within the Trust around the benefits and components of structured education and the referral process to increase awareness among HCPs.

IMPROVING SURGICAL HANOVER

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Background and Aims: Handover is an important part of safe, effective patient care. The on-call team covers General and Breast Surgery, with juniors also covering Urology and ENT. Problems were highlighted with a variation in length, quality and lead of handover, and standard of the handover list. The aim was to detect problems with surgical handover and to assess for compliance to ‘Safe handover: Guidance from the working time directive working party’ by RCS England.

Methods: A survey was sent to surgical doctors of all grades which included questions about the main issues with the handover and handover list, the order in which patients should be seen, and suggestions for change to enhance current practice. We received 22 responses in four weeks.

Results: Five common themes of issues were found:
- Professionalism
- Communication
- Patient Safety
- Structure and Organisation
- Urology, Breast and ENT

We received many suggestions, which guided the new handover list template and some of the changes implemented.

Discussion: Based on the results, a ‘3 steps to a safe surgical handover’ was created. This involves:
1) Introductions
2) FY/CT handover of Urology/ENT patients and jobs
3) Registrar handover starting with patients for theatre, unwell patients of which to be aware, followed by post-take patients

Posters of the ‘3 steps’ have been created for the handover room and printed as handouts. We have also created register and checklists to be completed at each handover in order to monitor changes and compliance. Part of the recommendations has been to include a lecture on how on-calls and handovers work for surgical and FY1 induction. A new list template has been made to clearly identify unwell patients and patients for theatre, as well as include useful contacts.

Conclusion: With the changes implemented, there has been an increase in structured handovers with fewer missed patients. This project is being reassessed with checklists and re-surveying.
**PREVALENCE OF THE Atherogenic Index AS A RISK FACTOR FOR THE DEVELOPMENT OF METABOLIC SYNDROME IN ADMINISTRATIVES AT CENTRO UNIVERSITARIO DE LA COSTA, PUERTO VALLARTA: PROPOSAL OF NUTRITIONAL APPROACH**

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**Background and Aims:** The atherogenic index (IA) is useful for predicting peripheral arterial disease linked to cardiovascular events. This is calculated by dividing total cholesterol (CT) and high-density lipoprotein cholesterol (HDL-C). In the same way, hypertension and hypertriglyceridemia are relevant abnormalities in the metabolic syndrome (MS). According to la Encuesta Nacional de Salud y Nutrición de Medio Camino (ENSANUT 2016), 28% of the Mexican population has high cholesterol problems. For this reason, this study aims to identify its frequency as a risk factor in MS and to propose strategies for its prevention and treatment.

**Methods:** A descriptive, cross-sectional study was carried out with a total of 76 participants aged between 21 and 43 years. Anthropometric measures were taken, fasting glucose and blood pressure were recorded, and serum tests were performed. In the first instance, the results were interpreted according to the International Diabetes Federation criteria to find at least one risk factor after the lipid profile test was applied.

**Results:** A high prevalence of AI was found in 10.52%. It was also found that 42% of the population had some of the risk factors for MS, of which 25% had hypertriglyceridemia, 15.6% had abnormal HDL, 34.37% had total cholesterol elevated, and 25% had an atherogenic index greater than 4.5.

**Discussion:** The consumption of monounsaturated fatty acids (MUFA) and polyunsaturated fatty acids (PUFA), as well as the Mediterranean diet and DASH, favour the control of blood pressure, coagulation, endothelial function and insulin resistance. This has beneficial effects on the prevention and treatment of MS, and such dietary suggestions should thus be implemented in the guidelines.

**Conclusion:** Both the prevalence of high AI and the frequency of dyslipidemia are similar in relation to that reported in recent years in our country. Hence, there is a need to act early with a specific nutritional approach.

**FRACTURED NECK OF FEMUR ADMISSION PROFORMA - A COST SAVING INTERVENTION WHICH IMPROVES PATIENT CARE**

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**Background and Aims:** Neck of femur fracture (NOF) carries a high mortality risk. Quality of NOF care in NHS Trusts in England has been audited against national standards, and financial incentives are provided to Trusts meeting these standards as Best Practice Tariff (BPT). We conducted a closed loop audit in an attempt to assess whether the care of NOF patients in our Trust meets national standards, and to determine its associated cost implications.

**Methods:** This closed loop audit was conducted over four-week periods by retrospectively analysing electronic patient records used in our Trust. Criteria assessed included: operation within 36 hours of admission; adequate pre-operative investigation (blood tests, ECG, and chest X-rays); analgesia; and documentation of Abbreviated Mental Test (AMTS) and pre-admission mobility. Following the first audit cycle, we introduced a printed departmental clerking proforma highlighting the essential criteria required for BPT, and stressed the mandatory completion of these forms for all NOF admissions.

**Results:** There were 33 and 35 patients in the first and second audit cycle, respectively. The department's performance (1st vs. 2nd audit cycle) was as follows: AMTS (58% vs. 89%); documentation of pre-admission mobility (73% vs. 89%); operation within 36 hours (76% vs. 86%); analgesia (91% vs. 97%); and pre-operative investigation (92% vs. 94%).

**Discussion:** Introduction of clerking proformas improved performance across all criteria monitored; more significantly in documentation of AMTS and pre-admission mobility, and time to surgery (within 36 hours). This also brought in £13000.00 as BPT incentive.

**Conclusion:** We demonstrated how the effective use of a simple, well-designed clerking proforma has led to (i) less duplication of work; (ii) better adherence to best practice; and (iii) cost savings, which allows for money to be invested in other facets of care. All these ultimately translate to better care for this group of patients with fragility fracture.
A COST-EFFECTIVENESS REVIEW OF MOHS MICROGRAPHIC SURGERY VERSUS SURGICAL EXCISION FOR TREATING PRIMARY NON-MELANOMA SKIN CANCERS

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Background: Non-melanoma skin cancers (NMSC) are the most prevalent skin cancers, with high financial burden for the NHS. It is generally accepted that Mohs micrographic surgery (MMS) is the treatment with the lowest recurrence rates.

Aims: To compare the cost-effectiveness of MMS vs. surgical excision (SE) for the treatment of NMSC.

Methods: We conducted a literature search on PubMed, Google Scholar and Springer-Link, using the search terms; ‘Mohs’, ‘Surgical Excision’, ‘Skin Cancer’, and ‘Cost’. We found 71 publications and, after applying our exclusion criteria, we were left with six papers to review (Figure 1).

Results: Summarised below (Table 1).

<table>
<thead>
<tr>
<th>Study</th>
<th>Year of Publication</th>
<th>Number of tumours (n)</th>
<th>NMSC type</th>
<th>End Point</th>
<th>Cost of MMS (US $)</th>
<th>Cost of SE (US $)</th>
<th>Difference (SE - MMS) (US $)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sebaratnam et al.</td>
<td>2016</td>
<td>n = 309</td>
<td>BCC</td>
<td>Initial margin clearance</td>
<td>$491.89</td>
<td>$445.65</td>
<td>-$46.24</td>
</tr>
<tr>
<td>Ravitskyi et al.</td>
<td>2012</td>
<td>n = 406</td>
<td>BCC and SCC</td>
<td>5 years</td>
<td>$804.72</td>
<td>$1025.83</td>
<td>$221</td>
</tr>
<tr>
<td>Cook and Zitelli</td>
<td>1998</td>
<td>n = 400</td>
<td>BCC and SCC</td>
<td>5 years</td>
<td>$1243</td>
<td>$1167</td>
<td>-$76</td>
</tr>
<tr>
<td>Bialy et al.</td>
<td>2004</td>
<td>n = 98</td>
<td>BCC and SCC</td>
<td>Initial margin clearance</td>
<td>$937</td>
<td>$1029</td>
<td>$92</td>
</tr>
<tr>
<td>Seidler et al.</td>
<td>2009</td>
<td>n = 98</td>
<td>BCC and SCC</td>
<td>5 years</td>
<td>$957</td>
<td>$1248</td>
<td>$292</td>
</tr>
<tr>
<td>Mosterd et al.</td>
<td>2008</td>
<td>n = 270</td>
<td>BCC</td>
<td>5 years</td>
<td>$1522.12</td>
<td>$1207.45</td>
<td>-$314.67</td>
</tr>
<tr>
<td>Mean</td>
<td></td>
<td>n = 263.5</td>
<td></td>
<td></td>
<td>$992.62</td>
<td>$1020.49</td>
<td>$28.02</td>
</tr>
</tbody>
</table>

(Rated used to exchange all figures into USD were AUD 1:0.79 USD and Euro 1:1.22 USD)

Table 1: Cost analysis of MMS versus SE

Discussion: The mean cost difference was $28.02 in favour of MMS. However, the range of values varied significantly, from MMS being cheaper by $292 to being more expensive by $314.67 (£258) compared to SE.

Conclusion: Generally, MMS has been found to have lower recurrences and higher QALY’s compared to SE, and may also be more cost-effective. In an era of increased NHS strain, more rigorous trials are required to determine cost-effectiveness.
The World Journal of Medical Education & Research (WJMER) is the online publication of the Doctors Academy Group of Educational Establishments. It aims to promote academia and research amongst all members of the multi-disciplinary healthcare team including doctors, dentists, scientists, and students of these specialties from all parts of the world. The journal intends to encourage the healthy transfer of knowledge, opinions and expertise between those who have the benefit of cutting-edge technology and those who need to innovate within their resource constraints. It is our hope that this interaction will help develop medical knowledge & enhance the possibility of providing optimal clinical care in different settings all over the world.