203. Assessment of a Modified Antibiotic Prophylaxis Open Fracture Protocol Brandon Hill, PharmD1; Kamla Sanasi-Bhola, MD2; Stella Okoye, MD3; Margaret Madera, PharmD Candidate1; Janie Ferren, PharmD, PharmD Candidate1; Julie Ann Justo, PharmD, MS, BCPS-AQ ID1; and P Brandon Bookstaver, PharmD, FCCP, FIDSA, AAHIVP1, Palmetto Health Richland, Columbia, South Carolina. 1University of South Carolina School of Medicine, Columbia, South Carolina, 2University of South Carolina College of Pharmacy, Columbia, South Carolina, 2Department of Clinical Pharmacy and Outcomes Sciences, University of South Carolina College of Pharmacy, Columbia, South Carolina

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Background. National guidelines support antibiotic prophylaxis for open fracture wounds. Our institution uses cefazolin based on fracture grade and comorbidities. The purpose of this study was to assess a modified adult antibiotic prophylaxis open fracture protocol (AOPF) which recommended weight-based cefazolin for low-grade fractures or ciprofloxacin plus vancomycin for high-grade fractures.

Methods. Adult patients with open fractures admitted to Palmetto Health Richland between January 2012 and December 2016 were screened for study inclusion. Exclusion criteria were receipt of antibiotics for reasons other than open fracture, death prior to wound closure, and local admission time >48 hours after time of injury. Compliance to all elements of AOPF was assessed. Clinical endpoints including open fracture infection rates, epidemiology, and drug-related adverse events were compared between pre-implementation (January 2012 – December 2012) and post-implementation period (November 2015 – December 2016; χ² and t-tests as appropriate were used to compare outcomes between groups.

Results. Following exclusions 189 patients were included in the analysis (90 pre- vs. 99 post-AOPF, respectively). Post-AOPF, a 17% (16/93) adherence rate to all AOPF elements was found. Appropriate agents were selected in 82.8% (77/93). The most common reasons for non-adherence were incorrect dosing and prolonged antibiotic duration. Infection rates were 23.3% (21/90) and 7.1% (7/99) in pre- vs. post-AOPF groups, respectively (P = 0.001). Infections primary caused by Gram-negative pathogens in pre-AOPF and Gram-negative organisms comprised 62% and 40% of open fracture site infections in pre- vs. post-AOPF groups, respectively. Incidence of acute kidney injury, Chlamydia difficile and staphylococcal infection rates were 33.3% (31/93) and 7.1% (7/99) in pre- vs. post-AOPF groups, respectively.

Conclusion. Overall adherence to all elements of the modified AOPF was low, yet the appropriate agent(s) was used in majority of cases. The modified AOPF was associated with a numerical decrease in infection rates post-open fracture and comparable AEs.

Disclosures. P. B. Bookstaver, Rock Pointe: Content Developer, Consulting fee

204. Pubic Osteomyelitis: Epidemiology and Factors Associated with Management Failure in Two French Reference Centers Tristan Ferry, MD, PhD1; Agathe Becker, MD, Florent Valour, MD, PhD2; Thomas Perpoint, MD1; Loïc Bousell, MD, PhD3; Alain Ruffion, MD, PhD3; Frederic Laurent, DPharm, PhD1; Eric Senneville, MD, PhD3; Christiana Chialiac, MD, PhD3; and Lyon BH study group1. 1Inserm 1111, UCELI, Hospices Civils de Lyon, Lyon, France, 2ID Department, Regional Reference Center for BJ, Hospices Civils de Lyon. Lyon, France, 3Hospices Civils de Lyon - Lyon Hosp. de la Croix-Rousse, Lyon, France, 4Hospices Civils de Lyon - Centre Hosp. de Lyon Sud-Elbeuf, Beneteau, France, 5Laboratory of Bacteriology, Regional Reference Center for BJ, Hospices Civils de Lyon, Lyon, France, 5Infectious Diseases, Dron Hospital, Tourcoing, France

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Background. Pubic osteomyelitis (PO) is a neglected bone and joint infections (BJI), as its management is still poorly codified. We aim to describe PO epidemiology and to look for factors associated with management failure.

Methods. We performed a retrospective cohort study in two French reference centers including patients with PO in 2010–2016. Treatment failure was defined by clinical (persistent or recurrence of clinical signs) and/or microbiological failure. Factors associated with treatment failure were determined by univariate Cox analysis (hazards ratio [HR] and 95% confidence interval calculations). Kaplan–Meier curve was compared between groups by log-rank test.

Results. Twenty-five patients were included over 13 years (median age 67 years; 19 men, median ASA score 3). Six (24%) had a PO from haematogenous origin. Those were all monomicrobial infection, due to S. aureus, mostly identified in young patients without comorbidities, especially in athletes. No surgery was required if no abscess or bone sequestrum were found. Nineteen patients (76%) had a post-operative chronic PO (developed from 1 month to 11 years after a pelvic surgery); 15 of them had history of pelvic cancer (60%). Twelve received radiotherapy at the site of infection (48%). Infection was polymicrobial in 68 % of cases, including 32 % of cases with multidrug-resistant pathogens. A clinical success was recorded in only 14 patients (56%). Treatment failure was always noticed in chronic post-operative forms. Potential risk factors for PO treatment failure pediatric and adult were similar: age at diagnosis, presence of comorbidities, CRP >100 mg/L, presence of multidrug-resistant pathogens.

Conclusion. This study highlights predominant chronic complex post-operative forms of PO. They are mostly plurimicrobial, sometimes associated with multi-drug resistance, occurring in fragile patients with pelvic cancer. It frequently leads to complex antibiotic therapy, with important risk of relapse. Aggressive surgical procedure with large bone resection is frequently required in patients who underwent pelvic radiotherapy.

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205. Reexamining BSA as a Preoperative Predictor of Risk of Prosthetic Joint Infection Celeste Russell, MPH1; Allina Nocon, MPH2; David Mayman, MD3; Geoffrey Westrich, MD4; Andy Miller, MD2; Barry Brause, MD2; and Michael Henry, MD2. 1Hosp. for Special Surgery, New York, New York, New York, 2Complex Joint Reconstruction Center, Hospital for Special Surgery, New York, New York, New York, 3Infectious Disease, Weill Cornell Medical College, New York, New York, 4Hosp. for Special Surgery, New York, New York, New York, Infectious Diseases, Hospital for Special Surgery, New York, New York

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Background. Prosthetic joint infection (PJI), a dreaded complication of arthroplasty, has been found to correlate with increasing body mass index (BMI) and body surface area (BSA). Recent data suggest that BSA may be a better predictive tool for assessing infection risk. We further evaluated this association in an orthopedic specialty hospital arthroplasty cohort to evaluate whether BSA is a predictor of prosthetic joint infection.

Methods. A retrospective cohort of hip and knee arthroplasty patients between 2009 and 2014 was identified through administrative hospital data using ICD-9 codes. Patients with a BMI <14 or >60 kg/m², height <142 or >200 cm, and weight <36 or >226 kg were excluded. BSA was calculated using the DuBois formula, and assessed both as a continuous and as a categorical variable. Univariate analyses were done using χ² tests and adjusted models were assessed using logistic regression.

Results. 17,859 knee and 18,128 hip patients were included. 1.1% of knees and 0.74% of hips were infected. [Mean BSA was 1.9 m² (10.2 m²). BSA was significantly associated with PJI in hips (p = 0.004), but not knees, when analyzed as a continuous variable in unadjusted models. However, this association lost its significance after adjusting for PJI risk factors. Additionally, when assessed as a categorical variable in a multivariable model, BSA in the highest quartile (>2.11) was not associated with PJI.

Conclusion. After evaluating BSA as a continuous and categorical variable, we failed to find an association between BSA and infection risk in THA or TKA. The impact of BSA decreased after multivariate adjustment. BSA may not be optimal as a predictor of preoperative risk.

Disclosures. All authors: No reported disclosures.

206. Variability in Management of Acute Osteoarticular Infections at a Children’s Hospital and Favorable Outcomes with Increasing Early Transition to Oral Therapy Nora Biary, MD, D1; Brian Wrotniak, PhD2; and Shamim Islam, MD, DTM&H3. 1University At Buffalo, State University of New York, Buffalo, New York, 2Emergency Medicine, University at Buffalo, Women and Children’s Hospital of Buffalo, Buffalo, New York, 3University at Buffalo, State University of New York, Buffalo, New York

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Background. To date, there is no established guideline for the treatment of acute pediatric osteoarticular infections (OA), and considerable variability in management exists amongst providers and across institutions. This study analyzed the recent management and clinical outcomes of OA at a children’s hospital.

Methods. Patients admitted with acute osteomyelitis and septic arthritis (OA) aged 2 months to 18 years at the Women & Children’s Hospital of Buffalo, between 1/1/2013 and 12/31/2016, were identified via billing databases. Patients with select comorbidities or >2 positive blood cultures were excluded. Patient demographics, anti-biotics used, culture results, time from intravenous (IV) to oral (PO) therapy, length of hospitalization (LOH), PICC (peripherally inserted central catheter) line use, and emergency room (ER) visits and readmissions within 6 months of discharge, were collected and analyzed. The study period was divided into 2-year groups, before and after 1/1/15, a juncture when a new provider began to promote early transition to oral therapy (through ID consults and informal discussion).

Results. Thirty-one patients were admitted during Time#1 (2013–14) and 43 during Time#2 (2015–16). Time to PO therapy was widely distributed (Figure 1) throughout the years, with 65% of Time#2 transition time <1 day (P = 0.02). Specifically for osteomyelitis, mean time to PO decreased from Time#1 to 7.62 from 11.5 to 8.3 days. Sixty-one percent of patients in Time#1 vs 25% in Time#2 (P = 0.01) received a PICC line, and PICC use was associated with 1.5 day greater LOH (P = 0.03). There was no relationship between time to PO and repeat ER visit/ readmission. Nearly 30% of patients in Time#1 had PICC or OA-related ER visits/ readmissions after discharge, while 0% did for Time#2 (P < 0.01).

Conclusion. Length of initial IV therapy and PICC line use continues to vary significantly for all elements of OA. In our children’s hospital, a shift towards an earlier transition to PO therapy has been adopted steadily – and prior to national and planned local guidelines – with a general decrease in LOH, duration of IV therapy, PICC line issues following discharge, and overall improved outcomes. Pediatric OA management represents an ideal focus for institutional quality and antibiotic stewardship efforts.