Association of Preoperative Vitamin D Levels and Severity of Fracture Among Orthopaedic Trauma Patients in a Single, Tertiary-Level Hospital *Christian Julius Patero Mendoza, MD*; *Mikhail Lew Perez Ver, MD*; *Andrew Gabriel Jacinto Tabberrah, MD*; *Mario Ver, MD St. Luke's Medical Center, Quezon City, Philippines*

Purpose: The high prevalence of hypovitaminosis D among trauma patients has been related to conflicting reports on risk of fractures, falls, nonunion, and poor clinical outcomes after surgery. There is limited evidence that ties up vitamin D levels with fracture severity with specific fractures. The primary objective of this study is to determine the association of preoperative level of vitamin D and the fracture severity among adult trauma patients. No previous study used the AO classification to stratify fracture severity and related this to vitamin D levels at time of injury.

Methods: All patients operatively treated for extremity fractures with preoperative vitamin D levels were reviewed. AO classification was assigned to all fractures present at injury. Demographics, presence of osteoporosis, mechanism of injury, and comorbidities were obtained for each patient. Follow-up clinical and radiographic data and fracture union were all recorded.

Results: 96 patients with 104 surgically treated extremity fractures were included. Patients presenting with more severe fractures were associated with lower levels of vitamin D and higher prevalence of hypovitaminosis D compared to patients with less severe fractures (χ 2 [4, N = 104] = 20.6, *P*<0.001). There was a strong, positive correlation between hypovitaminosis D and increasing fracture severity, which was statistically significant (rs [4] = 0.426, *P*<0.001). This association remains present in a subgroup analysis of patients without osteoporosis (*P* = 0.030), and in another subgroup of patients who sustained low-mechanism injuries (*P*<0.001). Union rate among our subjects is 97%.

Conclusion: Preoperative vitamin D level is associated with the severity of fracture as described in the AO classification sustained at the time of injury.

The FDA has stated that it is the responsibility of the physician to determine the FDA clearance status of each drug or medical device he or she wishes to use in clinical practice.