Nonopioid Pain Control Regimen After Open Reduction and Internal Fixation of Traumatic Fractures

NCT06113211

Status	RECRUITING
Phase	Phase 4
Sponsor	Henry Ford Health System
Enrollment	75 participants

Key Eligibility Criteria

Inclusion (1)

• All adult patients over age eighteen and scheduled for primary open reduction internal fixation following a traumatic fracture at Henry Ford Hospital (Detroit, Michigan, United States), and Henry Ford West Bloomfield Hospital (West Bloomfield, Michigan, United States) will be eligible for inclusion in this study. All patients will be met in our abulatory orthopedic clinics. All surgeries will be performed by a fellowship trained truama surgeons.

Exclusion (5)

• patients with a medical history of known allergies or intolerance to allergies or intolerance to Motrin, Lyrica, Tylenol, tramadol, Zanaflex

• substantial alcohol or drug abuse

• pregnancy

• history of narcotics within 6 months of surgery

• renal impairment, peptic ulcer disease, GI bleeding.

Locations (1 total)

Henry Ford Hospital System, Detroit, Michigan, United States