

Hematopoietic Stem Cell Dysfunction in the Elderly After Severe Injury

NCT02577731

Status	RECRUITING
Phase	Not Applicable
Sponsor	University of Florida
Enrollment	400 participants

Key Eligibility Criteria

Inclusion (14)

- Severe Trauma Population
- All adults (age ≥18 to 54)
- Blunt and/or penetrating trauma resulting in long bone or pelvic fractures requiring ORIF or closed reduction percutaneous pinning (CRPP).
- Blunt and/or penetrating trauma patient with either:
 - hemorrhagic shock defined by: i. systolic BP (SBP) < 90 mmHg or ii. mean arterial pressure < 65 mmHg or iii. base deficit (BD) ≥ 5 meq or iv. lactate ≥ 2
- ... and 9 more (see full listing online)

Exclusion (14)

- Patients not expected to survive greater than 48 hours.
- Prisoners.
- Pregnancy.
- Patients receiving chronic corticosteroids or immunosuppression therapies.
- Previous bone marrow transplantation.
- ... and 9 more (see full listing online)

Locations (1 total)

UF Health Shands Hospital at the University of Florida, Gainesville, Florida, United States

<https://clinicaltrials.gov/study/NCT02577731>

DISCLAIMER: This document is for informational purposes only and does not constitute medical advice. Always consult your healthcare provider before enrolling in any clinical trial. Information may not be up to date — verify details at [ClinicalTrials.gov](https://clinicaltrials.gov). Generated by [ClinicalTrialsFinder.org](https://clinicaltrialsfinder.org).