Treatment for Osteogenesis Imperfecta

OCR Number: OCR 6436

Description:

**PEDF and Derivative Peptide for Treatment of Osteogenesis Imperfecta**

- Absence of pigment epithelium-derived factor (PEDF) causes Osteogenesis Imperfecta (OI) in humans.
- OI Type VI is an autosomal recessive disease manifested by severely impaired bone mineralization and fractures in early childhood.
- PEDF is a regulator of MSC differentiation to the osteoblast lineage. PEDF modulates Wnt/β-catenin signaling to direct MSC fate toward osteoblasts. Restoration of PEDF in this PEDF KO mice corrected the bone phenotype (figure).
- Recently it was shown that PEDF treatment restores bone elasticity and reduces bone brittleness in the PEDF-KO mouse model (Unpublished data).

**Figure 1.** PEDF Treatment Increases Trabecular Bone Volume in a Mouse Model of OI Type VI. Micro-CT images of trabecular bone volume from three individual mice treated with vehicle or PEDF.

**PI:** Chuhan Chung

**Licensing Contact:** Hong Peng
hong.peng@yale.edu