

# Epimorphic Regeneration of Mesenchymal Stem Cells in Bone Marrow

Ken Muneoka\*

Department of Pharmacology, Texas A&M University, Texas, USA

## DESCRIPTION

Epimorphic regeneration, a remarkable process observed in certain organisms, allows the complete regeneration of complex tissues and structures after injury. This extraordinary ability has significant implications for human health, especially in relation to Mesenchymal Stem Cells (MSCs) found in bone marrow. MSCs have substantial regenerative potential due to their capacity to differentiate into various cell types and their role in tissue repair [1].

## Mesenchymal stem cells

MSCs are multipotent stromal cells primarily located in the bone marrow but also found in other tissues such as adipose tissue, umbilical cord blood, and dental pulp. These cells can differentiate into a range of cell types, including osteoblasts, chondrocytes, adipocytes, and myocytes, making them highly valuable for regenerative therapies [2]. MSCs play an important role in maintaining and repairing mesenchymal tissues. They support the homeostasis of the bone marrow microenvironment, aid in hematopoiesis, and participate in healing processes by migrating to injury sites and differentiating into necessary cell types. Their immunomodulatory properties are also essential in regulating the inflammatory response during tissue repair [3].

## Epimorphic regeneration

Epimorphic regeneration involves the ability of certain organisms to fully regenerate a lost or damaged body part, restoring its original structure and function [4]. This type of regeneration is seen in species such as salamanders and zebrafish, which can regenerate limbs, tails, and other complex structures. Upon injury, MSCs can dedifferentiate, reverting to a more primitive state. This is followed by proliferation, where the dedifferentiated cells rapidly divide to replace lost or damaged cells. Key signaling pathways involved in these processes include Wnt, Notch, and Hedgehog [5].

After proliferation, MSCs redifferentiate into specific cell types essential for tissue repair. In the bone marrow, this primarily involves differentiating into osteoblasts for bone formation and

chondrocytes for cartilage repair [6]. Additionally, MSCs remodel the Extra Cellular Matrix (ECM), which is crucial for providing structural support and regulating cell behavior during regeneration. This remodeling process ensures the proper environment for effective tissue regeneration and functional restoration [7].

The molecular pathways involved in MSC regeneration include some signaling pathways. The Wnt signaling pathway is critical for regulating MSC proliferation and differentiation, promoting osteogenic differentiation essential for bone formation and repair [8]. The Notch signaling pathway has a dual role in MSC regulation, controlling both proliferation and differentiation, and ensuring a balance between maintaining a pool of undifferentiated MSCs and promoting differentiation as needed. The Hedgehog pathway influences MSC proliferation and differentiation, particularly in bone repair, and interacts with other pathways like Wnt to coordinate tissue regeneration [9].

## Challenges

MSCs hold immense potential for therapeutic applications in regenerative medicine due to their ability to differentiate and their paracrine effects. They can potentially be used to treat various conditions including bone fractures, cartilage damage, and other degenerative diseases [10].

**Mechanistic insights:** Understanding the molecular and cellular mechanisms underlying MSC-mediated epimorphic regeneration is crucial. Key pathways and factors involved in MSC proliferation, differentiation, and signaling need to be elucidated.

**Clinical applications:** Translating the regenerative potential of MSCs into clinical applications involves addressing several challenges:

**Safety and efficacy:** Long-term studies are necessary to ensure the safety and efficacy of MSC-based therapies. Potential risks, such as unwanted differentiation or tumorigenesis, need to be thoroughly investigated [11].

**Correspondence to:** Ken Muneoka, Department of Pharmacology, Texas A&M University, Texas, USA, E-mail: ken1133@psu.edu

**Received:** 03-May-2024, Manuscript No. jcest-24-31619; **Editor assigned:** 06-May-2024, PreQC No. jcest-24-31619 (PQ); **Reviewed:** 20-May-2024, QC No. jcest-24-31619; **Revised:** 27-May-2024, Manuscript No. jcest-24-31619 (R); **Published:** 04-Jun-2024, DOI: 10.35248/2157-7013.24.15.456

**Citation:** Muneoka K (2024) Epimorphic Regeneration of Mesenchymal Stem Cells in Bone Marrow. J Cell Sci Therapy. 15:456.

**Copyright:** © 2024 Muneoka K. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

## CONCLUSION

The epimorphic regeneration of mesenchymal stem cells in bone marrow represents a promising area in regenerative medicine. Understanding the complex mechanisms that govern MSC behavior and their interactions within the bone marrow microenvironment is vital for developing effective therapies for bone and cartilage repair and harnessing their immunomodulatory properties.

## REFERENCES

1. Seifert AW, Muneoka K. The blastema and epimorphic regeneration in mammals. *Dev Biol.* 2018;433(2):190-199.
2. Londono R, Sun AX, Tuan RS, Lozito TP. Tissue repair and epimorphic regeneration: an overview. *Curr Pathobiol Rep.* 2018;6:61-69.
3. Stewart S, Rojas-Muñoz A, Belmonte JC. Bioelectricity and epimorphic regeneration. *Bioessays.* 2007;29(11):1133-1137.
4. Li C, Yang F, Sheppard A. Adult stem cells and mammalian epimorphic regeneration-insights from studying annual renewal of deer antlers. *Curr Stem Cell Res Ther.* 2009;4(3):237-251.
5. Carlson BM. Relationship between the tissue and epimorphic regeneration of muscles. *Am Zool.* 1970;10(2):175-186.
6. Easterling MR, Engbrecht KM, Crespi EJ. Endocrine regulation of epimorphic regeneration. *Endocrinology.* 2019;160(12):2969-2980.
7. Wong AY, Whited JL. Parallels between wound healing, epimorphic regeneration and solid tumors. *Development.* 2020;147(1):dev181636.
8. Mathew LK, Sengupta S, Franzosa JA, Perry J, La Du J, Andreasen EA, et al. Comparative expression profiling reveals an essential role for *raldh2* in epimorphic regeneration. *J Biol Chem.* 2009;284(48):33642-33653.
9. Ohgo S, Ichinose S, Yokota H, Sato-Maeda M, Shoji W, Wada N. Tissue regeneration during lower jaw restoration in zebrafish shows some features of epimorphic regeneration. *Dev Growth Differ.* 2019;61(7-8):419-430.
10. Laplace-Builhé B, Bahraoui S, Jorgensen C, Djouad F. From the basis of epimorphic regeneration to enhanced regenerative therapies. *Front cell dev biol.* 2021;8:605120.
11. Scaria SM, Frumm SM, Vikram EP, Easow SA, Sheth AH, Shamir ER, et al. Epimorphic regeneration in the mammalian tympanic membrane. *NPJ Regen Med.* 2023;8(1):58.