

Leveraging High Extracellular Vesicle Secretion for Cell Sorting and Optimization

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DESCRIPTION

Cell therapy is known as an effective therapy for many different kinds of diseases. It involves the transplantation of cells into a patient's body to replace or repair damaged cells or tissues. While cell therapy holds great potential, its efficacy can be influenced by several factors, including the quality and functionality of the transplanted cells. In recent years, Extracellular Vesicles (EVs) have gained significant attention for their role in cell-to-cell communication and their potential as therapeutic agents. This article explores the concept of optimizing cell therapy by sorting cells based on their high extracellular vesicle secretion. Extracellular vesicles are small membrane-bound particles released by cells into the extracellular space. They play a crucial role in intercellular communication by transferring proteins, lipids, and genetic material between cells. EVs have been found to have diverse functions, including immune modulation, tissue repair, and angiogenesis. Importantly, they can mediate the therapeutic effects of cell therapies by delivering bioactive molecules to target cells and tissues. To optimize cell therapy, it is crucial to identify and select cells with high extracellular vesicle secretion. Several techniques can be employed to achieve this goal, including flow cytometry, microfluidics, and nanotechnologies. These techniques allow for the isolation and enrichment of cells based on their EV secretion profiles. Flow cytometry is a widely used method that enables the identification and sorting of cells based on specific surface markers or fluorescent labels. By incorporating EV-specific markers, such as tetraspanins or specific Carrier Molecule, researchers can sort cells with higher EV secretion capacity. This sorting strategy ensures the selection of cells that are more likely to have a potent therapeutic effect. Microfluidic-based sorting platforms provide another avenue for isolating cells with high extracellular vesicle secretion. These platforms utilize microscale channels and precise fluid manipulation to separate cells based on various parameters, including size, density, and surface marker expression. By integrating EV-specific markers into

these platforms, it becomes possible to sort cells based on their EV secretion profiles efficiently. Furthermore, nanotechnologies, such as nanoscale sensors and sorting devices, offer exciting opportunities for isolating cells with high EV secretion. These technologies can provide real-time and sensitive detection of EVs, allowing for the identification and sorting of cells based on their EV secretion levels. By leveraging these advancements, researchers can obtain a highly purified population of cells with enhanced therapeutic potential. Sorting cells based on their high extracellular vesicle secretion can lead to several benefits in cell therapy. Firstly, it improves the therapeutic efficacy by selecting cells that possess a greater capacity for intercellular communication and tissue regeneration. These cells can release a higher quantity of EVs with the molecule with medicinal properties, enhancing their ability to modulate the immune response, promote angiogenesis, and stimulate tissue repair. Secondly, sorting cells based on EV secretion can enhance the safety profile of cell therapy. By selecting cells that secrete high levels of EVs, there is a reduced risk of potential adverse effects associated with direct cell transplantation. EVs, being smaller and less complex than whole cells, are less likely to cause immune reactions, tumorigenesis, or embolism. Therefore, using cells enriched for high EV secretion may mitigate safety concerns associated with traditional cell-based therapies. While sorting cells based on their EV secretion holds great promise for optimizing cell therapy, there are several challenges that need to be addressed. Firstly, standardization of sorting techniques and EV characterization methods is essential to ensure reproducibility and comparability of results across different research groups. This will facilitate the development of standardized protocols for EV-based cell therapy. Moreover, further research is needed to understand the functional heterogeneity of EVs and the specific Carrier Molecule responsible for their therapeutic effects. By identifying and characterizing these Carrier Molecule, it may be possible to engineer cells with enhanced EV secretion and targeting specific therapeutic targets with the results

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CONCLUSION

Optimizing cell therapy by sorting cells with high extracellular vesicle secretion offers tremendous potential for enhancing the therapeutic efficacy and safety of cell based treatments. By selecting cells that secrete a greater quantity of EVs, we can maximize their potential to communicate with target cells, deliver

bioactive molecules, and promote tissue repair. However, further research is needed to standardize sorting techniques, characterize EV carrier, and establish clinical protocols. With continued advancements in technology and deeper understanding of EV biology, We can realize the full potential of cell therapy and revolutionize the treatment of several medical conditions.