REVIEW

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Mitochondrial dysfunction in neurodegenerative disorders: Potential therapeutic application of mitochondrial transfer to central nervous system-residing cells

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Abstract

Mitochondrial dysfunction is reiteratively involved in the pathogenesis of diverse neurodegenerative diseases. Current in vitro and in vivo approaches support that mitochondrial dysfunction is branded by several molecular and cellular defects, whose impact at different levels including the calcium and iron homeostasis, energetic balance and/or oxidative stress, makes it difficult to resolve them collectively given their multifactorial nature. Mitochondrial transfer offers an overall solution since it contains the replacement of damage mitochondria by healthy units. Therefore, this review provides an introducing view on the structure and energy-related functions of mitochondria as well as their dynamics. In turn, we summarize current knowledge on how these features are desegulated in different neurodegenerative diseases, including frontotemporal dementia, multiple sclerosis, amyotrophic lateral sclerosis, Friedreich ataxia, Alzheimer's disease, Parkinson's disease, and Huntington's disease. Finally, we analyzed current advances in mitochondrial transfer between diverse cell types that actively participate in neurodegenerative processes, and how they might be projected toward developing novel therapeutic strategies.

Felipe Bustamente Barrientos and Noymar Lugue-Campos equally contribute as first authors in this manuscript.

Ania Maria Vega-Lettler Since July 17th 2021.

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Table of Contents Mitochondrial Dysfunction In Neurodegenerative Disorders

- 1. Understanding the eBook Mitochondrial Dysfunction In Neurodegenerative Disorders
 - The Rise of Digital Reading Mitochondrial Dysfunction In Neurodegenerative Disorders
 - Advantages of eBooks Over Traditional Books
- 2. Identifying Mitochondrial Dysfunction In Neurodegenerative Disorders
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
- 3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Mitochondrial Dysfunction In Neurodegenerative Disorders
 - User-Friendly Interface
- 4. Exploring eBook Recommendations from Mitochondrial Dysfunction In Neurodegenerative Disorders
 - Personalized Recommendations
 - Mitochondrial Dysfunction In Neurodegenerative Disorders User Reviews and Ratings
 - Mitochondrial Dysfunction In Neurodegenerative Disorders and Bestseller Lists

- 5. Accessing Mitochondrial Dysfunction In Neurodegenerative Disorders Free and Paid eBooks
 - Mitochondrial Dysfunction In Neurodegenerative Disorders Public Domain eBooks
 - Mitochondrial Dysfunction In Neurodegenerative Disorders eBook Subscription Services
 - Mitochondrial Dysfunction In Neurodegenerative Disorders Budget-Friendly Options
- 6. Navigating Mitochondrial Dysfunction In Neurodegenerative Disorders eBook Formats
 - o ePub, PDF, MOBI, and More
 - Mitochondrial Dysfunction In Neurodegenerative Disorders Compatibility with Devices
 - Mitochondrial Dysfunction In Neurodegenerative Disorders Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Mitochondrial Dysfunction In Neurodegenerative Disorders
 - Highlighting and Note-Taking Mitochondrial Dysfunction In Neurodegenerative Disorders
 - Interactive Elements Mitochondrial Dysfunction In Neurodegenerative Disorders
- 8. Staying Engaged with Mitochondrial Dysfunction In Neurodegenerative Disorders
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Mitochondrial Dysfunction In Neurodegenerative Disorders
- 9. Balancing eBooks and Physical Books Mitochondrial Dysfunction In Neurodegenerative Disorders
 - ∘ Benefits of a Digital Library
 - Creating a Diverse Reading Collection Mitochondrial Dysfunction In Neurodegenerative Disorders
- 10. Overcoming Reading Challenges
 - o Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Mitochondrial Dysfunction In Neurodegenerative Disorders
 - Setting Reading Goals Mitochondrial Dysfunction In Neurodegenerative Disorders
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Mitochondrial Dysfunction In Neurodegenerative Disorders
 - Fact-Checking eBook Content of Mitochondrial Dysfunction In Neurodegenerative Disorders
 - Distinguishing Credible Sources
- 13. Promoting Lifelong Learning

- Utilizing eBooks for Skill Development
- Exploring Educational eBooks
- 14. Embracing eBook Trends
 - Integration of Multimedia Elements
 - Interactive and Gamified eBooks

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