

DETECTION OF MITOCHONDRIAL DISEASES%0A

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These data confirm the specificity of mitochondrial O₂ detection by mitoSOX and illustrate that DHE detects cytoplasmic O₂, but not mitochondrial O₂. Separation of O₂ produced in either the mitochondria or the cytoplasm is therefore possible by HPLC measurements of 2-OH-E⁺ or 2-OH-Mito-E⁺.

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We thank all authors for their patience and cooperation that have made it possible to produce this unique collection of papers representing current knowledge on detection of mitochondrial causes of diseases. We especially thank Prof. N.S. Dhalla for making it possible to publish these contributions together in this focused issue and also as a hard-cover book.

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The MitoLight Apoptosis Detection Kit utilizes a lipophilic cation, termed as MitoLight, as a mitochondrial activity marker. MitoLight is a mitochondrial dye that stains mitochondria in living cells in a membrane potential-dependent fashion.

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Detection of Mitochondrial Diseases Edited by FRANK NORBERT GELLERICH Martin Luther Universität Halle-Wittenberg Neurologische Klinik und Poliklinik

Testing for Mitochondrial Disease Disorders ...

Testing for Mitochondrial Disease Disorders

Mitochondrial disorders are disorders in which the body lacks the ability to produce adequate energy to perform all of its functions. The body systems most dependent on this energy are the brain, muscle, heart, kidneys and liver.

Mitochondrial disease - Wikipedia

Mitochondrial diseases are a group of disorders caused by dysfunctional mitochondria, the organelles that generate energy for the cell. Mitochondria are found in every cell of the human body except red blood cells, and convert the energy of food molecules into the ATP that powers most cell functions.

Types of Mitochondrial Disease UMDF

KSS is a slowly progressive multi-system mitochondrial disease that often begins with drooping of the eyelids (ptosis). Other eye muscles eventually become involved, resulting in paralysis of eye movement. Degeneration of the retina usually causes difficulty seeing in dimly lit environments.

Detection of mitochondrial diseases (Book, 1997)

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Part 1: Noninvasive Detection of Mitochondrial Function. Part 2: Bioenergetic Investigation of Isolated Mitochondria, Skinned Muscle Fibers and Cells. Part 3: Mitochondrial Transition Pore, Radicals and Diseases. Part 4: Mitochondrial Genome and Diseases. Part 5: Ageing, Mitochondria and Diseases.

Detection of mitochondrial fission with orientation ...

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Mitochondrial diseases | Nature Reviews Disease Primers

Mitochondrial diseases are a group of genetic disorders that are characterized by defects in oxidative phosphorylation and caused by mutations in genes in the nuclear DNA (nDNA) and mitochondrial