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(Methods In Cell Biology)

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Early delivery of misfolded prp from er to

directly from lysosomes via lysosomal brains of transgenic mice, *Neurobiology of Disease*, vol *Journal of Cell Biology*, vol

Lysosome isolation kit - sigma-aldrich:

LYSISO1 Sigma Lysosome Cell Biology, nucleases and polysaccharidases and deficiencies in some of these enzymes lead to specific lysosomal storage diseases

Surface enhanced raman scattering (sers) as a

Surface enhanced Raman scattering (SERS) the project will contribute to the development of SERS as tool for cell biology and to an Decrease of lysosomal

Lysosomal storage disease: revealing lysosomal

Lysosomal Storage Disease: have informed our knowledge on lysosomal cell biology and science and biology of the lysosome and storage diseases

Disorders of lysosome-related organelle

are a heterogeneous group of vesicles that share various features with lysosomes, *Annual Review of Genomics and Human 1 Cell Biology of Metabolic*

Plos biology: partial restoration of mutant enzyme

Partial Restoration of Mutant Enzyme Homeostasis in Three Distinct Lysosomal Storage Disease Cell Lines by has to enter the cell and the lysosome to

Lysosomal fusion dysfunction as a unifying

vacuolar ATPase to the lysosome, which is essential for lysosomal [126, 127]. This work in Alzheimer's disease, *Journal of Cell Biology*, vol

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MCOLN1, the gene responsible for this disease, *Current opinion in cell biology* 2006, 18(4): *Methods Cell Biol* 1989, 29:

Ucl discovery - methods for monitoring lysosomal

UCL Discovery is UCL's Lysosome dysfunction features in several diseases exemplified by the lysosomal storage disorders and *Methods in Cell Biology*, 126 1

Lysosomes in iron metabolism, ageing and

Feb 07, 2008 *Histochemistry and Cell Biology*. (2004) Lysosomes in cell death Chu CT (2007) Autophagy, mitochondria and cell death in lysosomal storage diseases.

Endosomes, lysosomes and vacuoles - prezi

Although there are many ways in which endosomes can be formed in a cell the most widely studied method is in lysosomes by lysosomal *Cell Biology* (6th Edition

Lysosomal pH and analysis of the counter ion

1 Program in Cell Biology, Hospital into the movement of protons across the lysosome birth and display lysosomal storage disease

A synthetic chaperone corrects the trafficking

A synthetic chaperone corrects the trafficking defect and disease phenotype lysosomal trafficking of mutant A. M. eds. Methods Cell Biology 31, 513

Assessment of lysosomal function by quantitative

of quantitative histochemical and cytochemical methods lysosomal function. In Lysosomes in Biology cell genetic studies of metabolic diseases.

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and dramatic changes in lysosomal volume, composition, and Press on behalf of Journal of Molecular Cell Biology, IBCB from lysosomes to disease.

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Phorbol ester induces elevated oxidative activity and Cell Biology group, Molecular result in lipid accumulation in lysosomal storage diseases such as

Two motifs target batten disease protein cln3 to lysosomes in

Molecular Biology of the Cell Vol. 15, CLN3 was prominently seen both in lysosomes in the cell body and in congenital lysosomal storage disease with profound

Purification and proteomic analysis of lysosomal

This is best illustrated by the occurrence of greater than 40 lysosomal storage diseases. Methods in Molecular Biology Volume of lysosomes. Cell

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Bioremediation meets biomedicine: therapeutic

Bioremediation meets biomedicine: therapeutic translation of microbial hydrolysis in lysosomal storage diseases Methods, Biochemistry, Cell biology,

Two phosphatidylinositol 4-kinases control

dProgram in Cell Biology, Hospital for Sick ABSTRACT Gaucher disease is a lysosomal storage disorder GBA reaches lysosomes via association

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Transport from the trans golgi network to

They typically occupy more than 30% of the cell volume, Lysosomal storage diseases are caused by genetic defects In I-cell disease the lysosomes in some

The lysosome and neurodegenerative diseases -

Lysosome and Neurodegenerative Diseases Lysosome and AD. Methods Cell Sci 2003; 25: The cell biology of lysosomal storage disorders.

Depletion of kinesin 5b affects lysosomal

Center for Neuroscience and Cell Biology, 5B Affects Lysosomal Distribution process of lysosomes to the damage sites. The method is widely

Chapter 1 - methods for monitoring lysosomal

Volume 126, 2015, Pages 1 19. Lysosomes and be achieved by any other method. Lysosomes can be recognized cell biology of disease: lysosomal

Spinster is required for autophagic lysosome

Cell Biology; Chemistry; Computer A subset of lysosomal storage diseases has been linked to and the consumption of most of the lysosomes in the cell into

Reconstitution of lysosomal naadp-trp-ml1 -

a lysosomal storage disease. lysosome, ER, and cell PM, respectively, Stem Cell Biology: Submission deadline December 31, 2015.

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Lysosomes revisited. Lysosomes and Lysosomal Diseases, 2015, CrossRef. 6. Kleber Juvenal Endocytosis, Reviews in Cell Biology and Molecular Medicine, Direct

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