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Lysosomes and the connective tissue diseases

MALLUCCI L. HISTOCHEMICAL STUDIES OF LYSOSOMES AND LYSOSOMAL ENZYMES IN VIRUS-INFECTED Watts RW, Bitensky L, Chayen J Review Animal models for lysosomal

A Historical Perspective of the Glycosphingolipids

Two other major groups of lysosomal storage diseases (reviewed by Watts A historical perspective of the glycosphingolipids other lysosomal storage diseases

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Lysosomal storage diseases : biochemical and

Lysosomal storage diseases : biochemical and clinical aspects. Watts, R.W.E. Lysosomal storage diseases. London ; Philadelphia : Taylor & Francis, 1986

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Induced pluripotent stem cells derived from mouse

Most lysosomal storage diseases (LSDs) are life-threatening genetic diseases. The pathogenesis of these diseases is poorly understood. Induced pluripotent stem (iPS

Saposins A, B, C, and D in Plasma of Patients with

1 Lysosomal Diseases Research Unit, An introduction to the basic science and biology of the lysosome and storage diseases. Spellacy E, Watts RW.

Patent WO2001019955A2 - Lysosomal targeting

Lysosomal Storage diseases have been studied extensively and the enzymes (or lack thereof) The search identified a clone HS316G12(gi 4495019)

Inhibition of substrate synthesis as a strategy

for glycolipid lysosomal storage disease therapy required caused extensive inhibition of GI tract disaccharidases and resulted in osmotic diarrhoea.

Degenerative CNS Disease: Lysosomal Storage

Degenerative CNS Disease: Lysosomal Storage Disorders; GI; GU; Hematology / Oncology; Infectious Disease. Bacterial Infections;

The lysosome: from waste bag to potential

In addition, lysosomal contribution to diseases, including lysosomal storage disorders, neurodegenerative disorders, cancer Watts, 2012; Zhao, 2012).

Lysosomal metabolism of glycoproteins - Oxford

Defects in these pathways lead to lysosomal storage diseases. (Swainson and Watts, 1995) of extracellular proteins, and endocytosis by caveolae.

Lysosomal Storage Diseases in Adults -

Lysosomal Storage Diseases in Adults 761 Residual Accepted April 21, 1994 Key words: Lysosomes -Storage disease -Gangliosidosis -Gaucher GI Hurowitz, JM

Lysosomal storage disease: Gene therapy on both

Lysosomal storage diseases (LSDs) and their therapies. R.J. Watts, M.S. Dennis; Bispecific antibodies for delivery into the brain. Curr. Opin. Chem. Biol., 17

The Lysosome and Lysosomal Storage Disorders LSD -

The Lysosome and Lysosomal Storage Disorders (LSD). io.uwinnipeg.ca/~simmons/1115/cm1503/Image110.gi . Lysosomal Storage Diseases

T. Andrew Burrow , MD - Cincinnati Children's

T. Andrew Burrow, MD, is a faculty member of the Division of Human Genetics at Cincinnati Children's Hospital Medical Center The Lysosomal Storage Diseases. In:

Laboratory Diagnostics of Lysosomal Storage

tools in the diagnostics of inherited lysosomal storage diseases in future. References: 1) Watts, RWE The main groups of the inherited lysosomal storage diseases

Fabry disease - Wikipedia, the free encyclopedia

is a rare genetic lysosomal storage disease, GI tract pain is likely caused by accumulation of lipids in the small vasculature of the GI tract which

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