

**sphingolipids in disease pdf**

Fabry disease, also known as Anderson's Fabry disease, is a rare genetic disease that can affect many parts of the body including the kidneys, heart, and skin. Fabry disease is one of a group of conditions known as lysosomal storage diseases. The genetic mutation that causes Fabry disease interferes with the function of an enzyme which processes biomolecules known as sphingolipids, leading to ...

**Fabry disease - Wikipedia**

Tay-Sachs disease is a genetic disorder that results in the destruction of nerve cells in the brain and spinal cord. The most common type, known as infantile Tay-Sachs disease, becomes apparent around three to six months of age with the baby losing the ability to turn over, sit, or crawl. This is then followed by seizures, hearing loss, and inability to move.

**Tay-Sachs disease - Wikipedia**

5 5. POSTURAL INSTABILITY IN PARKINSON'S DISEASE: PERSPECTIVES OF EARLY DIAGNOSIS. EV Gracheva, AV Kudrevatykh, TV Sergeev, IV Miliukhina, Institute of Experimental Medicine RAS, St.

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