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Sphingolipidoses are a class of lipid storage disorders relating to sphingolipid metabolism. The main members of this group are Niemann-Pick disease, Fabry disease, Krabbe disease, Gaucher disease, Tay-Sachs disease and metachromatic leukodystrophy. They are generally inherited in an autosomal recessive fashion, but

notably Fabry disease is X-linked recessive. Taken together, sphingolipidoses have an incidence of approximately 1 in 10,000, but substantially more in certain ...

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