

Changing Treatment Paradigms to Optimize Outcomes for Patients With Type 1 Gaucher Disease: A Case-Based Approach

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Dr Pastores has extensive clinical and research experience in the diagnosis and management of lysosomal storage disorders and inherited diseases that primarily afflict individuals of Ashkenazi Jewish ancestry. He has also been engaged in the development and testing of treatment for Gaucher disease, Hurler syndrome and Morquio syndrome, Pompe disease, and a late (adult)-onset form of Tay–Sachs disease (GM2 gangliosidosis). His current interest lies in the molecular genetics of inherited diseases that involve the nervous system.