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Ganglioside sialidase and glycoprotein N-acetylgalactosaminyltransferase. II. Purification and properties of human brain ganglioside sialidase. Ganglioside sialidase (GM2/GD2 neuraminidase) was purified 1421-fold from human brain by gel filtration and ion-exchange chromatography. The activity was detected in postmortem brains of patients with inherited and acquired neurological disorders. The enzyme was a sialidase requiring Mn<sup>2+</sup> or Mg<sup>2+</sup> for activity, which was inhibited by high concentrations of lysosomal protease. The enzyme catalyzed the hydrolysis of gangliosides GM2, GD2 and GT1b, but neither N-acetylneuraminic acid- nor N-glycolylneuraminic acid-containing gangliosides. The purified enzyme appeared to be homogeneous based on its single band on dodecylsulfate-polyacrylamide gel electrophoresis. The isoelectric point (pI) of the enzyme was 4.4-4.5. The molecular mass, determined by gel filtration in the presence of sodium dodecylsulfate, was 135 kDa. The enzyme activity was inhibited by mercury, lead, silver, cobalt, cadmium, nickel, copper, zinc, and antimony salts as well as sulfhydryl group inhibitors. The amino acid composition of the enzyme was similar to that of the lysosomal sialidase purified from rat liver, i.e. 53% glutamic acid, 5% as c6a93da74d

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