Lysosomal storage diseases are a group of about 50 rare inherited metabolic disorders that result from defects in lysosomal function.

A problem of management of patients with lysosomal storage diseases in own experience with over 100 children with such diseases has been discussed. Symptomatic therapy of carpal tunnel syndrome, Pudenz valves, splenectomies, plasty of hernia, locomotive rehabilitation and various forms of cooperation with patients' families have been used in the treatment. An attempt of the treatment of the storage diseases with implantation of fetal membranes has been undertaken in view of the fact, that such membranes are the source of deficit enzyme [1)].

1)
Tylki-Szymańska A. [Treatment of patients with lysosomal storage diseases]. Pol Tyg Lek. 1992 May 18-25;47(20-21):450-2. Polish. PubMed PMID: 1437765.