:: Ketotic hypoglycemia

Diseases:

- Hyperglycerolemia (glycerol kinase deficiency)
- Hepatic glycogen synthase deficiency (glycogen storage disease type 0a, GSD type 0a)

The British Inherited Metabolic Disease Group (BIMDG) has published on its website guidelines for the emergency management of patients with inherited metabolic disorders.

Here are the ones for the management of a
Ketotic hypoglycemia in children with a hyperglycerolemia or a hepatic glycogen synthase deficiency.

- See children BIMDG guidelines (last reviewed in Feb 2012)

Further Information: see the Orphanet Abstracts for hyperglycerolemia and hepatic glycogen synthase deficiency.