:: Hyperammonemia due to citrullinemia or argininosuccinic aciduria

Diseases:

- citrullinemia type 1(argininosuccinate synthetase deficiency)
- Argininosuccinic aciduria (argininoscuccinase deficiency)



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 Hyperammonaemia due to citrullinemia type 1 or argininosuccinic aciduria in children and adults.

- ▶ See children BIMDG guidelines (last reviewed in Feb 2012)
- ▶ See <u>adults BIMDG guidelines</u> (last reviewed in 2012)

Further Information: see the Orphanet Abstracts for citrullinemia and argininosuccinic aciduria.