



Urea cycle disorders

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Most patients with urea cycle disorders who present as neonates, do so with deteriorating feeding, drowsiness and tachypnoea, following a short initial period when they appear well. The plasma ammonia should be measured at the same time as the septic screen in such patients. Ammonia levels above 200 $\mu\text{mol/l}$ are usually caused by inherited metabolic diseases and it is essential to make a diagnosis for genetic counselling, even if the patients die. The aim of treatment is to lower the ammonia concentrations as fast as possible. Sodium benzoate, sodium phenylbutyrate and arginine can exploit alternative pathways for the elimination of nitrogen but haemodialysis or haemofiltration should be instituted if ammonia concentrations are $>500 \mu\text{mol/l}$ or if they do not fall promptly. Long-term management involves drugs, dietary protein restriction and use of an emergency regimen during illness. Severe hyperammonaemia is usually associated with irreversible neurological damage, particularly if levels have been above 800 $\mu\text{mol/l}$ for >24 hours, and the option of withdrawing treatment should be discussed with the family.

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Introduction

The urea cycle is the final common pathway for the excretion of waste nitrogen in mammals (Fig. 1). Urea has low toxicity even at high concentrations, in contrast to its precursors, particularly ammonia. Urea cycle defects presenting in the neonatal period are usually associated with severe and rapidly worsening hyperammonaemia. This is a major emergency in neonates. Early recognition and aggressive treatment are essential to achieve a good outcome. Even with prompt intervention, the prognosis is poor for patients who present with symptoms in the neonatal period. A number of other disorders besides urea cycle defects can cause severe hyperammonaemia but most are inherited and every effort must be made to establish a diagnosis.

Clinical presentation

Patients with urea cycle disorders commonly present in the neonatal period but the symptoms and signs are not specific. Most of these babies are of normal birth weight and are initially healthy, but then after a short interval, that can be less than 24 hours, they become unwell. Common early symptoms are poor feeding, vomiting, lethargy and/or irritability and tachypnoea. The initial working diagnosis is almost invariably sepsis. Rather characteristically, these babies often have a mild but transient respiratory alkalosis at this stage that can be a useful diagnostic clue as there are few other causes in a baby not on a ventilator. They may also have neuro-muscular irritability and stridor but all these symptoms are usually only transient as generally the patients deteriorate rapidly. They develop more obvious neurological and autonomic problems, including changes of tone with loss of normal reflexes, vasomotor instability and hypothermia, apnoea and fits. The baby may soon become totally unresponsive and require full

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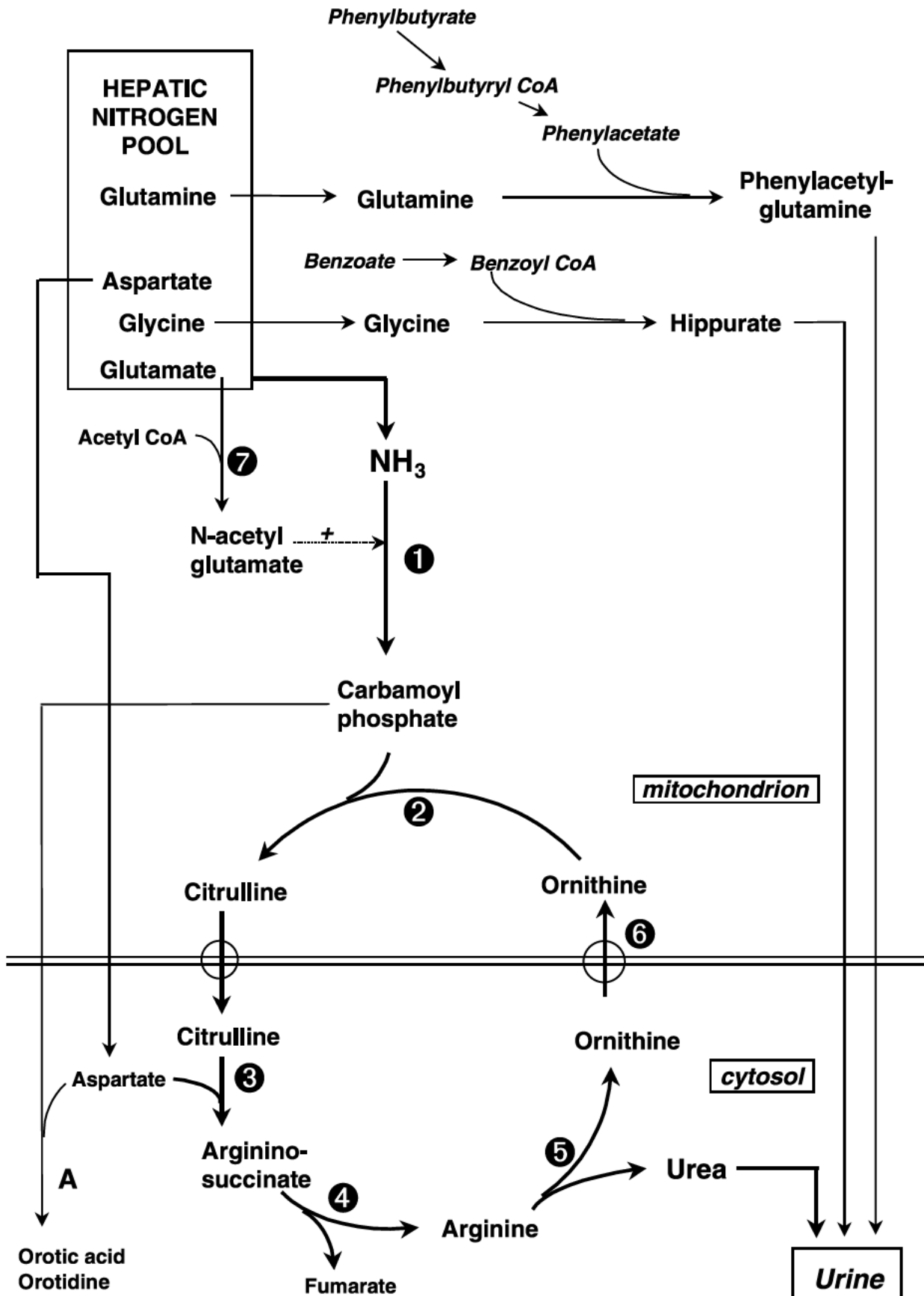


Figure 1. Pathways for the disposal of waste nitrogen. The urea cycle and alternative pathways of nitrogen excretion. Steps in the urea cycle: 1. Carbamoyl phosphate synthetase; 2. Ornithine transcarbamoylase; 3. Argininosuccinate synthetase; 4. Argininosuccinate lyase; 5. Arginase; 6. Mitochondrial ornithine carrier; 7. N-acetyl glutamate synthetase. A: Allouuridinol

intensive care. The babies then often develop a wide range of secondary complications such as disordered liver function that obscures the primary condition. Untreated, most babies will die, often with complications such as cerebral or pulmonary haemorrhage. Some survive but they are invariably handicapped, usually severely.

Patients with arginase deficiency usually present after the neonatal period with spasticity in the legs and developmental delay but seldom have symptomatic hyperammonaemia. On the other hand, neonatal hyperammonaemia is well recognized in patients with defects of the mitochondrial ornithine transporter, an essential component of the urea cycle (Hyperornithinaemia, Hyperammonaemia, Homocitrullinuria syndrome). Severe neonatal hyperammonaemia also occurs in patients with ornithine aminotransferase deficiency [1,2], a defect that more commonly presents in adults with cataracts and gyrate atrophy of the choroid and retina.

Differential diagnosis

The differential diagnosis of hyperammonaemia is wide and is summarized in Table 1. The most common differential diagnoses of severe hyperammonaemia are organic acidaemias, particularly propionic and methylmalonic acidaemia. It is important to recognize that patients with these disorders may have marked hyperammonaemia with a respiratory alkalosis without acidosis or ketosis. Transient hyperammonaemia of the newborn (THAN) is an ill-understood condition, possibly related to immaturity of liver metabolism or hepatic vascular disease. Plasma ammonia levels may be very high initially but no underlying metabolic disease is found. Although babies with THAN are often born prematurely with early onset of symptoms [3], it may be difficult to distinguish between urea cycle disorders and this disorder on clinical grounds. The incidence of THAN appears to have been falling over recent years in many centres around the world. Less severe hyperammonaemia is common, both in other metabolic disorders and acquired illness such as sepsis and perinatal asphyxia. Babies with systemic herpes simplex, particularly involving the liver, may have marked hyperammonaemia without obvious signs.

Investigations

Routine tests are not helpful for establishing the diagnosis of hyperammonaemia. The most

Table 1. Differential diagnosis of hyperammonaemia

Inherited disorders

Urea cycle enzyme defects

- Carbamoyl phosphate synthetase deficiency
 - Ornithine transcarbamoylase deficiency
 - Argininosuccinate synthetase deficiency (Citrullinaemia)
 - Argininosuccinate lyase deficiency (Argininosuccinic aciduria)
 - Arginase deficiency
 - N-acetylglutamate synthetase deficiency
- #### Transport defects of urea cycle intermediates
- Lysinuric protein intolerance
 - Hyperammonaemia – hyperornithinaemia – homocitrullinuria syndrome

Organic acidaemias

- Propionic acidaemia
- Methylmalonic acidaemia and other organic acidaemias

Fatty acid oxidation disorders

- Medium chain acyl-CoA dehydrogenase deficiency
- Systemic carnitine deficiency
- Long chain fatty acid oxidation defects and other related disorders

Other inborn errors

- Pyruvate carboxylase deficiency (neonatal form)
- Ornithine aminotransferase deficiency (neonates)

Acquired disorders

- Transient hyperammonaemia of the newborn
- Any severe systemic illness
- Herpes simplex – systemic infection
- Liver failure (rare in neonates)
- Infection with urease positive bacteria (if urinary tract stasis)

important diagnostic test in urea cycle disorders is measurement of the plasma ammonia concentration. In healthy neonates plasma ammonia is normally less than 65 $\mu\text{mol/l}$ [4], but may be raised as a result of a high protein intake, difficult venepuncture or a haemolysed blood sample. In sick neonates (for example, those with sepsis or perinatal asphyxia), plasma ammonia concentrations may increase up to 180 $\mu\text{mol/l}$. Patients with inborn errors presenting in the newborn period usually have concentrations greater than 200 $\mu\text{mol/l}$, often very much greater.

Ammonia levels can rise rapidly in patients with urea cycle disorders. Thus, plasma ammonia measurement should be repeated after a few hours, even if it is only modestly elevated.

In cases of significant hyperammonaemia, the following investigations should be performed immediately:

- Blood pH and gases
- Plasma urea and creatinine, electrolytes, glucose

Table 2. Diagnostic tests in urea cycle defects

Disorder	Alternative names	Plasma amino acid concentrations	Urine orotic acid	Tissue for enzyme diagnosis	Genetics (chromosome localization)
Carbamoyl phosphate synthetase deficiency	CPS deficiency	↑glutamine ↑alanine ↓citrulline ↓arginine	N	Liver	AR (chromosome 2p)
Ornithine transcarbamoylase deficiency	OTC deficiency	↑glutamine ↑alanine ↓citrulline ↓arginine	↑↑	Liver	X-linked (Xp21.1)
Argininosuccinic acid synthetase deficiency	Citrullinaemia	↑↑citrulline ↓arginine	↑	Liver/Fibroblasts	AR (chromosome 9q)
Argininosuccinic acid lyase deficiency	Argininosuccinic aciduria (ASA)	↑citrulline ↑argininosuccinic acid ↓arginine	↑	RBC/Liver/Fibroblasts	AR (chromosome 7q)
Arginase deficiency	Hyperargininaemia	↑arginine	↑	RBC/Liver	AR (chromosome 6q)
N-acetylglutamate synthetase deficiency	NAGS deficiency	↑glutamine ↑alanine	N	Liver	AR (not confirmed)

AR: autosomal recessive; RBC: red blood cells; N: normal.

- Liver function tests and clotting studies
- Plasma amino acids
- Urine organic acids, orotic acid and amino acids
- Plasma free and acylcarnitines

The plasma amino acids and urine organic acids are very urgent.

In all urea cycle disorders there is accumulation of glutamine and alanine. There are also increased concentrations of the amino acids immediately proximal to the block in the metabolic pathway and decreased concentrations of those beyond the block (Fig. 1). Thus, in citrullinaemia, argininosuccinic aciduria (ASA) and arginase deficiency, the plasma amino acids are usually diagnostic (Table 2).

Orotic acid and orotidine are excreted in excess in the urine if there is a metabolic block distal to the formation of carbamoyl phosphate. In these disorders carbamoyl phosphate accumulates, leaves the mitochondrion and enters the pathway for the de novo synthesis of pyrimidines in the cytosol (Fig. 1). Measurement of urinary orotic acid is particularly helpful for distinguishing ornithine transcarbamoylase (OTC) deficiency from carbamoyl phosphate synthetase (CPS) or N-acetyl glutamate synthetase (NAGS) deficiencies.

Diagnoses can generally be confirmed by measuring the enzyme activity in an appropriate tissue (Table 2). This is the only way to distinguish

between CPS and NAGS deficiencies. Assays of CPS are well-established but measurement of NAGS activity is not straightforward. Patients with NAGS deficiency generally show a clinical response to N-carbamyl glutamate, an orally active analogue of N-acetyl glutamate, but this is unreliable for diagnosis because a response is also seen in some patients with CPS deficiency [5].

Other investigations will detect complications. In the late stages of hyperammonaemia patients may have marked disturbances of liver function with disordered clotting, renal failure and hypocalcaemia. In the later stages of hyperammonaemic encephalopathy, brain imaging may show cerebral oedema or intracranial haemorrhage.

If the patient seems likely to die it is essential to collect the appropriate specimens, since otherwise the diagnosis cannot be confirmed:

- Plasma (heparinized, separated and deep frozen)
- Blood spots on filter paper for acylcarnitines
- Urine (deep frozen in a plain tube)
- Blood for DNA (anticoagulated with EDTA and deep frozen)
- Skin for fibroblast culture taken with sterile precautions into medium and stored at 4–8°C, not frozen
- Liver, snap frozen for enzymology

Table 3. The emergency treatment of neonatal hyperammonaemia

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- **General neonatal supportive care** e.g. ventilation (particularly prior to transfer) treatment of sepsis, seizures etc.
 - **Stop protein intake**
 - **Give a high energy intake**
 - either (a) oral
 - (i) 10% soluble glucose polymer (higher concentrations may be given if they are tolerated)
 - (ii) protein free formula (80056 (Mead Johnson); Duocal (SHS Ltd))
 - or (b) intravenously
 - (i) 10% glucose by peripheral infusion
 - (ii) 10–25% glucose by central venous line
- Fluid volumes may be restricted if there is concern about cerebral oedema
- **Alternative pathways for nitrogen excretion**
 - Sodium benzoate up to 500 mg/kg/day – oral or intravenously
 - Sodium phenylbutyrate up to 600 mg/kg/day
 - L-arginine
 - In citrullinaemia and ASA – up to 700 mg/kg/day
 - In OTC deficiency and CPS deficiency – up to 150 mg/kg/day
 - L-citrulline
 - In OTC deficiency and CPS deficiency up to 170 mg/kg/day instead of arginine
- For the emergency treatment of hyperammonaemia before the diagnosis is known, some centres consider the following to be a safer alternative:
- L-arginine 300 mg/kg/day
 - L-carnitine 200 mg/kg/day
- Both can be given orally or intravenously
- **Dialysis** (haemodialysis, haemodiafiltration or haemofiltration)
 - Start immediately if plasma ammonia >500 µmol/l or if ammonia does not fall with the above measures.
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Note these regimens are not nutritionally complete and will cause malnutrition. They must not be continued longer than absolutely necessary.

Pathogenic mechanisms

Ammonia induces many electrophysiological, vascular and biochemical changes in experimental systems but it is not known to what extent these are relevant to the problems of hyperammonaemia in man [6]. Ammonia increases the transport of tryptophan across the blood brain barrier, which then leads to an increased production and release of serotonin [7]. Some of the symptoms of hyperammonaemia can be explained on this basis and the dietary tryptophan restriction has reversed anorexia in some patients with urea cycle disorders [8].

Glutamine has been shown to accumulate in the brain during hyperammonaemia in experimental animals and in man in vivo, using proton nuclear magnetic resonance spectroscopy [9]. The concentrations are such that the increase in osmolality could be responsible for cellular swelling and cerebral oedema.

Acute management

In the neonatal period the immediate goal of treatment is to control the metabolic derangement. Once plasma ammonia concentrations are greater than 500 µmol/l this is urgent. The strategies used are to stop any dietary protein and give a high energy intake, reduce plasma ammonia concentrations with dialysis and utilize alternative pathways of nitrogen excretion. The emergency management of neonatal hyperammonaemia is summarized in Table 3.

Protein and energy intake

As soon as hyperammonaemia is suspected all intake of protein should be stopped and a high energy intake given either orally or intravenously.

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