Guidelines for acute management of hyperammonemia in the Middle East region

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Introduction
Acute hyperammonemia is defined as elevated plasma ammonia levels associated with muscular hypotonia, seizures, vomiting, and impaired consciousness.1 The clinical features are heterogeneous depending on the age of the patient and on the type and severity of the underlying cause.2 The early symptoms in all age groups are loss of appetite and vomiting, which could be reversible if recognized and treated early. In newborns, common early symptoms are poor feeding, vomiting, lethargy, hyperventilation resulting often in respiratory alkalosis, and irritability that rapidly progresses to seizures, deep coma, and even death if not urgently treated. In infants, vomiting mimics pyloric stenosis, cow milk intolerance, or gastroenteritis. In older children and adults, vomiting, protein aversion, ataxia, confusion, disorientation, hallucinations, or abnormal behavior point to central
nervous system or psychiatric disorders.\textsuperscript{2} If untreated, severe hyperammonemia will be neurotoxic and will cause irreversible brain damage both to the developing as well as to the mature brain.\textsuperscript{1,5} Hyperammonemic encephalopathy is associated with high mortality rates.\textsuperscript{6,7} The total duration of a hyperammonemic coma and the extent of hyperammonemia are the most relevant prognostic factors and are negatively correlated with the patient’s neurological outcome.\textsuperscript{6–4} Therefore, prompt identification and treatment of hyperammonemia are vital to optimize the outcome.

In the Middle East, the prevalence of urea cycle disorders and organic acidemias, which are the major causes of hyperammonemia, is probably higher than in other parts of the world.\textsuperscript{9–12} This is most likely due to the high rate of consanguinity leading to increased autosomal recessive disorders in the region.\textsuperscript{10–14} In addition to the high prevalence, there are several major challenges that must be considered as risk factors for the outcome of hyperammonemic individuals in the Middle East. These include lack of awareness of health care professionals about prompt diagnosis and adequate management of hyperammonemia, strained communication between physicians at primary, secondary, and tertiary hospitals, and lack of knowledge and availability of the diagnostic tools and medications required in the acute management of hyperammonemia. Finally, the recognition of late onset and milder cases diagnosed in adulthood would require the particular alertness among internists and adult intensive care units, a goal certainly not yet everywhere achieved. Based on these considerations, we regard the development of regional guidelines for the acute management of hyperammonemia as urgent and a chance to improve the patients’ outcome in the Middle East. In this article, therefore, we have developed consensus guidelines that in particular address the unique situation in the Middle East and have based them on the highest available level of evidence. The aim of these guidelines is to homogenize and harmonize the various regional treatment protocols currently used for patients with acute hyperammonemia, and to provide a resource for metabolic specialists and likewise for physicians who may come in contact with individuals with acute hyperammonemia.

**Method**

We have based our guidelines on the European guidelines,\textsuperscript{17} published in 2012, which followed a strict methodological protocol. To include all relevant recent information, we searched PubMed and Embase databases to include published materials from 2011 to 2014 that were not covered by the European guidelines.\textsuperscript{6,18–38} Development of the present guidelines followed the process of a Delphi conference and involved one preliminary meeting and two follow-up meetings with email exchanges between the Middle East Hyperammonemia and Urea Cycle Disorders Scientific Group regarding each draft of the manuscript. The Middle East Hyperammonemia and Urea Cycle Disorders Scientific Group include an expert panel of metabolic specialists from countries in the Middle East as well as renowned international advisors. The process of guideline development followed Grading of Recommendations Assessment, Development, and Evaluation methodology of classifying the evidence.\textsuperscript{39,40} Although these guidelines are developed with highest accuracy, such guidelines should be considered as recommendations that aim at guiding physicians, but should not be considered as a protocol that is blindly followed. In particular, as each patient is an individual and each situation may vary from what is described here, all medical decisions must be carefully made on an individual basis.

Evidence levels were classified in accordance with the Grading of Recommendations Assessment, Development, and Evaluation methodology.\textsuperscript{39–42}

**Results and discussion**

The first part of the results and discussion will focus on identifying unknown patients at risk for hyperammonemia. We regard an earlier identification of affected patients with high index of suspicion as a key to overall improved prognosis. The second part describes recommendations for the acute management of hyperammonemia and is divided into several sections according to practical considerations. Finally, other issues related to management and prognosis of hyperammonemia will be discussed. Embedded in the discussion are some answers to valid questions that may rise during the management of hyperammonemia in an emergency setting.

Although these guidelines are built upon previously published guidelines, the novelty of these guidelines is the focus on answers to many practical questions that were not discussed in depth in the previously published materials.\textsuperscript{17,43} Examples for practical aspects are: the meaning and definition of a high caloric intake, the normal ammonia levels according to the age, the amount of potassium in intravenous (IV) fluids when hyperammonemia scavengers are used. Additionally, the responsible person for insertion of the line(s) for dialysis, and the dosage of insulin if it is used during initial management of hyperammonemia are discussed. The protocol and suggested guidelines are summarized in Tables 1 and 2, respectively.
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give glucose polymers, or protein-free formulas (e.g., Pro-phree, Polycose, or Maxijul) through PO/NG as tolerated to give additional calories.

- Basic life support (CAB)
- Start carnitine, biotin, vitamin B12

in an undiagnosed acute case, start levocarnitine IV/PO 100 mg/kg/day divided q 6–8 h, hydroxycobalamin 1 mg IM/IV/PO, and biotin 10 mg IV/PO

- Consider starting insulin if hyperglycemia develops (glucose
- Prepare for CRRT

start 1.5 to double maintenance IVF as D10%, 0.45 NS

Table 1 Protocol for acute management of primary hyperammonemia based on ammonia level

<table>
<thead>
<tr>
<th>Ammonia level (µmol/L)</th>
<th>Undiagnosed case</th>
<th>Diagnosed case</th>
</tr>
</thead>
<tbody>
<tr>
<td>Above upper limit of normal</td>
<td>Stop protein intake</td>
<td>Same as Undiagnosed case</td>
</tr>
<tr>
<td></td>
<td>Give IV glucose at an appropriate dosage to prevent catabolism ± insulin</td>
<td></td>
</tr>
<tr>
<td>&gt;100 but &lt;250 (in neonate)</td>
<td>Same as above</td>
<td>Start medications and nitrogen scavengers according to the protocol of each disorder</td>
</tr>
<tr>
<td>&gt;150 but &lt;250</td>
<td>Start drug treatment with nitrogen scavengers (L-arginine and AMMONUL®)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Start carnitine, biotin, vitamin B12</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Start Carbaglu®</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Start lipid IV 2–3 g/kg to give higher calories</td>
<td></td>
</tr>
<tr>
<td>250–500</td>
<td>Same as above</td>
<td>Same as Undiagnosed case</td>
</tr>
<tr>
<td></td>
<td>Prepare for CRRT</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Begin CRRT, if no rapid drop of ammonia within 3–6 hours</td>
<td></td>
</tr>
<tr>
<td>&gt;500</td>
<td>Start CRRT with above measure</td>
<td></td>
</tr>
</tbody>
</table>

Abbreviations: IV, intravenous; CRRT, continuous renal replacement therapy.

Table 2 Summary of protocol of acute management of hyperammonemia

- Basic life support (CAB)
- Stop all source of protein both enteral and parenteral nutrition for a maximum of 24–48 hours
- Check glucose level (GlucoCheck)
- Insert an IV line (central and peripheral) and take blood for ammonia (NH₃), blood gases, Chem 1, and CBC, blood C/S (peripheral and central if patient has central line). Liver transaminases, Ca, alkaline phosphatase as STAT order
- Ammonia should be taken with precaution (without tourniquet, transported in ice water to the laboratory, separated within 30 minutes of collection, and analyzed immediately)
- Start 1.5 to double maintenance IVF as D10%, 0.45 NS + KCl 30 meq/L until serum K result is available, then adjust accordingly
- Call the pharmacy to prepare the medications and intralipid (see dosages in Tables 1 and 4)
- Call biochemical geneticist (metabolic) on call
- Consider starting insulin if hyperglycemia develops (glucose >10 mmol/L) at dose of 0.05–0.1 unit/kg/h and titrate up until blood glucose controlled (keep GlucoCheck 6.5–10 mmol/L). Total glucose requirements (mg/kg/min) depends on the age (0–1 year: 8–10, 1–3 years: 7–8, 4–6 years: 6–7, 7–12 years: 5–6, adolescent: 4–5, adults: 3–4)
- If ammonia >100 µmol/L in infants, children, and adults; and >150 µmol/L in neonates start loading dose of combined sodium benzoate and sodium phenylacetate (AMMONUL®) and arginine (see Tables 1 and 4)
- If the patient is on combined sodium benzoate and phenylacetate (AMMONUL®) or arginine give KCl 40 meq/L because they cause hyperchloremic hypokalemic metabolic acidosis. KCl can be given through peripheral line up to 60 meq/L; rate must not exceed 0.125 meq/kg/h
- Start dialysis if ammonia >300–500 µmol/L in neonates and children and there is no response to the medical treatment within 4 hours. Consult ICU and nephrology team if you anticipate starting dialysis in the next few hours
- Reloading has to be done carefully, in particular during the first 24 hours, as cumulative doses of >750 mg/kg/24 h of combined sodium benzoate and phenylacetate (AMMONUL®) have been shown to be associated with development of toxicity (vomiting, lethargy). Reloading only in neonates with severe disorders or those who are undergoing dialysis, and should be spaced at least 6 hours
- In an undiagnosed acute case also start N-carbamyl glutamate (Carbaglu®). It only exists as an enteral form, so it is generally given by NG tube. Give 100 mg/kg once followed by 50 mg/kg q 6 h
- In an undiagnosed acute case, start levocarnitine IV/PO 100 mg/kg/day divided q 6–8 h, hydroxycobalamin 1 mg IM/IV/PO, and biotin 10 mg IV/PO
- Give glucose polymers, or protein-free formulas (e.g., Pro-phree, Polycose, or Maxijul) through PO/NG as tolerated to give additional calories
- Do not decrease dextrose rate or amount and do not stop calorie delivery in the acute stage for any reason (e.g., medications, fluid bolus, or hyperglycemia) as this can precipitate hypoglycemia and catabolism, which will further worsen the patient’s condition
- Call metabolic dietitian on call
- If patient has a known diagnosis, do not stop other oral chronic medications (in case of vomiting, convert to IV forms if available)
- Antibiotics may be started if there is any evidence of sepsis. Ammonia, electrolyte, and blood gases analysis need to be done at regular intervals during this acceleration of management stage. The frequency is dictated by the patient’s condition and the speed at which results can be obtained
- Protein should be reintroduced within 24–48 hours of initiation of therapy even if the patient is on dialysis

Abbreviations: CBC, complete blood count; CAB, circulation, airway, breathing; Chem I, Na, K, Cl, creatinine; C/S, culture and sensitivity; Ca, calcium; NAGS, N-acetylglutamate synthase; CPS1, carbamoyl phosphate synthetase I; IVF, intravenous fluid; D10, dextrose 10%; NS, normal saline; NG, nasogastric; h, hours; IM, intramuscular; IV, intravenous; PO, peroral.
Identifying unknown patients at risk for hyperammonemia

When to order ammonia analysis?

Patients from all age groups who present with encephalopathy of unknown etiology should undergo measurements of plasma ammonia (NH₃). The signs and symptoms of hyperammonemia can vary according to age, but are nonspecific in all age groups. In neonates, lethargy, poor feeding, and vomiting are frequent; it may progress to central hyperventilation, seizures, coma, and death if left untreated. In older children, symptoms include unexplained change in consciousness, unexplained or unusual neurological or psychiatric illness, and acute liver failure. Chronically, hyperammonemia can present as recurrent vomiting, headache, ataxia, strange behavior, especially if it occurs episodically, developmental delay, and aversion to protein. It is strongly recommended to consider ammonia measurement in all encephalopathic patients, especially in newborns who require a septic screen and are sick looking at the same time. Hospitals may also decide to install an automatic red flag as warning for certain clinical circumstances as recently suggested.

Statement #1: strong recommendation, low quality evidence (1C)

Any individual at any age who presents with any of the following symptoms should undergo prompt plasma ammonia determination:

- Acute unexplained encephalopathy
- Acute neurological illness
- Suspected sepsis in a newborn
- Developmental delay of unknown cause
- Recurrent vomiting
- Acute liver failure, elevated transaminases of unknown cause
- Ataxia
- Headache, especially if episodic in women
- Coma or stupor
- Psychiatric symptoms
- Aversion to protein.

Initial management when hyperammonemia is suspected

Once hyperammonemia is suspected, the patient should be managed in a hospital with access to basic metabolic tests, first-line hyperammonemia medications, dialysis facilities, and metabolic specialists. If any of these elements are not available, the patient should be transferred without delay to a specialist center after stabilization according to basic life support (circulation, airway, and breathing), addressing vital signs as with any critically ill patient and including blood glucose monitoring. In addition, the following should be done:

1. Insert IV lines, if possible as a central venous access. If this cannot be achieved, an intraosseous access could be an alternative.
2. Maintain airway: intubate and ventilate if necessary.
3. Adequate rehydration using minimum of dextrose 10% glucose and high calorie intake, maintain normal blood pressure, and add vasopressors if necessary.
4. Take blood and urine samples as outlined in Table 3.

What ammonia levels require immediate action?

The normal ammonia level varies based on the patient’s age. However, in general hyperammonemia is considered if plasma ammonia is >50 µmol/L in infants, children, and adults and >100 µmol/L in newborns. Other authors have described the normal reference range as follows: up to 7 days: 94 µmol/L, 1 month–15 years: 48 µmol/L, and >15 years: 50 µmol/L. It should be highlighted that the values presented here should be considered as decision limits; the normal reference ranges of individual laboratories should be used for clinical interpretation. An alert system for nurses and clinicians should be active in the lab in case ammonia is above the reference range and this must be reported immediately to the health care professional in charge of the patient.

<table>
<thead>
<tr>
<th>Table 3 Laboratory testing that should be done in any patient suspected of having hyperammonemia</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ammonia</td>
</tr>
<tr>
<td>Glucose</td>
</tr>
<tr>
<td>Venous/arterial blood gas</td>
</tr>
<tr>
<td>Electrolytes (including calcium)</td>
</tr>
<tr>
<td>Anion gap</td>
</tr>
<tr>
<td>Blood urea nitrogen</td>
</tr>
<tr>
<td>Creatinine</td>
</tr>
<tr>
<td>Liver transaminases</td>
</tr>
<tr>
<td>Bilirubin</td>
</tr>
<tr>
<td>Albumin</td>
</tr>
<tr>
<td>Alkaline phosphatase</td>
</tr>
<tr>
<td>Lactate</td>
</tr>
<tr>
<td>Lipase (if symptoms indicate)</td>
</tr>
<tr>
<td>Blood culture</td>
</tr>
<tr>
<td>Complete blood count</td>
</tr>
<tr>
<td>Acylcarnitine profile (tandem mass spectrometry), total and free carnitine</td>
</tr>
<tr>
<td>Plasma amino acids</td>
</tr>
<tr>
<td>Save a blood sample for DNA banking and DNA testing (if genotype is not known)</td>
</tr>
<tr>
<td>Urine ketones (healthy infants should not have any)</td>
</tr>
<tr>
<td>Urine organic acids (obtained within 2 hours of presentation by any method of collection)</td>
</tr>
<tr>
<td>Urine for orotic acid</td>
</tr>
</tbody>
</table>
Statement #2: strong recommendation, low quality evidence (1C)
The normal reference ranges of individual laboratories should be used for clinical interpretation. In general, hyperammonemia requires further action, if plasma ammonia is >50 μmol/L in infants, children, and adults and >100 μmol/L in neonates. The given limits of plasma ammonia are decision limits (Table 1).

What precaution must be taken for testing of ammonia?
False-positive hyperammonemia is not uncommon; therefore, several precautions should be taken into consideration when collecting blood samples to measure ammonia:

- A free-flowing venous (or arterial) blood sample without tourniquet should be collected into a tube containing an anticoagulant (eg, lithium or heparin).
- The sample should be placed in ice water, transported to the laboratory, and analyzed immediately. Results should be available within 60 minutes of drawing the samples.

Statement #3: strong recommendation, moderate quality evidence (1B)
Ammonia should be measured in an emergency setting with results available in 60 minutes.

Statement #4: strong recommendation, low quality evidence (1C)
It is crucial to take blood for plasma amino acids and urine for organic acids and orotic acid before starting the treatment; however, urine sampling must not delay start of treatment.

What are the key elements of the initial management once hyperammonemia is identified?
Reversal of catabolism
a. Stop all sources of protein temporarily (for a maximum of 24–48 h).
b. Call the pharmacy to prepare the medications and glucose/lipid infusions (see dosages in Table 4).
c. Check blood glucose regularly (GlucoCheck).
d. Start high caloric intake in form of IV dextrose 10%, or higher if appropriate, at the rate of 1.5 times to double of maintenance requirements, with age-dependent glucose requirement (in mg/kg/min for 0–1 years: 8–10, 1–3 years: 7–8, 4–6 years: 6–7, 7–12 years: 5–6, adolescents: 4–5, adults: 3–4).47
e. If fatty acid oxidation defects are excluded, intralipid infusions should be started at 2–3 g/kg/day to give additional calories.17,48

What do we mean by high caloric intake?
It means covering at least 110% of the recommended daily allowance in order to shut down endogenous protein breakdown (recommended daily allowance: for newborn and infant: 110–120 kcal/kg/day, for 1–3 years: 100 kcal/kg/day, for 4–6 years: 90 kcal/kg/day, for 7–10 years: 70 kcal/kg/day, for 11–14 years: 50–55 kcal/kg/day).47

Statement #5: strong recommendation, low quality evidence (1C)
High index and early clinical suspicion as well as prompt diagnosis of hyperammonemia are vital to have a good outcome. The treatment of hyperammonemia should be started without delay unless a decision for withdrawal of treatment and for palliative care is made.

What drugs are used to treat hyperammonemia?
Sodium benzoate and sodium phenylacetate are ammonia scavengers that bypass the urea cycle by conjugation of benzoate with glycine to generate hippurate, and of phenylacetate with glutamine to generate phenylacetylglutamine, which are then excreted in urine. L-Arginine is a metabolite within the urea cycle and can help the urea cycle to run again if it is interrupted due to arginine deficiency resulting from a block within the urea cycle. Carbaglu® (N-carbamyl-L-glutamate) can replace N-acetylglutamate as an activator of mitochondrial carbamoyl phosphate synthetase 1, the first enzyme of the urea cycle.17,49,50 The dosages of medications, mechanism of action, route, and side effects are summarized in Table 4.

Dialysis
What is the indication of dialysis?

Neonates and children
Continuous venovenous hemodiafiltration should be started in neonates and children who have ammonia levels of >500 μmol/L or even at lower levels if there has been an inadequate response to medical management after 4 hours, which is the estimated time for preparing dialysis, including vascular access.51

Statement #6: strong recommendation, low quality evidence (1C)
Dialysis is indicated in neonates and children with ammonia levels 300–500 μmol/L. Dialysis should be strongly considered in neonates and children with ammonia >500 μmol/L or when there is no response to the treatment. Consult intensive care unit and nephrology team, if you anticipate starting dialysis in the next few hours.
Table 4 Medications used in acute management of hyperammonemia and their dosages

<table>
<thead>
<tr>
<th>Medications</th>
<th>Mechanism of action</th>
<th>Dosage</th>
<th>Route</th>
<th>Adverse reaction*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sodium benzoate</td>
<td>Conjugation with glycine to form hippuric acid</td>
<td>Weight ≤ 20 kg: 250 mg/kg as loading dose over 90 minutes followed by 250–500 mg/kg/day</td>
<td>IV</td>
<td>Cardiovascular: hypotension</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Weight &gt; 20 kg: 5.5 g/m² as loading dose over 90 minutes followed by 5.5 g/m²/day</td>
<td></td>
<td>Dermatologic: injection site reaction</td>
</tr>
<tr>
<td>Sodium phenylacetate</td>
<td>Conjugation with glutamine to form phenylacetylglutamine</td>
<td>Same as for Sodium benzoate</td>
<td>IV</td>
<td>Electrolytes imbalance: hyperglycemia,</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>hypokalemia, hypernatremia</td>
</tr>
<tr>
<td>AMMONUL®</td>
<td>Contains both sodium benzoate and sodium phenylacetate</td>
<td>Same as for Sodium benzoate</td>
<td>IV</td>
<td>Gastrointestinal: vomiting, diarrhea</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(AMMONUL® could be given through peripheral line on limited basis)</td>
<td></td>
<td>CNS: altered mental status, seizure,</td>
</tr>
<tr>
<td>L-Arginine</td>
<td>As an intermediate metabolite in the urea cycle. It can improve the flow through the urea cycle and thereby improves ammonia removal</td>
<td>250–400 mg/kg/day as loading dose over 90 minutes followed by 250–400 mg/kg/day</td>
<td>IV</td>
<td>Other: fever</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Same as for Sodium benzoate</td>
</tr>
<tr>
<td>Carbaglu® (carglumic acid)</td>
<td>Replace N-acetylglutamate as an activator of mitochondrial carbamoyl phosphate synthetase, the first enzyme of the urea cycle</td>
<td>100 mg/kg bolus per NG tube, then 25–62.5 mg/kg every 6 hours</td>
<td>NG</td>
<td>Hyperchloremic metabolic alkalosis,</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>hypokalemia, elevated BUN and creatinine</td>
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<td></td>
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<td></td>
<td></td>
<td>levels, flushing, nausea, vomiting,</td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td>abdominal cramps, bloating, numbness,</td>
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<td></td>
<td></td>
<td>headache</td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td>Abdominal pain, diarrhea, vomiting, anemia,</td>
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<td></td>
<td></td>
<td></td>
<td>otitis media, tinnitus, nasopharyngitis,</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>fever, headache</td>
</tr>
</tbody>
</table>

Notes: *It should be given through central line, however, could be given peripherally on limited basis. ^If on hemodialysis/hemodiafiltration, maintenance doses should be increased to 350 mg/kg/day (or proportional increase for body surface-based dose calculation). *It is supplied as a vial of 50 mL or 5,000 mg constituted of concentrated, aqueous 10% sodium benzoate and 10% sodium phenylacetate solution. Thus, each mL provides 100 mg of sodium benzoate and 100 mg of sodium phenylacetate in water. According to the prescribing information, AMMONUL must be diluted with sterile dextrose injection, 10% at ±25 mL/kg before administration. *It is supplied as 200 mg tablet; 5 or 60 tablets in a polypropylene bottle with polyethylene cap and desiccant unit. *Generally all the drugs are well tolerated and adverse reactions are only relevant if dosages are very high.

Abbreviations: BUN, blood urea nitrogen; IV, intravenous; NG, nasogastric; CNS, central nervous system.

Adults

In adults, hemodialysis (HD) or continuous renal replacement therapy (CRRT) in the form of continuous venovenous hemofiltration is the first-line therapy in case of acute hyperammonemia. Dialysis should be started as soon as possible when ammonia exceeds 200 µmol/L.17

Statement #7: strong recommendation, low quality evidence (1C)

Dialysis is the first-line treatment in acute hyperammonemia decompensations in adults.

What is the method of choice for dialysis?

Dialysis choices usually include HD and CRRT. Alternatively, but only in centers that lack the ability or expertise to perform extracorporeal therapy, peritoneal dialysis (PD) can be utilized. However, PD clears ammonia at a lower rate than hemodialysis modalities. HD is intermittent and gives highest ammonia clearance.18 However, in neonates, HD is difficult to perform due to technical challenges and a high risk of complications, for example, concerning the maintenance of an adequate blood pressure. Therefore, CRRT including continuous venovenous hemofiltration or continuous venovenous hemodiafiltration is the preferred method.17,18,52 It is very important to emphasize that at this stage, fast action is required to prevent brain injury; therefore, the line for dialysis should be inserted immediately. Every place should develop its own protocol for who should insert the dialysis line depending on the local resources. The catheter of dialysis can be removed after the plasma ammonia level has been stable in the normal range for at least a day. However, keeping the catheter in place should be balanced with the risk of maintaining the patient in an anticoagulated state.

Exchange transfusions should be avoided.17 Ammonia scavenger therapy needs to be continued during dialysis.

Statement #8: strong recommendation, moderate quality evidence (1B)

The method of choice for dialysis is CRRT, preferably hemodiafiltration. Peritoneal dialysis is a far less effective method. Exchange transfusions should be avoided.
1. Ondansetron (0.15 mg/kg) could be given to avoid vomiting when boluses of the ammonia scavengers are given, strong recommendation, low quality evidence (1C).

2. Laxative: Treat constipation aggressively since ammonia is also produced from urea breakdown by intestinal bacteria, strong recommendation, low quality evidence (1C).

3. If patient not diagnosed, consider additional administration of carnitine 100 mg/kg IV, hydroxycobalamin 1 mg intramuscular/intravenous, and biotin 10 mg IV/peroral, strong recommendation, moderate quality evidence (1B).

4. Antibiotics: Preferred to continue or initiate them as prophylaxis, strong recommendation, low quality evidence (1C).

5. Hyperventilation: When patient is ventilated, it is recommended to perform moderate hyperventilation to counteract cerebral edema, strong recommendation, low quality evidence (1C).

6. Mannitol: It has not been demonstrated to be effective in managing cerebral edema caused by hyperammonemia and therefore, it should not be given, strong recommendation, low quality evidence (1C).

7. Steroids should be avoided as they increase the amount of protein turnover and hence increase the nitrogen load, strong recommendation, low quality evidence (1C).

8. Glucose and insulin can serve as suppressors of catabolism, but their use require care. Consider insulin if hyperglycemia develops (blood glucose level >10 mmol/L) at dose of 0.05–0.1 unit/kg/h and titrate up until blood glucose controlled. The presence of glycosuria is an indication for continued administration of IV regular insulin at a rate that keeps glucose levels between 6.5 and 10 mmol/L, weak recommendation, low quality evidence (2C).

9. Valproic acid should be avoided in any patient who has known risk for hyperammonemia. It is known to decrease urea cycle function by inhibition of N-acetylglutamate synthase, strong recommendation, low quality evidence (1C).

10. NaHCO₃: Not recommended except if there is refractory acidosis, weak recommendation, low quality evidence (2C).

11. Citrulline: A clear diagnosis should be made before supplementing the patient with citrulline. Patients with arginosuccinate synthetase and argininosuccinate lyase deficiency have elevated citrulline, strong recommendation, moderate quality evidence (1B).

12. If the patient is on combined sodium benzoate and phenylacetate (AMMONUL®) or arginine give KCl 40 meq/L because they cause hyperchloremic hypokalemic metabolic acidosis. KCl can be given through peripheral line up to 60 meq/L, rate must not exceed 0.125 meq/kg/h, weak recommendation, low quality evidence (2C).

13. KCl can be given in the same line with lipid and ammonia scavengers, weak recommendation, low quality evidence (2C).
14. Combined sodium benzoate and sodium phenylacetate (AMMONUL®) can be given through peripheral lines, if no central line can be established, strong recommendation, low quality evidence (1C).

15. Reloading of ammonia scavengers should only be considered in neonates with severe hyperammonemic decompensation or those who are undergoing dialysis, and should be given with great caution to avoid adverse effects due to increased serum drug concentrations, strong recommendation, low quality evidence (1C).

Prognosis
Poor prognostic factors in such cases include the following:

• Hyperammonemic coma has lasted more than 3 days.\(^{17,57}\)
• Intracranial pressure is clearly increased.\(^{17}\)
• High level and long duration of hyperammonemia.\(^{17}\)

Statement #13: strong recommendation, low quality evidence (1C)
Neurodevelopmental prognosis is mainly dependent on total duration of coma, peak ammonia levels, and number of episodes of hyperammonemia. Additional studies are needed to identify other potential contributing factors.

Conclusion
In summary, there are several challenges and obstacles that clinicians face as they try to select the appropriate management protocol for acute hyperammonemia. These suggested guidelines aim to ease these challenges. However, the rarity of these diseases has resulted in mostly low evidence level for the statements made here, which corresponds to inferences derived from none analytical studies, such as case reports or case series or from expert opinion. Therefore, the recommendations contained herein should not be considered infallible or absolute.

Acknowledgments
The authors would like to thank Erik Brouwer from SOBI, Swedish Orphan Biovitrum for his valuable contribution. The Middle East Hyperammonemia and Urea Cycle Disorders Scientific Group meetings were organized with the help and financial support of SOBI, Swedish Orphan Biovitrum, but the company has no influence on the scientific contents of these guidelines.

Disclosure
The authors report no conflicts of interest in this work.

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