PDR® 60 EDITION 2006

PHYSICIANS' DESK REFERENCE®

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whether to discontinue nursing or to discontinue the drug, taking into account the importance of the drug to the mother.

Pediatric Use: Safety and effectiveness of TUSSIONEX Pennkinetic Extended-Release Suspension in pediatric patients under six have not been established (see WARNINGS).

Geriatric Use: Clinical studies of TUSSIONEX did not include sufficient numbers of subjects aged 65 and over to determine whether they respond differently from younger subjects. Other reported clinical experience has not identified differences in responses between the elderly and younger patients. In general, dose selection for an elderly patient should be cautious, usually starting at the low end of the dosing range, reflecting the greater frequency of decreased hepatic, renal, or cardiac function, and of concomitant disease or other drug therapy.

This drug is known to be substantially excreted by the kidney, and the risk of toxic reactions to this drug may be greater in patients with impaired renal function. Because elderly patients are more likely to have decreased renal function, care should be taken in dose selection, and it may be useful to monitor renal function.

ADVERSE REACTIONS

Central Nervous System: Sedation, drowsiness, mental clouding, lethargy, impairment of mental and physical performance, anxiety, fear, dysphoria, euphoria, dizziness, psychic dependence, mood changes.

Dermatologic System: Rash, pruritus.

Gastrointestinal System: Nausea and vomiting may occur; they are more frequent in ambulatory than in recumbent patients. Prolonged administration of TUSSIONEX Pennkinetic Extended-Release Suspension may produce constipation.

Genitourinary System: Ureteral spasm, spasm of vesicle sphincters and urinary retention have been reported with opiates.

Respiratory Depression: TUSSIONEX Pennkinetic Extended-Release Suspension may produce dose-related respiratory depression by acting directly on brain stem respiratory centers (see OVERDOSAGE).

Respiratory System: Dryness of the pharynx, occasional tightness of the chest.

DRUG ABUSE AND DEPENDENCE

TUSSIONEX Pennkinetic Extended-Release Suspension is a Schedule III narcotic. Psychic dependence, physical dependence and tolerance may develop upon repeated administration of narcotics; therefore, TUSSIONEX Pennkinetic Extended-Release Suspension should be prescribed and administered with caution. However, psychic dependence is unlikely to develop when TUSSIONEX Pennkinetic Extended-Release Suspension is used for a short time for the treatment of cough. Physical dependence, the condition in which continued administration of the drug is required to prevent the appearance of a withdrawal syndrome, assumes clinically significant proportions only after several weeks of continued oral narcotic use, although some mild degree of physical dependence may develop after a few days of narcotic therapy.

OVERDOSAGE

Signs and Symptoms: Serious overdosage with hydrocodone is characterized by respiratory depression (a decrease in respiratory rate and/or tidal volume, Cheyne-Stokes respiration, cyanosis), extreme somnolence progressing to stupor or coma, skeletal muscle flaccidity, cold and clammy skin; and sometimes bradycardia and hypotension. Although miosis is characteristic of narcotic overdose, mydriasis may occur in terminal narcosis or severe hypoxia. In severe overdosage apnea, circulatory collapse, cardiac arrest and death may occur. The manifestations of chlorpheniramine overdosage may vary from central nervous system depression to stimulation.

Treatment: Primary attention should be given to the reestablishment of adequate respiratory exchange through provision of a patent airway and the institution of assisted or controlled ventilation. The narcotic antagonist naloxone hydrochloride is a specific antidote for respiratory depression which may result from overdosage or unusual sensitivity to narcotics including hydrocodone. Therefore, an appropriate dose of naloxone hydrochloride should be administered, preferably by the intravenous route, simultaneously with efforts at respiratory resuscitation. Since the duration of action of hydrocodone in this formulation may exceed that of the antagonist, the patient should be kept under continued surveillance and repeated doses of the antagonist should be administered as needed to maintain adequate respiration. For further information, see full prescribing information for naloxone hydrochloride. An antagonist should not be administered in the absence of clinically significant respiratory depression. Oxygen, intravenous fluids, vasopressors and other supportive measures should be employed as indicated. Gastric emptying may be useful in removing unabsorbed drug.

DOSAGE AND ADMINISTRATION

Shake well before using.

Adults: 1 teaspoonful (5 mL) every 12 hours; do not exceed 2 teaspoonfuls in 24 hours.

Children 6-12: 1/2 teaspoonful every 12 hours; **do not exceed 1 teaspoonful in 24 hours.**

Not recommended for children under 6 years of age (see PRECAUTIONS).

HOW SUPPLIED

TUSSIONEX Pennkinetic (hydrocodone polistirex and chlorpheniramine polistirex) Extended-Release Suspension is a gold-colored suspension.

NDC 53014-548-67 473 mL bottle

Shake well. Dispense in a well-closed container. Store at 59°-86°F (15°-30°C).

Celltech Pharmaceuticals, Inc.
Rochester, NY 14623 USA
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® Celltech Manufacturing, Inc.
Tussionex® Pennkinetic® Extended-Release Suspension:
US Patent No. 4,762,709.2
Current as of 06/2005 LR242A
Rev. 12/02

VICON FORTE® Capsules ℞

[vī'kon for'tā]

Therapeutic Vitamins-Minerals

℞ only

DESCRIPTION

Each black and orange VICON FORTE® capsule for oral administration contains:

Vitamin A	8,000 IU
Vitamin E	50 IU
Ascorbic acid	150 mg
Zinc sulfate, USP*	80 mg
Magnesium sulfate, USP†	70 mg
Niacinamide	25 mg
Thiamine mononitrate	10 mg
d-Calcium pantothenate	10 mg
Riboflavin	5 mg
Manganese chloride	4 mg
Pyridoxine hydrochloride	2 mg
Folic acid	1 mg
Vitamin B_{12} (Cyanocobalamin)	10 mcg

* As 50 mg dried zinc sulfate.
† As 50 mg dried magnesium sulfate.

Each capsule also contains Edible ink, FD&C Blue No. 1, FD&C Red No. 40, FD&C Yellow No. 6, gelatin, lactose, magnesium stearate, silicon dioxide, sodium lauryl sulfate, and titanium dioxide.

HOW SUPPLIED

Orange and black capsules imprinted with "ucb" and "316" in bottles of 60 (NDC 50474-316-22) and 500 (NDC 50474-316-24) and unit-dose packs of 100 (NDC 50474-316-27).

Dispense in tight, light-resistant container with a child-resistant closure.

Manufactured for
UCB Pharma, Inc.
Smyrna, GA 30080
by Mallinckrodt Inc.
Hobart, NY 13788

Current as of 05/2005 1E 10/2003

ZAROXOLYN® TABLETS ℞

[zar "ox 'uh-lin]

(metolazone tablets, USP)

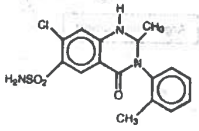
℞ Only

DO NOT INTERCHANGE: DO NOT INTERCHANGE ZAROXOLYN TABLETS AND OTHER FORMULATIONS OF METOLAZONE THAT SHARE ITS SLOW AND INCOMPLETE BIOAVAILABILITY AND ARE NOT THERAPEUTICALLY EQUIVALENT AT THE SAME DOSES TO MYKROX® TABLETS, A MORE RAPIDLY AVAILABLE AND COMPLETELY BIOAVAILABLE METOLAZONE PRODUCT. FORMULATIONS BIOEQUIVALENT TO ZAROXOLYN AND FORMULATIONS BIOEQUIVALENT TO MYKROX SHOULD NOT BE INTERCHANGED FOR ONE ANOTHER.

DESCRIPTION

ZAROXOLYN Tablets (metolazone tablets, USP) for oral administration contain 2½, 5, or 10 mg of metolazone, USP, a diuretic/saluretic/antihypertensive drug of the quinazoline class.

Metolazone has the molecular formula $C_{16}H_{16}ClN_3O_3S$, the chemical name 7-chloro-1, 2, 3, 4-tetrahydro-2-methyl-3-(2-methylphenyl)-4-oxo-6-quinazolinesulfonamide, and a molecular weight of 365.83. The structural formula is:



Metolazone is only sparingly soluble in water, but more soluble in plasma, blood, alkali, and organic solvents.

Inactive Ingredients: Magnesium stearate, microcrystalline cellulose and dye: 2½ mg-D&C Red No. 33; 5 mg-FD&C Blue No. 2; 10 mg-D&C Yellow No. 10 and FD&C Yellow No. 6.

HOW SUPPLIED

ZAROXOLYN Tablets (metolazone tablets, USP) are shallow biconvex, round tablets, and are available in three strengths:

2½ mg, pink, debossed "ZAROXOLYN" on one side, and "2½" on reverse side.

NDC 53014-975-71	Bottle of 100's
NDC 53014-975-90	Bottle of 1000's
NDC 53014-975-72	Carton of 100's, unit dose

5 mg, blue, debossed "ZAROXOLYN" on one side, and "5" on reverse side.

NDC 53014-850-71	Bottle of 100's
NDC 53014-850-90	Bottle of 1000's
NDC 53014-850-72	Carton of 100's, unit dose

10 mg, yellow, debossed "ZAROXOLYN" on one side, and "10" on reverse side.

NDC 53014-835-71	Bottle of 100's
NDC 53014-835-90	Bottle of 1000's
NDC 53014-835-72	Carton of 100's, unit dose

Store at 25°C (77°F); excursions permitted to 15°-30°C (59°-86°F) [See USP Controlled Room Temperature]. Protect from light. Keep out of the reach of children.

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Current as of 06/2005

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AMMONUL® ℞

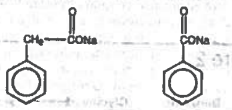
[ăm-mŏn-ūl]

(sodium phenylacetate and sodium benzoate) Injection 10% / 10%

Rx Only

DESCRIPTION

AMMONUL® (sodium phenylacetate and sodium benzoate) Injection 10% / 10% is a sterile, concentrated, aqueous solution of sodium phenylacetate and sodium benzoate, used for the treatment of hyperammonemia in urea cycle disorders. The pH of the solution is between 6 and 8. Sodium phenylacetate is a crystalline, white to off-white powder with a strong, offensive odor. It is soluble in water. Sodium benzoate is a white and odorless, crystalline powder that is readily soluble in water.



Sodium Phenylacetate Sodium Benzoate

Sodium phenylacetate has a molecular weight of 158.13 and the molecular formula $C_8H_7NaO_2$. Sodium benzoate has a molecular weight of 144.11 and the molecular formula $C_7H_5NaO_2$.

Each mL of AMMONUL® contains 100 mg of sodium phenylacetate and 100 mg of sodium benzoate, and Water for Injection. Sodium hydroxide and/or hydrochloric acid may have been used for pH adjustment.

AMMONUL® injection is a sterile, concentrated solution intended for intravenous administration via a central line only after dilution (see **DOSAGE AND ADMINISTRATION**). AMMONUL® is packaged in single-use vials.

CLINICAL PHARMACOLOGY

Sodium phenylacetate and sodium benzoate are metabolically active compounds that can serve as alternatives to urea for the excretion of waste nitrogen. Phenylacetate conjugates with glutamine in the liver and kidneys to form phenylacetylglutamine, via acetylation. Phenylacetylglutamine is excreted by the kidneys via glomerular filtration and tubular secretion. The nitrogen content of phenylacetylglutamine per mole is identical to that of urea (both contain two moles of nitrogen). Similarly, preceded by acylation, benzoate conjugates with glycine to form hippuric acid, which is rapidly excreted by the kidneys by glomerular filtration and tubular secretion. One mole of hippuric acid contains one mole of waste nitrogen. It has been shown that phenylacetylglutamine and hippurate can serve as alternative vehicles to effectively reduce waste nitrogen levels in patients with deficiencies of urea cycle enzymes and, thus, attenuate the risk of ammonia and glutamine-induced neurotoxicity.

Continued on next page

Ammonul—Cont.

Urea cycle disorders can result from decreased activity of any of the following enzymes: *N*-acetylglutamate synthetase (NAGS), carbamyl phosphate synthetase (CPS), argininosuccinate synthetase (ASS), ornithine transcarbamylase (OTC), argininosuccinate lyase (ASL), or arginase (ARG). The most frequently observed initial presenting symptoms in neonates include lethargy, seizures, poor feeding, neurologic changes, edema, and respiratory distress. Patients with milder forms of enzyme deficiencies may not present until late childhood, adolescence, or adulthood. Hyperammonemic crisis with lethargy, delirium, and coma, in these patients, are often precipitated by viral illness, high protein diet, stress, or trauma.

Plasma and urine amino acid analyses are used to diagnose ASS and ASL and to provide a preliminary diagnosis of CPS, OTC, or ARG. Blood citrulline levels are very low or absent in OTC and CPS, very high in ASS, and normal to moderately high in ASL and ARG. ASL may be distinguished by the presence of high levels of the unusual amino acid argininosuccinic acid (ASA) in the urine. It should be noted, however, that ASA tends to co-elute initially with other amino acids (such as leucine and isoleucine) in chromatographs, and may be missed on initial examination. ARG is characterized by high urine levels of arginine. A definitive diagnosis of CPS and OTC require a liver biopsy, and red blood cell enzyme analysis is needed to confirm a diagnosis of ARG. Patients suspected of having a urea cycle disorder, based on family history, should have documented hyperammonemia prior to administration of AMMONUL®.

Mechanism of Action

Figure 2 is a schematic illustrating how the components of AMMONUL®, phenylacetate and benzoate, provide an alternative pathway for nitrogen disposal in patients without a fully functioning urea cycle. Two moles of nitrogen are removed per mole of phenylacetate when it is conjugated with glutamine, and one mole of nitrogen is removed per mole of benzoate when it is conjugated with glycine.

[See figure 2 below]

Pharmacokinetics

The pharmacokinetics of intravenously administered AMMONUL® were characterized in healthy adult volunteers. Both benzoate and phenylacetate exhibited nonlinear kinetics. Following 90 minute intravenous infusion mean AUC_{last} for benzoate was 20.3, 114.9, 564.6, 562.8, and 1599.1 mcg/mL following doses of 1, 2, 3.75, 4, and 5.5 g/m^2, respectively. The total clearance decreased from 5.19 to 3.62 L/h/m^2 at the 3.75 and 5.5 g/m^2 doses, respectively.

Similarly, phenylacetate exhibited nonlinear kinetics following the priming dose regimens. AUC_{last} was 175.6, 713.8, 2040.6, 2181.6, and 3829.2 mcg•h/mL following doses of 1, 2, 3.75, 4, and 5.5 g/m^2, respectively. The total clearance decreased from 1.82 to 0.89 mcg•h/mL with increasing dose (3.75 and 4 g/m^2, respectively).

During the sequence of 90 minute priming infusion followed by a 24 hour maintenance infusion, phenylacetate was detected in the plasma at the end of infusion (T_{max} of 2 hr at 3.75 g/m^2) whereas, benzoate concentrations declined rapidly (T_{max} of 1.5 hr at 3.75 g/m^2) and were undetectable at 14 and 26 hours following the 3.75 and 4 g/m^2 dose, respectively.

A difference in the metabolic rates for phenylacetate and benzoate was noted. The formation of hippurate from benzoate occurred more rapidly than that of phenylacetylglutamine from phenylacetate, and the rate of elimination for hippurate appeared to be more rapid than that for phenylacetylglutamine.

Pharmacokinetic observations have also been reported from twelve episodes of hyperammonemic encephalopathy in seven children diagnosed (age 3 to 26 months) with urea cycle disorders who had been administered AMMONUL® intravenously. These data showed peak plasma levels of phenylacetate and benzoate at approximately the same times as were observed in adults. As in adults, the plasma levels of phenylacetate were higher than benzoate and were present for a longer time [1].

The pharmacokinetics of intravenous phenylacetate have been reported following administration to adult patients with advanced solid tumors. The decline in serum phenylacetate concentrations following a loading infusion of 150 mg/kg was consistent with saturable enzyme kinetics. Ninety-nine percent of administered phenylacetate was excreted as phenylacetylglutamine [2,3].

Special Populations

Gender:

Pharmacokinetic parameters of AMMONUL® were compared in healthy males and females. Bioavailability of both benzoate and phenylacetate was slightly higher in females than in males. However, conclusions cannot be drawn due to the limited number of subjects in this study.

Hepatic Insufficiency:

Limited information is available on the metabolism and excretion of sodium phenylacetate and sodium benzoate in patients with impaired hepatic function. However, as the liver is one of the two organs (the other is the kidney) in which the metabolic conjugation of sodium phenylacetate and sodium benzoate is known to take place, care should be used in administering AMMONUL® to patients with hepatic insufficiency.

Renal Impairment:

For effective AMMONUL® drug therapy, renal clearance of the drug metabolites and subsequently ammonia is required. Therefore, patients with impaired renal function should be closely monitored.

Dialysis:

Intravenous use of AMMONUL® is complementary with the use of dialysis[4,5]. In the non-neonatal study patient population treated with AMMONUL® , dialysis (standard hemodialysis, peritoneal dialysis, arteriovenous hemofiltration, or other dialysis) was required in 13% of hyperammonemic episodes. Standard hemodialysis was the most frequently used dialysis method. High levels of ammonia can be reduced quickly when AMMONUL® is used with dialysis, as the ammonia-scavenging of AMMONUL® suppresses the production of ammonia from catabolism of endogenous protein[6] and dialysis eliminates the ammonia and ammonia conjugates.

Drug Interactions:

Formal drug interaction studies have not been performed with AMMONUL®.

Pharmacodynamics

In patients with hyperammonemia due to deficiencies in enzymes of the urea cycle, AMMONUL® has been shown to decrease elevated plasma ammonia levels and improve encephalopathy and survival outcome compared to historical controls. These effects are considered to be the result of reduction in nitrogen overload through glutamine and glycine scavenging by AMMONUL® in combination with appropriate dietary and other supportive measures.

Clinical Data

The efficacy of AMMONUL® in improving patient survival of acute hyperammonemic episodes was demonstrated in an analysis of 316 patients (1045 episodes of hospitalization) treated between 1981 and 2003.

The demographic characteristics and diagnoses of the patient population are shown in Table 1.

Table 1 Baseline Characteristics and Diagnoses of Study Population

		Patients* N=316
Gender	Male	158 (51%)
	Female	150 (49%)
Age (years)	N	310
	Mean (SD)	6.2 (8.54)
	Min–Max	0.0–53.0
Age groups	0–30 days	104 (34%)
	31 days–2 years	55 (18%)
	> 2–12 years	90 (29%)
	> 12–16 years	30 (10%)
	> 16 years	31 (10%)
Enzyme deficiency	OTC	146 (46%)
	ASS	71 (22%)
	CPS	38 (12%)
	ASL	7 (2%)
	ARG	2 (< 1%)
	THN	2 (< 1%)
	Other**	56 (18%)

OTC = ornithine transcarbamylase deficiency; ASS = argininosuccinate synthetase deficiency; CPS = carbamyl phosphate synthetase deficiency; ASL = argininosuccinate lyase deficiency; ARG = arginase deficiency; THN = transient hyperammonemia of the newborn

*For the summary at the patient level, data obtained at first episode used.

**Diagnosis unknown or pending (33 episodes), acidemia (14 episodes), HHH syndrome (6 episodes), carnitine translocase deficiency (4 episodes), liver disease (3 episodes), HMG CoA lyase deficiency (1 episode), non-ketotic hyperglycinemia (1 episode), suspected fatty acid oxidation deficiency (1 episode), and valproic-acid-induced hyperammonemia (1 episode).

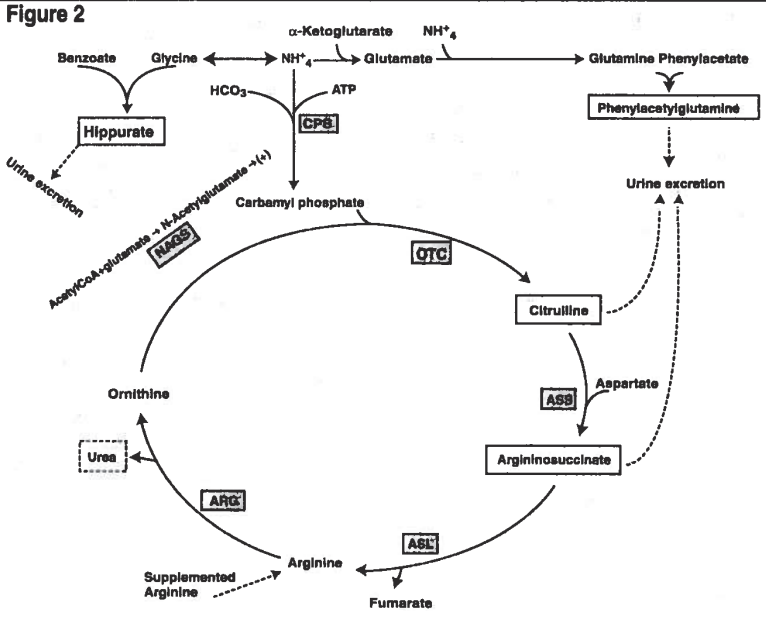
On admission to the hospital, patients with hyperammonemia or a potential urea cycle disorder (UCD) were treated with a bolus dose of 0.25 g/kg (or 5.5 g/m^2) sodium phenylacetate + 0.25 g/kg (or 5.5 g/m^2) sodium benzoate over a period of 90 minutes to 6 hours, depending on the specific UCD. Infusions also contained arginine; the dose of arginine depended on the specific UCD. After completion of the bolus dose, maintenance infusions of the same dose over 24 hours were continued until the patient was no longer hyperammonemic or oral therapy could be tolerated. The mean (SD) duration of treatment was 4.6 (6.45) days per episode, and ranged from 1 to 72 days.

Survival was substantially improved after AMMONUL® treatment compared with historical values (estimated 14% 1-year survival rate with dietary therapy alone) [10] and with dialysis (estimated 43% survival of acute hyperammonemia) [11].

Ninety-four percent (981 of 1045) of hyperammonemic episodes treated with AMMONUL® resulted in patients being discharged from the hospital. Eighty percent of patients (252 of 316) survived their last episode. Of the 64 patients who died, 53 (83%) died during their first hyperammonemic episode. Of the 104 neonates (<30d) treated with AMMONUL®, 34 (33%) died during the first hyperammonemic episode.

Ammonia levels decreased from very high levels (> 4 times the upper limit of normal [ULN]) to lower levels in 91% of episodes after treatment. In patients responding to therapy, mean ammonia concentrations decreased significantly within four hours of initiation of AMMONUL® therapy and

Figure 2



CPS = carbamyl phosphate synthetase; OTC = ornithine transcarbamylase; ASS = argininosuccinate synthetase; ASL = argininosuccinate lyase; ARG = arginase; NAGS = N-acetylglutamate synthetase

were maintained. Dialysis is recommended for those patients who fail to have a significant reduction in plasma ammonia levels within 4 to 8 hours after receiving AMMONUL®. A shift from high (≤ 4 times ULN) to very high (> 4 times ULN) levels was observed in only 4% of the episodes.

Improvements in neurological status endpoints were observed in most episodes and patients. Overall, investigators rated neurological status as improved, much improved, or the same in 93% of episodes, and overall status in response to treatment as improved, much improved, or the same in 97% of episodes. Recovery from coma was observed in 97% of episodes where coma was present at admission (111 of 114 episodes).

INDICATIONS AND USAGE

AMMONUL® is indicated as adjunctive therapy for the treatment of acute hyperammonemia and associated encephalopathy in patients with deficiencies in enzymes of the urea cycle. In acute neonatal hyperammonemic coma, in moderate to severe episodes of hyperammonemic encephalopathy, and in episodes of hyperammonemia which fail to respond to an initial course of AMMONUL® therapy, hemodialysis is the most rapid and effective technique for removing ammonia [12,13]. In such cases, the concomitant administration of AMMONUL® can help prevent the re-accumulation of ammonia by increasing waste nitrogen excretion [4,5,13].

CONTRAINDICATIONS

AMMONUL® should not be administered to patients with known hypersensitivity to sodium phenylacetate or sodium benzoate.

WARNINGS

Any episode of acute symptomatic hyperammonemia should be treated as a life-threatening emergency. Treatment of hyperammonemia may require dialysis, preferably hemodialysis, to remove a large burden of ammonia. Uncontrolled hyperammonemia can rapidly result in brain damage or death, and prompt use of all therapies necessary to reduce ammonia levels is essential.

Management of hyperammonemia due to inborn errors of metabolism should be done in coordination with medical personnel familiar with these diseases. The severity of the disorder may necessitate the use of hemodialysis combined with nutritional management and medical support. The multidisciplinary nature of the treatment usually requires the facilities of a tertiary or quaternary care center.

Ongoing monitoring of plasma ammonia levels, neurological status, laboratory tests, and clinical response in patients receiving AMMONUL® is crucial to assess patient response to treatment. Because urine potassium loss is enhanced by the excretion of the nonreabsorbable anions, phenylacetylglutamine and hippurate, plasma potassium levels should be carefully monitored and appropriate treatment given when necessary. Serum electrolyte levels should be monitored and maintained within the normal range.

AMMONUL® contains 30.5 mg of sodium per mL of undiluted product. Thus, AMMONUL® should be used with great care, if at all, in patients with congestive heart failure or severe renal insufficiency, and in clinical states in which there is sodium retention with edema. If an adverse reaction does occur, discontinue administration of AMMONUL®, evaluate the patient, and institute appropriate therapeutic countermeasures.

Administration must be through a central line. Administration through a peripheral line may cause burns.

Bolus infusion flow rates are relatively high, especially for infants (see **DOSAGE AND ADMINISTRATION**). Extravasation of AMMONUL® into the perivenous tissues may lead to skin necrosis. If extravasation is suspected, discontinue the infusion and resume at a different infusion site, if necessary. Standard treatment for extravasation can include aspiration of residual drug from the catheter, limb elevation, and intermittent cooling using cold packs [14]. The infusion site must be monitored closely for possible infiltration during drug administration. Do not administer undiluted product.

Due to structural similarities between phenylacetate and benzoate to salicylate, AMMONUL® may cause side effects typically associated with salicylate overdose, such as hyperventilation and metabolic acidosis. The clinician is advised to perform blood chemistry profiles, and frequent blood pH and pCO_2 monitoring.

PRECAUTIONS

General:

AMMONUL® is a concentrated solution and must be diluted before administration via a central line. Because sodium phenylacetate and sodium benzoate are metabolized in the liver and kidney, and since phenylacetylglutamine and hippurate are primarily excreted by the kidney, use caution when administering AMMONUL® to patients with hepatic or renal insufficiency. AMMONUL® infusion has been associated with nausea and vomiting. An antiemetic may be administered during AMMONUL® infusion.

Because of prolonged plasma levels achieved by phenylacetate in pharmacokinetic studies, repeat loading doses of AMMONUL® should not be administered.

Use of corticosteroids may cause the breakdown of body protein and, thereby, potentially increase plasma ammonia levels in patients with impaired ability to form urea.

Neurotoxicity of Phenylacetate:

Neurotoxicity was reported in cancer patients receiving intravenous phenylacetate, 250-300 mg/kg/day for 14 days, repeated at 4-week intervals. Manifestations were predominantly somnolence, fatigue, and lightheadedness, with less frequent headaches, dysgeusia, hypoacusis, disorientation, impaired memory, and exacerbation of a preexisting neuropathy. These adverse events were mainly mild. The acute onset of symptoms upon initiation of treatment and reversibility of symptoms when the phenylacetate was discontinued suggest a drug effect [2,3].

In animal studies, subcutaneous administration to rat pups of 190-474 mg/kg of phenylacetate caused decreased proliferation and increased loss of neurons, and reduced central nervous system (CNS) myelin. Cerebral synapse maturation was retarded, and the number of functioning nerve terminals in the cerebrum was reduced, which resulted in impaired brain growth [15]. Pregnant rats were given phenylacetate at 3.5 Åμmol/g/day subcutaneous from gestation day 7 through normal delivery. Prenatal exposure of rat pups to phenylacetate produced lesions in layer 5 cortical pyramidal cells; dendritic spines were longer and thinner than normal and reduced in number [16].

Drug Interactions:

Some antibiotics such as penicillin may compete with phenylacetylglutamine and hippurate for active secretion by renal tubules, which may affect the overall disposition of the infused drug.

Probenecid is known to inhibit the renal transport of many organic compounds, including aminohippuric acid, and may affect renal excretion of phenylacetylglutamine and hippurate [13].

There have been reports that valproic acid can induce hyperammonemia through inhibition of the synthesis of N-acetylglutamate, a co-factor for carbamyl phosphate synthetase [14]. Therefore, administration of valproic acid to patients with urea cycle disorders may exacerbate their condition and antagonize the efficacy of AMMONUL® [15].

Carcinogenesis, Mutagenesis, Impairment of Fertility:

Carcinogenicity, mutagenicity and fertility studies of sodium phenylacetate have not been conducted. Sodium benzoate has been extensively tested as a food preservative. Results indicate that sodium benzoate is not mutagenic or carcinogenic, and does not impair fertility.

Pregnancy:

Pregnancy Category C. Animal reproduction studies have not been conducted with AMMONUL®. It is not known whether AMMONUL® can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity. Thus, AMMONUL® should be given to a pregnant woman only if clearly needed.

Labor and Delivery:

The effects of AMMONUL® on labor and delivery are unknown.

Nursing Mothers:

It is not known whether sodium phenylacetate, sodium benzoate, or their conjugation products are excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when AMMONUL® is administered to a nursing woman.

Pediatric:

AMMONUL® has been used as a treatment for acute hyperammonemia in pediatric patients including patients in the early neonatal period (see **DOSAGE AND ADMINISTRATION**).

ADVERSE REACTIONS

The safety data were obtained from 316 patients who received AMMONUL® as emergency (rescue) or prospective treatment for hyperammonemia as part of an uncontrolled, open-label study. The study population included patients between the ages of 0 to 53 years with a mean (SD) of 6.2 (8.54) years; 51% were male and 49% were female who had the following diagnoses: OTC (46%), ASS (22%), CPS (12%), ASL (2%), ARG (< 1%), THN (< 1%), and other (18%).

Table 2 Adverse Events Occurring in ≥ 3% of Patients Treated with AMMONUL®

	Patients N=316
No. patients with any adverse event	163 (52%)
Blood and lymphatic system disorders	35 (11%)
Anemia NOS	12 (4%)
Disseminated intravascular coagulation	11 (3%)
Cardiac disorders	28 (9%)
Gastrointestinal disorders	42 (13%)
Diarrhea NOS	10 (3%)
Nausea	9 (3%)
Vomiting NOS	29 (9%)
General disorders and administration-site conditions	45 (14%)
Injection-site reaction NOS	11 (3%)
Pyrexia	17 (5%)
Infections	39 (12%)
Urinary tract infection NOS	9 (3%)
Injury, poisoning and procedural complications	12 (4%)
Investigations	32 (10%)
Metabolism and nutrition disorders	67 (21%)
Acidosis NOS	8 (3%)
Hyperammonemia	17 (5%)
Hyperglycemia NOS	22 (7%)
Hypocalcemia	8 (3%)
Hypokalemia	23 (7%)
Metabolic acidosis NOS	13 (4%)
Nervous system disorders	71 (22%)
Brain edema	17 (5%)
Coma	10 (3%)
Convulsions NOS	19 (6%)
Mental impairment NOS	18 (6%)
Psychiatric disorders	16 (5%)
Agitation	8 (3%)
Renal and urinary disorders	14 (4%)
Respiratory, thoracic and mediastinal disorders	47 (15%)
Respiratory distress	9 (3%)
Skin and subcutaneous tissue disorders	19 (6%)
Vascular disorders	19 (6%)
Hypotension NOS	14 (4%)

Clinically Important Adverse Reactions

Adverse events occurred most frequently in the following system organ classes: nervous system disorders (22% of patients), metabolism and nutrition disorders (21% of patients), and respiratory, thoracic and mediastinal disorders (15% of patients). The most frequently reported adverse events were vomiting (9% of patients), hyperglycemia (7% of patients), hypokalemia (7% of patients), convulsions (6% of patients), and mental impairment (6% of patients).

Adverse events leading to study drug discontinuation occurred in 4% of patients. Metabolic acidosis and injection-site reactions each led to discontinuation in 2 patients (< 1%). Adverse events leading to discontinuation in 1 patient included bradycardia, abdominal distension, injection-site extravasation, injection-site hemorrhage, blister, overdose, subdural hematoma, hyperammonemia, hypoglycemia, clonus, coma, increased intercranial pressure, hypercapnia, Kussmaul respiration, respiratory distress, respiratory failure, pruritis, and maculo-papular rash.

Subpopulation and Risk Factor Data

Adverse events were reported with similar frequency in patients with OTC, ASS, CPS, and diagnoses categorized as "other." Nervous system disorders were more frequent in patients with OTC and CPS, compared with patients with ASS and patients with "other" diagnoses. Convulsions and mental impairment were reported in patients with OTC and CPS. These observations are consistent with literature reports that patients with enzyme deficiencies occurring earlier in the urea cycle (i.e., OTC and CPS) tend to be more severely affected.

Adverse event profiles did differ by age group. Patients ≤ 30 days of age had more blood and lymphatic system disorders and vascular disorders (specifically hypotension), while patients > 30 days of age had more gastrointestinal disorders (specifically nausea, vomiting and diarrhea).

Other Less Common Adverse Events Occurring in < 3% of Patients

Less common adverse events that could represent drug-induced reactions or are characterized as severe are listed below by body system.

BLOOD AND LYMPHATIC SYSTEM DISORDERS: coagulopathy, pancytopenia, thrombocytopenia

CARDIAC DISORDERS: atrial rupture, cardiac or cardiopulmonary arrest/failure, cardiogenic shock, cardiomyopathy, pericardial effusion

EYE DISORDERS: blindness

GASTROINTESTINAL DISORDERS: gastrointestinal hemorrhage

GENERAL DISORDERS AND ADMINISTRATION SITE CONDITIONS: asthenia, brain death, chest pain, multiorgan failure, edema

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Ammonul—Cont.

HEPATOBILIARY DISORDERS: cholestasis, hepatic artery stenosis, hepatic failure/ hepatotoxicity, jaundice
INFECTIONS AND INFESTATIONS: sepsis/septic shock
INJURY, POISONING AND PROCEDURAL COMPLICATIONS: brain herniation, subdural hematoma
INVESTIGATIONS: blood carbon dioxide changes, blood glucose changes, blood pH increased, cardiac output decreased, pCO_2 changes, respiratory rate increased
METABOLISM AND NUTRITION DISORDERS: alkalosis, dehydration, fluid overload/retention, hyperkalemia, hypernatremia, alkalosis, tetany
NEOPLASMS BENIGN, MALIGNANT AND UNSPECIFIED: hemangioma acquired
NERVOUS SYSTEM DISORDERS: areflexia, ataxia, brain infarction, brain hemorrhage, cerebral atrophy, clonus, depressed level of consciousness, encephalopathy, nerve paralysis, intracranial pressure increased, tremor
PSYCHIATRIC DISORDERS: acute psychosis, aggression, confusional state, hallucinations
RENAL AND URINARY DISORDERS: anuria, renal failure, urinary retention
RESPIRATORY, THORACIC AND MEDIASTINAL DISORDERS: acute respiratory distress syndrome, dyspnea, hypercapnia, hyperventilation, Kussmaul respiration, pneumonia aspiration, pneumothorax, pulmonary hemorrhage, pulmonary edema, respiratory acidosis or alkalosis, respiratory arrest/failure
SKIN AND SUBCUTANEOUS TISSUE DISORDERS: alopecia, pruritis generalized, rash, urticaria
VASCULAR DISORDERS: flushing, hemorrhage, hypertension, phlebothrombosis/thrombosis

OVERDOSAGE

Overdosage has been reported during AMMONUL® treatment in urea cycle-deficient patients [17]. All patients in the uncontrolled open-label study were to be treated at the same dose of AMMONUL®. However, some patients received more than the dose level specified in the protocol. In 16 of the 64 deaths, the patient received a known overdose of AMMONUL®. Causes of death in these patients included cardiorespiratory failure/arrest (6 patients), hyperammonemia (3 patients), increased intracranial pressure (2 patients), pneumonitis with septic shock and coagulopathy (1 patient), error in dialysis procedure (1 patient), respiratory failure (1 patient), intractable hypotension and probable sepsis (1 patient), and unknown (1 patient). Additionally, other signs of intoxication may include obtundation (in the absence of hyperammonemia), hyperventilation, a severe compensated metabolic acidosis, perhaps with a respiratory component, large anion gap, hypernatremia and hyperosmolarity, progressive encephalopathy, cardiovascular collapse, and death.
In case of overdose of AMMONUL®, discontinue the drug and institute appropriate emergency medical monitoring and procedures. In severe cases, the latter may include hemodialysis (procedure of choice) or peritoneal dialysis (when hemodialysis is unavailable) [17].

DOSAGE AND ADMINISTRATION

Administration must be through a central line. Administration through a peripheral line may cause burns.
General
AMMONUL® is administered intravenously as a loading dose infusion administered over 90 to 120 minutes, followed by an equivalent maintenance dose infusion administered over 24 hours. AMMONUL® may not be administered by any other route. Administration of analogous oral drugs, such as Buphenyl® (sodium phenylbutyrate), should be terminated prior to AMMONUL® infusion.
Hyperammonemic coma (regardless of cause) in the newborn infant should be aggressively treated while the specific diagnosis is pursued. All patients should be promptly hemodialyzed as the procedure of choice using the largest catheters consistent with the patient's size. A target blood flow of 150 mL/min/m^2 may be attained using a 7F catheter. (Ammonia clearance [mL/min] is similar to the blood flow rate [mL/min] through the dialyzer). Clearance of ammonia is approximately ten times greater by hemodialysis than by peritoneal dialysis or hemofiltration. Exchange transfusion is ineffective in the management of hyperammonemia. Hemodialysis may be repeated until the plasma ammonia level is stable at normal or near normal levels.
AMMONUL® infusion should be started as soon as the diagnosis of hyperammonemia is made. Treatment of hyperammonemia also requires caloric supplementation and restriction of dietary protein. Non-protein calories should be supplied principally as glucose (8-10 mg/kg/min) with Intralipid added. Attempts should be made to maintain a caloric intake of greater than 80 cal/kg/d. During and after infusion of AMMONUL®, ongoing monitoring of neurological status, plasma ammonia levels, clinical laboratory values, and clinical responses are crucial to assess patient response to treatment. The need for other interventions to control hyperammonemia must be considered throughout the course of treatment. Patients with a large ammonia burden or who are not responsive to AMMONUL® administration require aggressive therapy including hemodialysis (see WARNINGS).
AMMONUL® must be diluted with sterile Dextrose Injection, 10% (D10W) before administration. The dilution and dosage of AMMONUL® are determined by weight for neonates, infants and young children, and by body surface area for larger patients, including older children, adolescents, and adults (Table 3). Maintenance infusions may be continued until elevated plasma ammonia levels have been normalized or the patient can tolerate oral nutrition and medications.
AMMONUL® solutions are physically and chemically stable for up to 24 hours at room temperature and room lighting conditions. No compatibility information is presently available for AMMONUL® infusion solutions except for Arginine HCl Injection, 10%, which may be mixed in the same container as AMMONUL®. Other infusion solutions and drug products should not be administered together with AMMONUL® infusion solution. AMMONUL® solutions may be prepared in glass and PVC containers. AMMONUL® solutions should be inspected visually for particulate matter and discoloration before administration.
[See table 3 below]
Arginine Administration:
Intravenous arginine is an essential component of therapy for patients with carbamyl phosphate synthetase (CPS), ornithine transcarbamylase (OTC), argininosuccinate synthetase (ASS), or argininosuccinate lyase (ASL) deficiency. Because a hyperchloremic acidosis may ensue after high-dose arginine hydrochloride administration, plasma levels of chloride and bicarbonate should be monitored and appropriate amounts of bicarbonate administered.
Pending a specific diagnosis, intravenous arginine (6 mL/kg of Arginine HCl Injection, 10%, over 90 minutes followed by the same dose over 24 hours) should be given to hyperammonemic infants suspected of having a urea cycle disorder for two reasons: 1) infants with deficiencies in enzymes of the urea cycle (apart from arginase deficiency) are usually arginine-deficient; 2) hyperammonemia in infants with ASS or ASL deficiency usually respond favorably to arginine administration. If deficiencies of ASS or ASL are excluded as diagnostic possibilities, the intravenous dose of arginine HCl should be reduced to 2 mL/kg/d Arginine HCl Injection, 10%.
Converting To Oral Treatment:
Once elevated ammonia levels have been reduced to the normal range, oral therapy, such as sodium phenylbutyrate, dietary management and protein restrictions should be started or reinitiated.

Table 3. Dosage and Administration

Patient Population	Components of Infusion Solution AMMONUL® must be diluted with sterile dextrose injection 10% at ≥ 25 mL/Kg before administration.		Dosage Provided		
	AMMONUL®	Arginine HCl Injection, 10%	Sodium Phenylacetate	Sodium Benzoate	Arginine HCl
0 to 20 kg:					
CPS and OTC Deficiency					
Dose Loading: over 90 to 120 minutes Maintenance: over 24 hours	2.5 mL/kg	2.0 mL/kg	250 mg/kg	250 mg/kg	200 mg/kg
ASS and ASL Deficiency					
Dose Loading: over 90 to 120 minutes Maintenance: over 24 hours	2.5 mL/kg	6.0 mL/kg	250 mg/kg	250 mg/kg	600 mg/kg
> 20 kg:					
CPS and OTC Deficiency					
Dose Loading: over 90 to 120 minutes Maintenance: over 24 hours	55 mL/m^2	2.0 mL/kg	5.5 g/m^2	5.5 g/m^2	200 mg/kg
ASS and ASL Deficiency					
Dose Loading: over 90 to 120 minutes Maintenance: over 24 hours	55 mL/m^2	6.0 mL/kg	5.5 g/m^2	5.5 g/m^2	600 mg/kg

HOW SUPPLIED

AMMONUL® (sodium phenylacetate and sodium benzoate) Injection 10% / 10% is supplied in single-use glass vials.
NDC-62592-720-50 single use vial containing 50 mL of sodium phenylacetate and sodium benzoate injection 10% / 10%.
Storage: Store at 25°C (77°F), excursions permitted to 15°-30°C (59°-86°F).

KEEP OUT OF REACH OF CHILDREN

Non-pyrogenic.
Rx Only

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