Hyperammonemic Encephalopathy and Lipid Dysmetabolism in a Critically Ill Patient after A Short Course of Amiodarone

Maximilien Cappe¹, Philippe Hantson¹, Mina Komuta², Marie-Françoise Vincent³, Pierre-François Laterre¹, Ismail Ould-Nana¹

¹ Department of Intensive Care, Cliniques St-Luc, Université catholique de Louvain, 1200 Brussels, Belgium
² Department of Pathology, Cliniques St-Luc, Université catholique de Louvain, 1200 Brussels, Belgium
³ Department of Clinical Chemistry, Cliniques St-Luc, Université catholique de Louvain, 1200 Brussels, Belgium

INTRODUCTION

Amiodarone is a class III antiarrhythmic agent that is widely used in clinical practice to treat supraventricular and ventricular arrhythmias. Among the adverse effects of amiodarone, acute hepatotoxicity is a rare but potentially fatal complication [1-6]. A case of severe but reversible metabolic disturbances occurring soon after a short-course of intravenous amiodarone in a critically ill patient, is described.

The clinical pattern was distinct from the hepatic failure exceptionally reported after a short-course of intravenous amiodarone. This acute liver dysfunction may also have been precipitated by co-existing factors such as acquired carnitine deficiency, severe obesity, and a long-term course of pancreatitis complicated by abdominal infections.

CASE REPORT

A 39-year-old woman weighing 87 kg weight and a body mass index of 34.9 kg/m², was admitted to the intensive care unit (ICU) of a tertiary teaching hospital. (Cliniques universitaires St-Luc, Brussels, Belgium), for an acute pancreatitis secondary to biliary perforation after endoscopic resection of a duodenal ampulla. The patient had a past medical history of Gardner syndrome (a form of familial adenomatous polyposis associated with mutations in the \( \text{APC} \) gene) and had previously undergone total colectomy and unilateral adrenalectomy. At the time of admission, laboratory data did not reveal any change in lipid metabolism or in liver function tests including ammonemia. However, two years previously, diffuse liver steatosis had been suspected following a computed tomography examination. Lipid profile was normal at that time.

The clinical course over the first weeks of hospitalisation was characterized by multiple episodes of pancreatitis-related intra-abdominal infections that were treated with antimicrobial therapy, surgery or percutaneous drainage.

Enteral nutrition was started immediately following admission but was limited by a slow gastric emptying and a reduced intestinal transit. Parenteral nutrition was introduced on hospital day twenty-eight. On day thirty-three, the patient presented with recurrent episodes of paroxysmal atrial fibrillation and intravenous amiodarone hydrochloride (CORDARONE*, Sanofi, DOI: 10.2478/jccm-2019-0026
Diegem, Belgium) was started with a loading dose of 300 mg and a maintenance dose of 450 mg over a single day. Two days after the loading dose of amiodarone, the patient became encephalopathic and hyperammonemia was documented. The medications administered over the week preceding amiodarone hydrochloride prescription are listed in Table I.

At the same time, there was a significant rise in liver enzymes, namely alkaline phosphatase and gamma-glutamyl transferase, and an increase in plasma triglycerides (Table II).

Parenteral nutrition had been stopped the day before amiodarone had been prescribed. Amino acids were measured by ion-exchange chromatography and post-column derivatization with ninhydrin, and spectrophotometric detection using a Biochrom 30 amino acid analyser™ (Biochrom LTD, Cambridge, UK).

A significant increase in glutamine plus glutamate, consistent with the observed hyperammonemia was recorded. A significant decrease in citrulline was also noted. Organic acids were measured by gas chromatography (Hewlett Packard 5973™, Littleton, CO 80127 USA) after extraction from urine by standard ethyl acetate diethyl ether extraction and derivatization to trimethylsilyl (TMS) derivatives. Lactic and pyruvic aciduria was recorded, as was a moderate hyper excretion of 3-hydroxybutyric acid. Arterial lactate increased to 15 mmol/L.

The treatment for this outcome consisted of carnitine supplementation (Cliniques St-Luc Hospital Pharmacy, Brussels, Belgium) 100 mg/kg as a single dose, and sodium benzoate (Cliniques St-Luc Hospital Pharmacy, Brussels, Belgium) administration, 12 g/day for seven days. The total daily dose of glucose infusion was

| Table I. List of the medications prescribed over the week preceding amiodarone hydrochloride prescription |
|---------------------------------------------------------------|
| **Medications**                                               |
| Vancomycin (Mylan, Hoeilaart, Belgium)                        |
| Ceftazidim (Kefadim®, Eurocept Pharmaceuticals, Ankeeven, The Netherlands) |
| Piperacillin-tazobactam (Mylan, Hoeilaart, Belgium)           |
| Amoxicillin (Clamoxyl®, Sandoz, Vilvoorde, Belgium)           |
| Metronidazole (B Braun Medical, Diegem, Belgium)              |
| Cefuroxime (Zinacef®, GlaxoSmithKline Pharmaceuticals, Wavre, Belgium) |
| Temocillin (Negaban®, Eumedica SA, Manage, Belgium)           |
| Insulin (Actrapid®, Novo Nordisk, Brussels, Belgium)          |
| Norepinephrine (Aguettant SA/NV, Brussels, Belgium)           |
| Nadroparine (Fraxiparine®, Movianto, Aspen, USA)              |

| Table II. Laboratory data before Day 0, and after amiodarone administration |
|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|
|                     | Day 0 | Day 2 | Day 3 | Day 4 | Day 5 | Day 6 | Day 7 | Day 8 | Day 9 | Day 10 | Day 11 | Day 12 |
|---------------------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| GGT (NV=<40 IU/L)   | 65    | 126   | 305   | 700   | 1172  | 1640  | 1874  | 1507  | 1520  | 1160  | 1267  | 990   |
| Alkaline phosphatase (NV=35-105 IU/L) | 125 | 250   | 369   | 540   | 659   | 734   | 736   | 636   | 626   | 560   | 673   | 549   |
| AST (NV=13-35 IU/L) | 29    | 114   | 80    | 35    | 28    | 23    | 24    | 20    | 22    | 22    | 21    | 17    |
| ALT (NV=7-35 IU/L)  | 15    | 19    | 41    | 56    | 57    | 50    | 41    | 40    | 37    | 35    | 37    | 37    |
| Total bilirubin (NV=<1.2 mg/dL) | 0.9 | 1.1   | 1.1   | 1.3   | 1.4   | 1.3   | 1.2   | 1.2   | 1.1   | 0.9   | 0.9   | 0.9   |
| INR (NV=0.80-1.20) | 2.09  | 2.11  | 1.57  | 1.59  | 1.47  | 1.47  | 1.36  | 1.35  | 1.32  | 1.32  | 1.26  | 1.27  |
| Ammonaemia (NV=<90 µg/dL) | -   | 179   | 167   | 155   | 139   | 153   | 99    | 79    | 83    | 111   | 63    |       |
| Triglycerides (NV=<150 mg/dL) | -   | -     | 454   | 438   | 498   | 608   | 604   | -     | 444   | 338   | 303   | 223   |
| Arterial lactate (NV=<2.2 mmol/L) | -   | 10.4  | 2.9   | 2.0   | 1.3   | 1.4   | 1.9   | 2.0   | 1.5   | 1.4   | 1.9   | 1.9   |
| Urine lactate (NV=<50 mmol/mmol creatinine) | -   | 1190  | -     | -     | -     | -     | -     | -     | -     | -     | -     | -     |
| Parenteral nutrition | Stop | -     | -     | -     | -     | -     | -     | -     | -     | -     | -     | -     |
| Enteral nutrition | No | No     | 500   | No     | 500   | 500   | 500   | 500   | 500   | 500   | 500   | 500   |
| Glucose supplementation | Yes | Yes   | Yes   | Yes   | Yes   | Yes   | Yes   | Yes   | Yes   | Yes   | Yes   | Yes   |

GGT, gamma-glutamyl transferase; AST, aspartate aminotransferase; ALT, alanine aminotransferase; INR, International Normalized Ratio. Enteral nutrition (Peptamen HN, Nestlé® HealthScience): proteins 20% (33 g/500 ml), glucose 47%, lipids 33% (24.5 g/500 ml, 70% of fat as medium-chain triglycerides). Enteral nutrition consisted of Peptamen HN (Nestlé® HealthScience Belgique, Brussels, Belgium): proteins 20% (33 g/500 ml), glucose 47%, lipids 33% (24.5 g/500 ml, 70% of fat as medium-chain triglycerides).
increased, with additional supplementation of hypoglycemic glucose to control episodes of hyperglycemia.

Three days after the administration of amiodarone, the enteral nutrition was progressively reintroduced.

Continuing correction of metabolic disorders characterized the further biological course. Parenteral nutrition, including a moderate dose of lipids, was resumed on day fourteen, to complete enteral nutrition. This approach was well tolerated.

**Discussion**

In addition to the common metabolic disorders observed during the course of acute pancreatitis complicated by multiple infections, this patient presented less common metabolic manifestations. The primary documented disorders were hypertriglyceridermia, hypoglycemia, hyperlactatemia and hyperammonemia. These findings were suggestive of disorders of fatty acids oxidation (FAO), hypertriglyceridermia and hypoglycemia, together with inhibition of the mitochondrial respiratory chain (hyperlactatemia) and urea cycle (hyperammonemia).

The common causes for these disorders were investigated and particularly the potential role of drug prescription and nutritional status. Hypertriglyceridermia and hyperammonemia were associated with a rise in liver enzymes, mainly alkaline phosphatase, and hepatic dysfunction was therefore suspected.

Oral long-term use of amiodarone may be complicated by an asymptomatic rise in serum transferase concentration in up to 25% of treated patients [7]. The steatotic pattern of drug-induced liver injury is well-known in patients with long-term administration of amiodarone and occurs after several months or years of treatment [8]. Steatosis during amiodarone therapy may result from impaired β-oxidation [8-9]. By contrast, acute hepatotoxicity following intravenous loading appears exceptional, and the possible role of the solvent has been discussed [10-15]. Usually, liver injury is rapidly reversible after discontinuation of amiodarone [2]. However, amiodarone can also cause fulminant hepatic failure, especially following a high-dose intravenous administration. Lethal cases presented massive hepatic necrosis on their histology, which was not observed in the current case [1-6].

Drugs associated with mitochondrial toxicity and failure of aerobic metabolism usually produce microvesicular steatosis with minimal inflammation and necrosis at liver biopsy [16]. These changes may develop within days of the administration of aspirin) or weeks in the case of linezolid. By contrast, amiodarone is usually considered to cause chronic liver injury. Amiodarone can induce steatohepatitis-like liver damage which can eventually lead to cirrhosis [17]. Amiodarone-induced liver injury is associated with both microvesicular steatosis and macrovascular steatosis as a consequence of impaired mitochondrial function and inhibition of fatty acids oxidation [8,9].

Amiodarone is an amphiphilic drug with protonable amine moiety that favours its accumulation inside the mitochondrial matrix. In amiodarone, the benzofuran-phenyl methanone moiety could be the chemical structure responsible for mitochondrial dysfunction [18]. The drug has been shown to have a dual action on fatty acids oxidation [18-19]. Amiodarone, at least in some in vitro models, is a direct inhibitor of carnitine-palmitoyl transferase 1 (CPT1) [20]. This would result in a reduced availability of acylcarnitine, with in turn, reduced plasma ketone bodies, accumulation of plasma acylcarnitine derivatives and urine dicarboxylic acids, and severe hypoglycemia [17,22]. The decrease of acetyl-CoA formation would also reduce the availability of N-acetyl glutamic acid (NAGA), a substrate for carbamoyl phosphate synthetase (CPS1), and reduce ammonia clearance. Amiodarone could also impair fatty acids oxidation by the inhibition of the mitochondrial respiratory chain activity at the level of complexes I and II [19-21] It is also to be noted that in our patient, the endogenous synthesis of carnitine could have been reduced by malabsorption, sepsis and organ failure [23].

The possible role of pre-existing liver steatosis and obesity may also be considered [24,25]. Obese individuals may present a higher risk of drug-induced liver injury, but this is unlikely in the case for amiodarone [26]. Additionally, a review of the literature suggests that, among the medications prescribed to the patient during the week before the onset of the most recent and unexpected metabolic disorders, no drug other than amiodarone had the ability to inhibit CPT1.

In the present case report, the mechanism causing hyperammonemia remains speculative (Figure 1). Ammonia production by intestinal bacteria was unlikely as previously the patient had a total colectomy. The absence of an inborn error of metabolism in the urea cycle was also verified. Carnitine is also indirectly required for the proper functioning of the urea cycle.
The synthesis of N-acetyl glutamic acid (NAGA), an important cofactor of carbamoyl phosphate synthetase (CPS1), produced from acetyl-CoA and glutamate by NAGA synthetase, is decreased.

Finally, there is no documented relationship between anomalies in the FAO or urea cycle and Gardner’s syndrome.

**Conclusion**

In conclusion, even a short course of amiodarone therapy may disturb some metabolic pathways, mainly FAO. The reasons why the present patient was so susceptible to this remains speculative as amiodarone is widely and safely prescribed in ICU patients with various medical conditions. Genetic predisposition could not be excluded. Other factors are probably involved and this would be particularly the case for critically ill patients presenting with associated causes for acquired carnitine deficiency such as sepsis or a catabolic state, or pre-existing disorder, including non-alcoholic liver fatty disease.

Reversibility was observed after amiodarone discontinuation.

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**Fig. 1. (Adapted from [22,23]). Possible mechanisms of amiodarone-induced hypertriglyceridemia and hyperammonemia.** Amiodarone is recognized as a potential inhibitor of carnitine-palmitoyl transferase 1 (CPT1). This will result in a reduced availability of acylcarnitine, with impaired beta-oxidation and acetyl-CoA production. Amiodarone also impairs the mitochondrial respiratory chain activity at the level of complexes I and II. The decrease in acetyl-CoA may also reduce the availability of N-acetyl glutamic acid (NAGA), a substrate for carbamoyl phosphate synthetase (CPS1).
INFORMED CONSENT

Written consent was obtained from the patient.

CONFLICTS OF INTEREST

The authors declare that there is no conflict of interest regarding the publication of this article.

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