Motilitone toxicity in a dog

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A three-month-old, intact male Maltese dog was presented to the hospital with lethargy after taking a human medication, Motilitone. Physical examination, including a neurological examination, revealed no remarkable findings, but cholinergic crisis symptoms appeared gradually. Blood and radiological examinations showed no remarkable findings. The dog was tentatively diagnosed with a cholinergic crisis associated with Motilitone intake. Treatment included intravenous administration of atropine (0.02 mg/kg) every 30 minutes and supportive fluid therapy. After 12 hours of treatment, the patient's clinical signs were resolved. This is the first case report describing Motilitone toxicity in a dog.

Keywords: cholinergic crisis; functional dyspepsia; herbal combination; Motilitone; toxicity

Cholinergic crisis refers to a clinical status manifested by the overstimulation of muscarinic or nicotinic receptors at the neuromuscular junction caused by the excessive accumulation of acetylcholine (ACh). The excessive accumulation of ACh is usually caused by the inhibition of the acetylcholinesterase (AChE), an ACh-degrading enzyme [1]. The most common causes are the excessive use of medications for myasthenia gravis or exposure to organophosphates, such as herbicides, insecticides, and other pesticides [1]. The over-accumulation of ACh at the neuromuscular junction causes symptoms related to the muscarinic and nicotinic receptors [1,2]. Clinical findings related to the muscarinic receptors are parasympathetic symptoms, such as miosis, bradycardia, bronchospasm, bronchorrhea gastrointestinal (GI) secretion, lacrimation, and urination. The symptoms related to the nicotinic receptors are skeletal and respiratory muscular weakness, tachycardia, muscle fasciculation, and flaccid paralysis. Death may result from respiratory compromise and paralysis. The clinical findings also involve stimulation of the central nervous system, resulting in agitation, dullness, seizure, coma, and ataxia [1-3].

Functional dyspepsia (FD) is a complex of symptoms recognized as discomfort in the upper abdomen and includes recurrent upper abdominal pain, excessive fullness, and bloating [4,5]. Drugs, such as metoclopramide, a dopamine antagonist 1st generation drug, domperidone, a dopamine antagonist 2nd generation drug, and cisapride, a serotonin agonist 3rd generation drug, were developed to treat FD [5]. However, several side effects of the drugs have been reported [4].

Motilitone was developed in Korea in 2011 as a 4th generation human FD treatment and was formulated as a combination of Corydalis Tuber (the root of Corydalis yanhusuo) and Pharbitidis Semen (the seed of Pharbitis nil). Motilitone's ac-
tivity mechanism has been revealed through many studies. Its gastric fundus relaxation effect appears to be related to the increased gastric capacity by 5-HT\textsubscript{1A} agonism [4], whereas its visceral analgesic effect appears to be related to the antinociceptive effect of visceral stimuli through adrenergic \alpha\textsubscript{2} agonism [5]. Dopamine causes inhibition of GI motility, but Motilitone is a D\textsubscript{2} antagonist that promotes GI motile functions [6]. In addition, Motilitone’s 5-HT\textsubscript{1A} agonism is related to the enhanced GI motility and visceral analgesia [5].

To the best of our knowledge, there are no reports on the effect of Motilitone in veterinary medicine. This case study describes the management of a cholinergic crisis in a young dog that consumed Motilitone, which induces parasympathetic activity.

A three-month-old, intact male 1.3 kg Maltese dog was presented to the Veterinary Teaching Hospital of Kangwon National University with lethargy after consuming Motilitone (Motilitone pill form; Dong-A Pharm. Co., Ltd., Korea). Physical and neurological examinations revealed no remarkable findings on the first visit. However, over time, the dog’s heart rate decreased (bradycardia), postural reflexes decreased, and the cranial nerve response gradually weakened. Other clinical symptoms, such as pinpoint miosis, salivation, tachypnea, and persistent vocalization, were also presented. Blood and radiological examinations showed no remarkable findings. The dog was diagnosed tentatively with acute cholinergic crisis based on the history and the symptoms related to the enhanced parasympathetics. Treatment started with stabilization of the airway, breathing, and circulation. Oxygen supplementation, warming, and supportive intravenous (IV) fluid therapy (0.9% NaCl) were provided. To overcome the cholinergic crisis, atropine administration was performed. After the first 0.04 mg/kg IV atropine bolus administration, 0.02 mg/kg IV atropine was administered at 30-minute intervals, six times, until no cholinergic crisis-related symptoms were detected. During treatment, the patient’s vitals were monitored in real-time. No side effects from atropine, such as tachycardia or mydriasis, were induced. Twelve hours after the first atropine treatment, most of the clinical symptoms had improved, and the patient left the hospital in a healthy state one day after treatment.

For a definitive diagnosis, the concentrations of free ACh and prolactin in the patient’s serum and that of four healthy male young (1-3 years) dogs were measured and compared. Client consent to testing the serum of all of the study dogs was obtained. All experimental protocols in this study were based on ethical procedures and were approved by the Kangwon National University Institutional Animal Care and Use Committee (approval number: KW-201117-1).

The levels of free ACh [7] and prolactin [8] in the serum were estimated using ELISA kits (MBS011705 for free ACh, MBS2604244 for prolactin; MyBioSource, Inc., USA). For ELISA, the serum dilution factors were 1:100 in the ACh ELISA and 1:1 in the prolactin ELISA. All ELISA procedures were conducted according to the manufacturer’s instructions. Upon completion of the ELISA procedures, the plate wells were read at 450 nm using a SpectraMax ABS Plus Microplate Reader (Molecular Devices, LLC, USA). Serum was obtained from four healthy dogs, and the average free ACh level was 20.969 ± 9.903 nmol/L in healthy dogs. Before treatment, the patient’s ACh level was 31.771 nmol/L, which was higher than the healthy group average. Twelve hours after treatment, the patient’s ACh level was 23.903 nmol/L. Twenty-four hours after treatment, the patient’s ACh level was 19.391 nmol/L, which was similar to the average level of the healthy group (Fig. 1).

The average prolactin concentration in the four healthy dogs was 2.383 ± 0.323 ng/mL. Before treatment, the patient’s prolactin level was 2.811 ng/mL, which was higher than the average of the healthy group. Twelve hours after treatment, the patient’s prolactin level was 2.684 ng/mL. Twenty-four hours after treatment, the level was 2.311 ng/mL, which was similar to the average level of the healthy group (Fig. 2).

Motilitone is a widely used botanical drug for treating FD in humans in Korea and is made from Corydalis Tuber and Pharbitidis Semen. Both of those materials have been used traditionally in Korea, China, and Japan for digestive diseases.

Previous studies have shown that Corydalis Tuber and Pharbitidis Semen, the raw materials of Motilitone, have AChE inhibition effects [9,10]. The suppression of AChE results in the accumulation of ACh in the brain and other tissues, and it may appear in blood [11].

Dopamine is a hormone that suppresses the secretion of prolactin. Therefore, the action of Motilitone as a D\textsubscript{2} receptor antagonist can lead to hyperprolactinemia. Kwon and Son [5] examine blood prolactin levels after the oral administration of Motilitone and reported that the concentration of prolactin in the blood increased with an increasing drug dosage in rats.

The present study confirmed that Motilitone caused an initial increase in ACh and prolactin in the patient’s blood followed by decreases in the concentrations of ACh and prolactin over time. Based on the results of this study, a diagnosis of a cholinergic crisis caused by Motilitone toxicity was made. The diagnosis was supported by the evidence of Motilitone poisoning obtained by measuring the serum concentrations of ACh and prolactin.

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Several studies have examined the effects of Motilitone in animal models. A gastric prokinetic effect was observed in rats [12] and dogs [12,13], a gastric relaxation effect was confirmed in dogs [6,12] and rats [4,6], and a visceral analgesic effect was demonstrated in rats [14]. Various doses have been applied in studies evaluating the efficacy of FD treatments through animal models. In the reported studies, doses of 0.03 to 10 mg/kg orally (PO) in rats and 0.1 to 3 mg/kg PO in dogs were used, and no
side effects were observed. The Motilitone manufacturer (Dong-A Pharm. Co., Ltd.) reported that after repeated administration for 26 weeks in rats, the no observable adverse effect level (NOAEL) was 150 mg/kg and the LD50 (lethal dose 50%) was greater than 2,000 mg/kg as a single administration in rats. In dogs, it was reported that the NOAEL was 100 mg/kg/day when administered repeatedly for 13 weeks and 200 mg/kg/day with repeated administration for 26 weeks [4]. In this study's patient, the ingested dose was approximately 46 mg/kg, which is less than that reported in a previous toxicity study. On the other hand, considering the patient's age-related metabolic function status, it was presumed that the ingested dose was sufficiently toxic. Additional veterinary safety, toxicity, and dosage studies of Motilitone should be undertaken.

The possible treatment options for a cholinergic crisis are administering anticholinergic drugs, such as atropine, catecholamines, such as dopamine, dobutamine, noradrenaline, and adrenaline, or performing hemodialysis or mechanical ventilation in the case of inhaled gases [1,3]. Atropine is an anti-muscarinic anticholinergic drug that competes with ACh on post-synaptic muscarinic receptors and is used widely to treat cholineric crisis in human medicine [3,15]. In this patient, atropine was administered along with supportive care, with the atropine dosage based on the occurrence of organophosphate poisoning and in accord with the clinical signs [15].

Cholinergic crisis is a life-threatening condition that rarely occurs in human and veterinary fields. This case study is the first report of a cholinergic crisis in a young dog that occurred after taking Motilitone, a human FD medication.

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