Delirium After Chloralhydrate in A Child– A Case Report

Çocukta Kloralhidrat Sonrası Deliryum - Olgu Sunumu

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ÖZ


Anahtar Kelimeler: kloralhidrat, deliryum, çocuk

ABSTRACT

Chloralhydrate is a drug that is used as sedative - hypnotic drug for imaging methods where analgesia is not required and sedation is sufficient in children. It is widely used because of its wide therapeutic index, relatively low respiratory depression, ease of administration and less frequent side effects. Although delirium was mentioned as the side effect of chloral hydrate in many places, no case report could be detected. In this case report, a child who developed delirium after the therapeutic use of chloralhydrate was mentioned.

Keywords: chloralhydrate, delirium, pediatric

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INTRODUCTION

It is difficult to achieve immobility in children in situations such as radiological imaging, tooth extraction, EEG, ECG, and suturing. In such cases, the use of chloral hydrate for sedation is still practiced in many centers (1). Chloral hydrate is a drug used as a sedative-hypnotic drug in imaging methods that do not require analgesia and sedation is sufficient in children. It is widely used because of its broad therapeutic index, relatively low respiratory depression, ease of administration, and fewer side effects. The preferred dose is 25 to 50 mg/kg up to a maximum of 1 g (2). Chloral hydrate, a commonly used sedative in children, has many serious side effects in high or toxic doses. Some of those: prolonged sedation, paradoxical excitement, delirium, and unpleasant taste sensations (3).

Delirium is defined as a disturbance in attention, awareness, and cognition over a short period of time that is not explained by a pre-existing neurocognitive disorder, or a decreased level of arousal attributed to a medical condition, intoxication, withdrawal, or a medication side effect (4).

Although delirium is mentioned in many places as a side effect of chloral hydrate, no case report has been found. Our case is a child who developed delirium after administration of chloral hydrate.

CASE REPORT

Written consent has been obtained from the patient's relatives.

A 5.5-year-old girl with a weight of 20 kg was admitted to the outpatient clinic due to occasional nonspecific headache. Brain magnetic resonance imaging of the patient was requested, but chloral hydrate 50 mg/kg/dose was administered 2 times, one hour apart, for sedation so that he would not move during the procedure. Half an hour after the second dose, there was a slowing in her speech and movements, numbness in her hands and legs. Afterwards, she has acted in a manner that hurts herself and her surroundings, has been rebellious and she was brought to the emergency room of the pediatric emergency department with these complaints. The patient was not making eye contact with his mother. The child's actions weren't purposeful, she was unaware of her surroundings, restless and inconsolable. We measure/observe delirium with The Pediatric Anesthesia Emergence Delirium (PAED) scale in our clinic. Her PAED score was 18.

With the recommendation of the poison counselor, after the active charcoal was given, the patient was admitted to the service for 24-hour follow-up. All laboratory tests, systemic and neurological examinations were normal. She had agitation for 2-3 hours before and she had euphoric behaviors for 8-9 hours. During this period, vital signs were stable. She started to behave completely normal in about 12-13 hours. She discharged without problems.

DISCUSSION

In children, delirium is an altered state of consciousness, which occurs with anesthesia and continues until early recovery. Awareness or attention disorder about the child's environment and manifests itself as disorientation, hyperactive behavior and hypersensitivity in the acute period after anesthesia (5).

Since the late 1800s, chloral hydrate, has been widely used in pediatric sedation, especially in newborns and infants for the last three decades. Also in our clinic, chloral hydrate is used routinely if necessary. Chloral hydrate can be administered orally or rectally and absorbed from the gastrointestinal tract with the highest serum concentrations within 30-60 minutes. In pediatric patients, the dose of chloral hydrate is 25 to 50 mg/kg and up to 1 gram (1, 2). In vivo, it converts to trichloroethane, which is a sedative and active form. Its half-life is about 8-12 hours, but its acute overdose and in intoxications half-life can be up to 35 hours. It is recommended not to be used in children older than 4 years or in children with neurodevelopmental disorders due to the increased risk of adverse effects and treatment failure (1).

From 1996 to 2009, the Institute for Safe Drug Administration (ISMP) stated that there were 8 deaths and a number of misuse related use of faulty chloral hydrate for pediatric sedation. In 2014, the institute reported that a 4-year-old girl could not be awakened from sleep for a long time after chloral hydrate from 70 mg/kg for dental treatment and was ex-home. In the report, it was state that a 3-year-old child was given 6000 mg chloral hydrate accidentally, after vomiting, respiratory arrest,
intubated and monitored and again it was stated that 100 mg / kg dose was given to another patient for 15 months, stridor and apnea developed after vomiting, followed up for 12 hours and discharged (1).

Chloralhydrate is no longer recommended for sedation in children and is not available in many countries, including the United States. Some countries have removed chloralhydrate from national health formulas because of their potential carcinogenicity (6).

Due to the delayed initial action of chloralhydrate, long duration of action and the high frequency of side effects, given the availability of better alternatives, the use of chloralhydrate is no longer recommended (7, 8). Italy and France were also banned in 2016, in the United States in 2012 due to carcinogenicity and genotoxicity (9).

There are studies suggesting that preoperative anxiety decreases with oral chloral hydrate and has little effect on postoperative delirium and postoperative maladaptive behavior (2). In some studies, 4% to 7% of participants were reported that other adverse effects such as sedation failures (20%), prolonged sedation (19%), excessive drowsiness, airway obstruction, respiratory depression, hypercapnia, post-discharge sedation, respiratory arrest, hypotension and cardiopulmonary arrest (1). The risk of resedation that lasts longer than 24 hours may occur at any age in children, even in therapeutic doses (10).

In Uptodate and the drug package insert, it is indicated that it has some side effects even when taken in appropriate doses. These; In cardiovascular system: Atrial arrhythmia, depression of myocardial contractility, hypotension, torsades de pointes, shortening of refractory periods, ventricular arrhythmia. In the central nervous system: Abnormal gait, ataxia, confusion, delirium, dizziness, drowsiness, drug addiction (if used for a long time), hallucinations, hangover effect, weakness, nightmares, paradoxical excitation, vertigo. Dermatologic: Skin rash. Endocrine and metabolic: Acute porphyria, ketonuria. In the gastrointestinal tract: diarrhea, gas, stomach irritation, nausea, vomiting. Hematological and oncologic: acute porphyrria, eosinophilia, leukopenia. In the eyes: Allergic conjunctivitis, blepharoptosis, keratooconjunctivitis. In Ear: Increased middle ear pressure (infants and children). In the respiratory system: Airway obstruction (young children), laryngeal edema (children). Potential cardiotoxic effects such as ventricular dysrhythmias and hypotension may occur at high or overdose. Unfortunately, there is no known antidote of chloral hydrate (1).

Delirium in children is usually for a short time after recovery from anesthesia (5), but our patient developed delirium after chloral hydrate without any sedation.

In our patient, delirium developed after 100 mg / kg chloralhydrate, activated charcoal was given and hydration was provided. There were no abnormalities in blood values and vital signs during follow-up.

CONCLUSION

Chloralhydrate is prohibited in many countries. In many countries, it is one of the most frequently used sedative drugs that are used as controlled. However, due to the long duration of the effects, the lack of controlled follow-up, and the serious side effects leading to death, even if given in the appropriate dose, the unit with intensive care conditions should be given under clinical supervision and then followed up.

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REFERENCES


