ACUTE LAMOTRIGINE OVERDOSE IN ADULTS: A CASE REPORT

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INTRODUCTION
Self-treatment with Lamotrigine rarely ends with toxicity, regardless of the suicidal intentions of the patient. The authors hereby present an illustrative case of the patient who has been treated with epilepsy therapy with Dandy-Walker syndrome and congenital epidermolysis bullosa (potentially skin-unwanted). Lamotrigine is a phenyltriazine-class, broad-spectrum antiepileptic and therapy of bipolar depression. Dandy-Walker syndrome is a pathological entity and represents the set of developmental, cerebral, but also other abnormalities of the organism. Epidermolysis bullosa is a hereditary, non-inflammatory skin disease with a mucous membrane of characteristic “bubbles”.

Case report: Our patient, a 37-year-old male was first admitted to the hospital department of Urgent Medicine of Clinical Center Kragujevac because he consumed two boxes of Lamotrigine tablets. In the receiving clinic, the patient showed respiratory failure and was urgently intubated. From medical documentation and hetero-anamnesis (obtained by his father), the authors found out that he was treated for epilepsy, Dandy-Walker syndrome, and congenital epidermolysis bullosa, which deteriorated with the use of Lamotrigine through potentially undesirable skin effects. During clinical observation, a lavage of gastric contents was conducted. The medical coal was used via nasogastric intubation as a detoxification method because of the patient’s comatose state. Causative metabolic pathway of lamotrigine, the hemodialysis was performed.

Conclusion: The case report of our patient points to the necessity of a multidisciplinary approach of the expert team, consisting of the clinical pharmacologist and toxicologist, neurologist, dermatologist, nephrologists, and other specialists, if necessary. Patients with Dandy-Walker syndrome require adequate socio-medical care.

Keywords: Lamotrigine, acute overdose, Dandy-Walker syndrome, congenital epidermolysis bullosa, clinical manifestations, diagnostic and therapeutic interventions
tion of anti-epileptics (respiratory depression, cardiac arrest, coma, and death) which required endotracheal intubation as an effective procedure in overdose cases (1). This syndrome diagnosis (her father provided diagnosis) was followed by clinical symptoms, that have previously been reported as epilepsy, intellectual incompetence, and other. The case report illustrates the complexity of the observation of the syndrome in an adult male, which presents the diagnostic and the therapeutic challenge in acute overdose with Lamotrigine. The constellation of symptoms and structural abnormalities of the brain should be taken into consideration when dealing with patients with neuropsychiatric manifestations, systemic diseases, and specific skin diseases, especially if diagnosed in childhood and adolescence and chronically treated with Lamotrigine.

**CASE REPORT**

A patient of 37 years was conducted as an emergency admission to The Department of Urgent Medicine of Clinical Center Kragujevac in Kragujevac because he had consumed two boxes of Lamotrigine tablets (overdose). From his medical documentation and hetero-anamnesis, the authors discovered that he was treated for epilepsy, Dandy-Walker syndrome, and congenital (epidermolysis bullosa), which worsened with the use of Lamotrigine, through undesirable skin effects.

At admission, the patient was disoriented, agitated, a-febrile of medium osteomuscular build and nutrition, no signs of hemorrhagic syndrome and peripheral adenopathy. Auscultatory over lungs weakened respiratory murmur. Rhythmic heart action, well-audible tones, no noise; the abdomen was soft, palpably painless, without defense, and peritoneal reaction. Bullous changes bleeding when touched were seen on the mucous membrane of the oral cavity, two erosions covered with thin crusts were seen on the inner side of the left hand, and one bulla filled with serous contents was observed in the pubic area (Figure 1).

**Figure 1.** Epidermolysis bullosa (Lamotrigine-overdose), as potentially undesirable skin effects.

Initially, due to the patient’s non-cooperation, it was not possible to perform an electrocardiographic recording, which was subsequently performed and shallow negative T-waves from V1 to V4 were recorded, with sinus rhythm and heart rate 90/min. Immediately, after the examination, breathing ceased, and the patient was urgently intubated and connected to an assisted ventilation device. Biochemical laboratory tests were regular except for inflammations parameters. The gas analyses indicated hypoxemia (Table 1).

During the clinical observation, the lavage of gastrointestinal contents was performed. Medical coal was used via nasogastric intubation as a detoxification method since the patient was in a comatose state. Causative metabolic pathway of Lamotrigine the double hemodialysis with the minimal ultrafiltration was also performed.

Additionally, a computerized tomography (CT) scan of the brain discovered: cerebral hemiatrophy including cerebellum hypo-atrophy with skull thickening/widening of the vermis (dysgenesis) and communication of ventricle (IV) with arachnoid-cisternal spaces in fossa crani posterior (Figure 2).

As for clinical characteristics, a radiography of the heart and lungs performed several times showed regular findings. The concentration of Lamotrigine in blood was determined at a reference institution in Belgrade of Serbia (toxicological laboratory of Military Medical Academy), which was elevated to a toxic concentration of 20.82 g/l (reference value 4-11 g/l).

On the fourth day of hospitalization, the state of consciousness stabilized, and gas exchange normalized, after which the patient was separated from assisted ventilation and extubated. The patient had normal vital signs: blood pressure was 110/55 mm Hg, heart rate was 54/min, respiratory rate was 15/min, and the temperature was about 36°C. Ophthalmological signs showed minor nystagmus, degenerative myopia, and characteristic retinal defects with vitreous fenestrated membranes in the retinal periphery. The virus tests were negative, as well as the psychoactive substance test. Due to clinical improvement and normalization of all basic-laboratory analyzes, so no control level of Lamotrigine was performed. Finally, since Lamiktal® tablets (brand name of medicament) can cause skin changes exacerbating its condition, the doctors decided to opt for gradual exclusion of this medicament while continuing the treatment of epilepsy with Kepra® tablets (brand name). To treat depression, Elice® tablets (brand name) were introduced. The patient was discharged for 16 days of hospitalization in good general condition.
DISCUSSION

Our case reported the illustrates-nature of Dandy-Walker syndrome and congenital epidermolysis bullosa in an adult male and represented the diagnostic and therapeutic challenge in acute overdose of Lamotrigine. Our patient was of depressed mood resulting in suicidal ideation/suicidal attempts and developed the status of epileptic.

The spectrum of clinical effects of Lamotrigine in acute overdose was not precisely established. Several cases of overdose had serious effects such as coma, respiratory depression, and intraventricular conduction disturbances.

Serum evaluation revealed high Lamotrigine levels without any other etiology for mental dysfunction. After the prompt supportive treatment with early intubation, use of potassium chloride for hypokalemia, and the administration of sodium bicarbonate, the condition of the overdosed patients improved (2). Encephalopathy is the secondary cause of serious Lamotrigine-toxicity, as the clinical manifestation (3). Lamotrigine-overdose was usually benign, mild, or with no toxicity, but large exposures were associated with severe central nervous system depression, cardiac conduction delays, seizures, and death (4).

Other toxic effects include hypersensitivity reactions, QRS-complex prolongations, rhabdomyolysis, serotonin syndrome, seizures, and/or agitation (5), with these effects of the other author’s opinion, we also agree. Today, many document treatment-refractory Lamotrigine cardiotoxicity among dogs by applying intra-lipid emulsion therapy (6). Dandy-Walker malformation or syndrome occurs sporadically. Some patients remain clinically asymptomatic for years, while others may exhibit a variety of co-morbidities leading to earlier diagnosis and multidisciplinary research. Treatments are generally focused on posterior fosse symptoms, often including surgical interventions, like ventriculoperitoneal and cystoperitoneal shunting (7), but the surgical possibility of intervention was excluded from our patient.

In case reports, many authors described the neuropsychological and behavioral profile of patients, usually in adult males (8), as in our case report.

Brain computed tomography and brain magnetic resonance imaging have shown cyst in posterior fosse, hydrocephalus, hypoplasia of corpus callosum, syringomyelia, absence of cerebellar vermis, etc. Surgery involving arachnoid adhesiolysis and endoscopic third ventriculostomy was performed (9).

The progression of associated morbidities in Dandy-Walker syndrome requires an early, multidisciplinary diagnosis, so that clinically asymptomatic cases, over the many years (10,11), would not remain undiagnosed, as in our case.

Bullous dermatitis in infants and adults is a clinical term used for several disorders associated with primary neonatal pemphigus. The common symptoms of the disorder regardless of etiologic factors are redness of skin and formation of bubbles of various sizes filled with serous or serous-bloody content. Bursting bubble patches peel off, leaving bare, sometimes oozing surface (12). Bullous dermatitis in neonates and adults associated with primary neonatal pemphigus should be ruled out as a differential diagnosis. It is characterized by irritation and hyperemia of dry skin, with the formation of bubbles with serous or serous-bloody contents, which are emptied and moisturize skin (12).

Immune-Fluorescence-Antigen-Mapping testing is morphological verification of diagnosis and targeted genetic analysis of the mutations by the molecular
AKUTNO TROVANJE LAMOTRIGINOM KOD ODRASLIH: PRIKAZ SLUČAJA

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Sažetak

Uvod: Samotrovanje lamotriginom je retko trovanje, nezavisno od suicidne namere bolesnika. Autori prikazuju ilustrativni slučaj bolesnika, koji je lečen (potencijalno kožno-nezeljenom terapijom) za epilepsiju, Dandy-Walkerovim sindromom i kongenitalnom buloznom dermolizom. Lamotrigin je antiepileptik klase feniltriazina u terapiji bipolarne depresije. Dandy-Walkerov sindrom je patološki entitet i predstavlja skup razvojnih, moždanih, ali i drugih abnormalnosti organizma. Bulozna epidermolize je nasledna, nezapaljena bolesti kože i sluzokože sa karakterističnim “mehurićima”.

Prikaz bolesnika: Naš bolesnik, star 37 godina prvi put je primljen u hospitalni odsek Centra za urgentnu medicinu, Kliničkog Centra Kragujevac u Kragujevcu, jer je konzumirao dve kutije Lamotrigina tableta U prijemnoj ambulanci bolesnik pokazuje respiratornu insuficijenciju, prestanak disanja i u besvesnom stanju je hitno intubiran.

Iz referente medicinske dokumentacije i heteroanamneze (od njegovog oca) saznajemo da je lečen od epilepsije, Dandy-Walkerovim sindromom i kongenitalne bulozne dermolize, koja se se sa terapijom Lamotriginom pogoršala, kroz potencijalno neželjeno dejstvo na koži. U toku kliničke observacije urađena je lavaža želudačnog sadržaja. Kao metod detoksikacije primenjen je medicinski ugalj putem nazogastrične sonde zbog komatoznog stanja bolesnika. Uzročno metabolitičkom putu lamotrigina, sprovedena je hemodializa.

Zaključak: Prikaz slučaja ovog bolesnika ukazuje na neophodnost multidisciplinarnog pristupa tima eksperta, u sastavu kliničkog farmakologa i toksikologa, neurologa, dermatologa, nefrologa i drugih specijalista. Pacijenti sa Dandi-Walkerovim sindromom zahtevaju adekvatnu društveno medicinsku brigu.

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Table 1. The gas, blood and laboratory analyzes (biochemical parameters).

| Gas analyzes | Blood analyzes | Laboratory analyzes |
|--------------|----------------|---------------------|
| Ph 7.24      | Le 11.10       | Glycosa 5.1         |
| pO₂ 6.2      | Er 3.95        | Urea 1.6            |
| pCO₂ 3.7     | Hgb 124        | Kreatinin 72        |
| Na 139       | Hct 0.327      | CK 195              |
| K 3.3        | Tr 182         | CRP 15.5            |
| Ca 1.14      | PTT 36.3       |                     |
| HCO₃ 19.3    | INR 1.590      |                     |
| SpO₂ 87%     | Albumin 44     |                     |

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