

## Baclofen-induced toxic encephalopathy in end-stage renal disease: should we be more careful?

Abdul S. Mohammed, Abdalla Hassan\*, Prajwal Boddu

Advocate Illinois Masonic Medical Center, Department of Internal Medicine, Chicago, Illinois

### ABSTRACT

Baclofen is a highly used centrally acting GABA agonist that continues to be an effective therapy for symptomatic relief of skeletal muscle spasm and spasticity in traumatic spinal cord lesions, multiple sclerosis, cerebral palsy, stroke and chronic hiccups and has recently become popular as recreational drug. The renally dependent excretion determines the circulating concentrations and guides effective dosing to decrease adverse reactions. Caution should be considered in administering baclofen to patients with decreased renal function. We present a patient with end stage renal disease on hemodialysis with recent baclofen ingestion who presented with toxic encephalopathy that was resolved with additional dialysis sessions.

**Keywords:** Baclofen; Encephalopathy; Dialysis; End-stage renal disease

### Introduction

Baclofen (parachlorophenyl gamma-aminobutyric acid) is an agonist of the neurotransmitter gamma-aminobutyric acid (GABA) [1] with centrally functioning mechanism of action has been used to treat spasticity and more recently chronic hiccups [2, 3]. Baclofen is well absorbed in the gastrointestinal tract producing the determined therapeutic levels of 80–400 ng/mL [4]. The primary excretion is through the kidney (70–80%) with the remainder metabolized in the liver or processed through the gastrointestinal tract [4]. Only a small portion is able to cross the blood brain barrier, creating the desired effects. The half-life is approximately 6.8 hours [3]. Patients with decreased renal function have an extended half-life with more crossing of the blood brain barrier. The generalized central nervous system (CNS) depression effects are then further pronounced producing fatigue, syncope, hypotension, ataxia, psychological disturbances, and cardiovascular and respiratory depression [2, 4]. In patients with limited renal function, there have been reports of toxic side effects with the initial dosing of 5 mg three times a day within only a few days [5] and baclofen effects can be prolonged in these patients. Baclofen toxicity has been shown to resolve with

hemodialysis resembling excretion rates similar to normal renal function [6]. Here we report a case of baclofen-induced encephalopathy in a dialysis patient with recovery following hemodialysis sessions.

### Case presentation

Our patient is a 73-year-old female with a past medical history of end stage renal disease on hemodialysis, diabetes mellitus type 2, hypertension, coronary artery disease status post coronary artery bypass graft, hyperlipidemia and peripheral vascular disease status post angioplasty who presented for altered mental status. The patient had a recent primary care physician visit with concern for low back pain due to muscle spasm and was started on baclofen 10 mg twice a day. On the day of admission, the patient was found to be speaking incoherently, repeatedly mumbling words like “okay” and “hold on”, confused and unable to concentrate. The vital signs were the following: temperature 99 °F, respiratory rate 18 breaths/minute, pulse 81 beats/minute, blood pressure 142/88 mmHg and pulse oxygenation 99% on room air. On initial presentation, the patient was lethargic but arousable and unable to answer questions. She was not alert or oriented to herself or her surroundings. She did not have fever, headache, neck stiffness, nausea, vomiting, seizures or head injury. She had not traveled recently or had any sick contacts. Laboratory tests showed the following: sodium 138 mEq/L, potassium 4.3 mEq/L,

\*Correspondence

**Dr. Abdalla Hassan**

Advocate Illinois Masonic Medical Center, Department of Internal Medicine, Chicago, Illinois

**E Mail:** [abdalla.hassan@advocatehealth.com](mailto:abdalla.hassan@advocatehealth.com)

**Mohammed et al**

ASIAN PACIFIC JOURNAL OF HEALTH SCIENCES, 2016; 3(1): 53-55

[www.apjhs.com](http://www.apjhs.com)

chloride 97 mEq/L, bicarbonate 29 mEq/L, blood urea nitrogen 42 mg/dL, creatinine 4.9 mg/dL. The white blood cell count was 8.5 mg/dL, hemoglobin 11.1 mg/dL, and platelet count 237/mm<sup>3</sup>. Hepatic function panel had a total bilirubin of 0.7 mg/dL, AST 29 u/L, ALT 39 u/L, alkaline phosphatase 120 u/L, creatine kinase 350 u/L, thyroid stimulating hormone 1.4 and ammonia 23 u/L. Her arterial blood gas was normal and toxicology screen was negative. A head computed tomography and MRI was performed with no acute intracranial abnormalities and chest x ray was normal as well. Compliance with her last hemodialysis session was confirmed. With high suspicion for baclofen-induced encephalopathy, the patient was admitted for emergent dialysis. She received hemodialysis for two sessions with significant improvement in mental status without any residual symptoms. Upon discharge, the patient had baclofen discontinued

### Discussion

Baclofen was originally found to be effective in treatment of spasticity concerning spinal cord lesions in the 1960s [7]. The mechanism of action is a centrally acting presynaptic agonist of GABA to enhance tone reduction and reduce spasticity [1]. In recent studies, it has been shown effective in treating persistent hiccups [3]. Baclofen is efficiently absorbed in the gastrointestinal tract and with variations of the half-life dependent largely on renal function being the primary mechanism of excretion [4]. A high correlation between renal clearance and creatinine clearance was shown to exist in a excretion kinetic study [8]. Baclofen is a moderately lipophilic molecule and with increased circulating concentration crossing the blood brain barrier has a correlating amplified centrally acting response [9]. These adverse reactions include hypotension, bradycardia, respiratory depression, and toxic encephalopathy [5, 9]. There have been previous cases with chronic renal failure and symptoms of neurotoxicity. Serum concentrations of baclofen were measured to have progressively decreased during the sessions of hemodialysis with resolution of the neurologic sequel [5]. The effectiveness of hemodialysis was additionally seen in another case of baclofen overdose with normal renal function. In this case, a patient consumed 200 mg of extended release baclofen with subsequent ventilator dependent respiratory failure. Two sessions resolved the side effects with successful extubation and resolution of additional symptoms [9]. In a study of the efficiency in using hemodialysis, the filtration rate for baclofen was as effective as normal kidneys [6]. This high efficiency

of filtration is because only 30% of the circulating concentration is bound to protein [6]. In case reports, there is a several-hour delay prior to improvement in symptoms with the theorized necessity of redistribution from the CNS to the intravascular system and filtration by dialysis [10]. Interestingly, the therapeutic range of baclofen 80–400 ng/mL has been found to cause negative side effects in patients with impaired renal function. The exact therapeutic level with the intended decrease in dosing remains unclear with continued susceptibility to adverse side effects [11]. Electroencephalography (EEG) has been used in initial evaluations for the etiology of unconsciousness in baclofen overdose. These changes include pseudo periodic sharp waves to periodic high-amplitude discharges and generalized slow waves [12]. Resolution of these EEG changes was seen following treatment [13]. These changes did not indicate the capacity to induce epilepsy and have been recommended not to treat with antiepileptic medications [13]. Baclofen renal dosing continues to be nonspecific with renal adjustment recommendations. A study in 2011 highly recommended changes in labeling with improved guidelines in dosing. There was a proposal to avoid this medication with a GFR < 30 mL/min/1.73 m<sup>2</sup>. This was further detailed with a decrease in dosing and titrating in a GFR > 30 to 60 mL/min/1.73 m<sup>2</sup> [14]. In following the reported history of decreased renal function and side effects, this continues to be a highly appropriate recommendation. Our case represents toxic encephalopathy with inappropriate dosing of baclofen that was resolved with hemodialysis. Special consideration is required with this renally excreted medication and more specific guidelines would likely further alleviate incidence of side effects.

### Conclusion

Special consideration is required with this renally excreted medication and more specific guidelines would likely further alleviate incidence of side effects. It's paramount that physicians are aware of such an interaction in ESRD patients.

### Reference

1. R. Gerkin, S. C. Curry, M. V. Vance, P. W. Sankowski, and R. D. Meinhart. First-order elimination kinetics following baclofen overdose. *Annals of Emergency Medicine*. 1986; 15(7): 843–846.

2. A. Dario and G. Tomei. A benefit-risk assessment of baclofen in severe spinal spasticity. *Drug Safety*. 2004; 27(11): 799-818.
3. C. Zhang, R. Zhang, S. Zhang, M. Xu, and S. Zhang. Baclofen for stroke patients with persistent hiccups: a randomized, double-blind, placebo-controlled trial. *Trials*. 2014; 15: 295.
4. R. R. Young and P. J. Delwaide. Drug therapy: spasticity (second of two parts). *The New England Journal of Medicine*. 1981; 304 (2): 96–99.
5. C.-L. Chou, C.-A. Chen, S.-H. Lin, and H.-H. Huang. Baclofen-induced neurotoxicity in chronic renal failure patients with intractable hiccups. *Southern Medical Journal*. 2006; 99(11): 1308–1309.
6. M. Brvar, M. Vrtovec, D. Kovač, G. Kozelj, T. Pezdir, and M. Bunc. Haemodialysis clearance of baclofen. *European Journal of Clinical Pharmacology*. 2007; 63(12):1143–1146.
7. N. E. F. Cartlidge, P. Hudgson, and D. Weightman. A comparison of baclofen and diazepam in the treatment of spasticity. *Journal of the Neurological Sciences*. 1974; 23(1): 17–24.
8. E. W. Wuis, M. J. M. Dirks, E. F. S. Termond, T. B. Vree, and E. Van der Kleijn. Plasma and urinary excretion kinetics of oral baclofen in healthy subjects. *European Journal of Clinical Pharmacology*. 1989; 37(2): 181–184.
9. L. S. Dias, G. Vivek, M. Manthappa, and R. V. Acharya. Role of hemodialysis in baclofen overdose with normal renal function. *Indian Journal of Pharmacology*. 2011; 43(6): 722–723.
10. K.-S. Chen, M. J. Bullard, Y.-Y. Chien, and S.-Y. Lee. Baclofen toxicity in patients with severely impaired renal function. *Annals of Pharmacotherapy*. 1997;31(11): 1315–1320.
11. V.-C. Wu, S.-L. Lin, S.-M. Lin, and C.-C. Fang. Treatment of baclofen overdose by haemodialysis: a pharmacokinetic study. *Nephrology Dialysis Transplantation*. 2005;20(2): 441–443.
12. B. Sauneuf, H. K. Totouom, B. Savary et al., Clinical and EEG features of acute intrathecal baclofen overdose. *Clinical Neurology and Neurosurgery*. 2012;114(1): 84–86.
13. T. Fakhoury, B. Abou-Khalil, and B. Blumenkopf. EEG changes in intrathecal baclofen overdose: a case report and review of the literature. *Electroencephalography and Clinical Neurophysiology*. 1998; 107(5): 339–342.
14. A. El-Husseini, A. Sabucedo, J. Lamarche, C. Courville, and A. Peguero. Baclofen toxicity in patients with advanced nephropathy: proposal for new labeling. *American Journal of Nephrology*. 2011; 34(6):491–495.

**Source of Support: Nil**

**Conflict of Interest: None**