Sodium Valproate versus Continuous Infusion of Haloperidol in Management of Agitated Critically Ill Patients

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Abstract

AIM: Describe the efficacy and safety of valproate and haloperidol infusion in controlling agitation in the intensive care unit (ICU).

MATERIAL AND METHODS: Prospective study on 100 critically ill patients with agitation in Kasralainy Hospital over the period from May 2016 to June 2017. Patients were divided into two groups, each group included 50 patients, 1st group patients received Depakene orally, and 2nd group patients received haloperidol by i.v infusion for 72 h. Richmond agitation sedation score and doses of additional sedative drugs were noted and calculated daily in the first three days.

RESULTS: Our study showed that valproate was equal in efficacy in controlling agitation; decreasing the RAAS significantly after 48 h from initiation (2.52 ± 0.61 vs 0.28 ± 0.54 with p < 0.001) for Depakene and (2.6 ± 0.67 vs 0.34 ± 0.48 with p < 0.001) for haloperidol. There was also a decrease in the doses of additional sedative drugs used to control agitation (midazolam & propofol) after 48 h from drug initiation. Both drugs therapy was associated with decrease in heart rate (89 ± 20 vs 86.6 ± 13.6 with p = 0.002 for valproate and 99.8 ± 23.3 vs 91 ± 16.7 with p < 0.001 for haloperidol). They did not affect blood pressure. Haloperidol therapy was associated with significant QTc prolongation.

CONCLUSION: Valproate was equal in efficacy as haloperidol infusion in controlling agitation in ICU and decreasing the doses of additional sedative drugs used after 48 h from initiation.

Introduction

Agitation occurs in up to 70% of critically ill patients and is a significant source of distress for patients, families, and providers (Fraser GL et al., 2000). Sedatives are administered to 50% of intensive care unit (ICU) patients to alleviate agitation [1]. Choice of sedative is complex and largely driven by patient context. No sedative has consistently been shown to be superior to the rest, and alternative agents are greatly needed [2]. Most ICU patients, especially those requiring mechanical ventilation, are treated with opioids, propofol, and/or benzodiazepines [1], [3]. Use of these agents is limited by adverse effects (e.g., hemodynamic derangement) for safe administration [4]. New therapies for treating agitation are rarely introduced into practice, with dexmedetomidine being the most recent in 1999. Consequently, providers have increasingly repurposed older pharmacologic agents as ICU sedatives (e.g., clonidine, Phenobarbital, and valproate (Depakene) [5], [6].

Haloperidol is a butyrophenone that works by blocking D2 receptors, probably in the mesolimbic region [7]. Its side effects include extrapyramidal symptoms, and rarely, the neuroleptic malignant syndrome [8]. The patient should be monitored for precipitation of arrhythmias such as torsade de points [9] and haloperidol should be used with caution in patients with a QTc interval of over 450 msec. National guidelines for the management of agitation/delirium recommend short term haloperidol if non-pharmacological measures are not effective.

Depakene is an antiepileptic and mood stabiliser approved for the treatment of seizures, manic episodes associated with bipolar disorder, and migraine prophylaxis. Mechanistically, it blocks voltage-dependent sodium and calcium channels, increases γ-aminobutyric acid (GABA) synthesis, potentiates GABA activity at postsynaptic receptors, blocks GABA degradation, and attenuates the activity of glutamate upon N-methyl-D-aspartate receptors [10], [11].
Recently, valproate has been administered to critically ill patients to treat agitation and delirium, but there are few published reports to support this practice [12], [13], [14]. Valproate is an emerging treatment for ICU agitation because it allows patients to interact with their caregivers; can be administered outside of the ICU; has both an intravenous (IV) and enteral formulation; has a low drug acquisition cost; and has not been associated with respiratory depression, hemodynamic derangements, or delirium. In this study, we describe the use of Depakene & haloperidol for agitation in critically ill patients and examine their safety.

We aimed to evaluate the adding effect of a calcium sensitiser (levosimendan) compared to the conventional inotropic and vasoactive agent used in the patient with poor left ventricular function undergoing cardiac surgery on different measured hemodynamic variables and the effect on the outcome.

Material and Methods

Study design

Our study was a prospective one on Agitated critically ill patients in Kasr Alainy hospital, critical care department over the period from May 2016 to June 2017. Patients included if they were above 18 years old and have severe agitation (score ≥ 2 on RAAS) patients were excluded if they have advanced cardiac diseases, hepatic diseases, advanced malignancy or if they were on antiepileptic. Our study was conducted on 100 patients divided in two groups; each group included 50 patients. Patients in group I received Depakene orally, and patients in group II received haloperidol by i.v. Infusion for 72 h.

Demographic data and patients’ characteristics

Patient demographic data included age, sex, weight, history of psychiatric diagnosis, alcohol or substance abuse, the reason for ICU admission, need for mechanical ventilation. Clinical outcomes were descriptive and included ICU length of stay and ICU mortality.

Efficacy outcomes

Efficacy data were collected from day of drug (valproate or haloperidol) initiation and continuing for 72 h or until discontinuation, whichever came first. The 3-day interval was selected to allow a reasonable time to observe efficacy or lack thereof, the agitated patients were examined daily and Richmond Agitation-Sedation Scale (RASS) was assessed daily from drug initiation up to 3 days. Also, the doses of additional sedative drugs were noted and documented daily during drug therapy.

Safety outcomes

Safety parameters were examined for hospital duration of valproate and haloperidol therapy. Records were specifically reviewed for possible hepatotoxicity, hematologic toxicity for both drugs and QTc prolongation for haloperidol. The number of patients who had discontinued because of a suspected adverse event was also recorded. Hepatotoxicity was defined as a new alanine aminotransferase 3 times the upper limit of normal (120 U/L), alkaline phosphatase 2 times the ULN (234 U/L), total bilirubin 2 times the ULN (2 mg/dL), or a doubling of the baseline value if it was already abnormal following drug initiation. Suspected cases of hepatotoxicity were further assessed using the validated Roussel Uclaf Causality Assessment Method (RUCAM) [15]. Hematologic toxicity was defined as a new leukocyte count 4200 cells/mm3, absolute neutrophil count 2400 cells/mm3, or platelet count 140 000 cells/mm3 or platelet drop by 50% if platelets were already 140 000 cells/mm3 following valproate initiation.

Results

Demographic data and patient characteristics

One hundred patients were included in the study. The mean age of the studied population was 63.7 ± 15 years; the youngest was 22 years; they included 54 Males and 46 Females. Regarding Comorbidities; in our study 36 Smokers, 42 with pre-existing Cardiac patients (Ischemic heart disease and chronic heart failure) and 39 with pre-existing Renal patients (serum creatinine ≥ 1.4 mg/dl). End organ failure (45%), sepsis (40%) and Stroke (hemorrhagic and non-hemorrhagic) (15%) were the most common reasons for ICU admissions. The mean length of ICU stay was 7 days, 21 patients died in the ICU.

| Table 1: Baseline characteristic data of the total population |
|-------------------------|----------------|
| Age (mean ± SD) years   | 63 ± 15        |
| Gender-females n        | 46             |
| Weight (mean ± SD) kg   | 76.5 ± 14      |
| Smokers n (%)           | 36             |
| Pre existing Cardiac disease n (%) | 42 |
| Pre existing Renal disease n (%) | 39 |
| ICU Mortality (%)       | 21%            |
| ICU stay (mean) days    | 7              |
| Causes of admission     |                |
| End Organ failure (%)   | 45%            |
| Sepsis (%)              | 40%            |
| Stroke (%)              | 15%            |
Efficacy outcomes

Regarding valproate group; the severity of agitation decreased after 48 h from starting valproate therapy; RASS score decrease from 2.52 ± 0.61 to 0.28 ± 0.54 before and after depakene initiation respectively with P value < 0.001. The doses of concomitant sedative drugs used to control agitation (midazolam & propofol) decreased after 48h from drug initiation (77.2 ± 82.4 mg/day vs. 0 mg/day) and (3.5 ± 2.0 gm/day vs. 0.35 ± 0.49 gm/day) with P value 0.004 and 0.135 respectively. Regarding haloperidol group; the severity of agitation decreased after 48 h from starting valproate therapy; RASS score decrease from 2.6 ± 0.67 Vs 0.34 ± 0.48 before and after depakene initiation respectively with P value < 0.001. The doses of concomitant sedative drugs used to control agitation (midazolam & propofol) decreased after 48 h from drug initiation (101 ± 141 mg/day vs. 16 ± 33 mg/day) and (1.6 ± 1.9 gm/day vs. 0.33 ± 0.7 gm/day) with P value < 0.001.

Hemodynamic response

The Depakene group showed a statistically significant difference regarding HR after 48 h from valproate therapy (89 ± 20 vs 86.6 ± 13.6) with P-value 0.002, there was no effect on systolic or diastolic blood pressure. Haloperidol group showed a statistically significant difference regarding HR after 48 h from haloperidol (99.8 ± 23.3 vs 91 ± 16.7) with P-value < 0.001, there was no effect on systolic or diastolic blood pressure.

Safety outcomes

All patients had complete blood picture & liver function tests monitored during drug therapy. In valproate group; Hemoglobin level and Total leucocytic count showed significant decrease after 48 h from valproate therapy with p = 0.002 & 0.025 respectively. Liver enzymes (ALT & AST) showed a significant increase after 48 h from valproate therapy with p < 0.0001. One patient developed hepatotoxicity (elevation of ALT ≥ 120 IU/ml) but was unrelated to the drug as his RUCAM score was 3. There was significant QTc prolongation after 48 h from haloperidol initiation with p < 0.001, but this was not associated with major cardiac events (VT or VF).

<table>
<thead>
<tr>
<th>Lab</th>
<th>Day 0</th>
<th>Day 2</th>
<th>Day 0</th>
<th>Day 2</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hemoglobin g/ml</td>
<td>10.6 ± 1.9</td>
<td>10.2 ± 1.5</td>
<td>10.6 ± 1.9</td>
<td>10 ± 1.7</td>
</tr>
<tr>
<td>Total leucocytic count x 10^9</td>
<td>10.7 ± 9</td>
<td>9.2 ± 7</td>
<td>13 ± 8</td>
<td>10 ± 5</td>
</tr>
<tr>
<td>Platelets x 10^9</td>
<td>253 ± 103</td>
<td>245 ± 102</td>
<td>209 ± 88</td>
<td>212 ± 94</td>
</tr>
<tr>
<td>ALT IU/ml</td>
<td>44 ± 32</td>
<td>49 ± 100</td>
<td>49 ± 83</td>
<td>39 ± 36</td>
</tr>
<tr>
<td>AST IU/ml</td>
<td>39 ± 25</td>
<td>48 ± 126</td>
<td>47 ± 66</td>
<td>39 ± 33</td>
</tr>
<tr>
<td>Bilirubin IU/ml</td>
<td>0.62 ± 0.5</td>
<td>0.58 ± 0.47</td>
<td>1.32 ± 2.5</td>
<td>1.2 ± 2.3</td>
</tr>
</tbody>
</table>

Discussion

Regarding efficacy outcomes in Depakene & haloperidol groups of our study, there was a significant decrease in the severity of agitation after 48 h from drug initiation with Richmond agitation sedation scores p < 0.001. Also, there was a marked decrease in the doses of concomitant sedative drugs used to control agitation (Midazolam & propofol) after 48 h from drug initiation. Similarly, Gagnon et al. (2017) performed a retrospective cohort study evaluated critically ill adults treated with Depakene for agitation on Fifty-three patients and showed that the incidence of agitation significantly decreased following the initiation of Depakene from 96% to 61% on Depakene day 3 (P < 0.001). Also, in this study, Depakene therapy was associated with reduced doses of concomitant sedative drugs [16]. Riker et al. (1994) evaluated the Continuous infusion of haloperidol in critically ill agitated patients. This study was performed on eight patients required mechanical ventilation who had severe agitation which was refractory to intermittent bolus treatment with benzodiazepines, narcotics, and haloperidol. These patients received Continuous infusion of haloperidol to maintain adequate sedation. This study showed that there was a significant decrease in the Sedation-Agitation Scale after 48 h from haloperidol initiation (+2.4 vs +0.8) and (P = 0.06). Also, there was a marked reduction in the daily total of non-haloperidol sedatives after 48h of continuous infusion of haloperidol with (P-value = 0.15) [17].

Regarding Hemodynamics response after Depakene & haloperidol initiation, our study showed that there was significant decrease in patients heart rate after 48 h from drug initiation with p = 0.002 & < 0.001 respectively, blood pressure showed no significant changes before and after drug initiation. Sinha et al. (2000) reviewed hospital records of 13 patients with status epileptics and hypotension who received IV Depakene therapy. There were no significant changes in blood pressure, pulse, or use of vasopressors. The data suggest that Depakene
loading is well tolerated, even in patients with cardiovascular instability [18], also Riker, et al. (1994) evaluated the Continuous infusion of haloperidol in eight critically ill agitated patients. There were no hypotension episodes noted [17].

Regarding safety outcomes, our study showed that hepatotoxicity (ALT ≥ 120 IU/ml) had developed for one patient (2%) hepatotoxicity is unlikely to be related to the drug (Depakene or haloperidol) as RUCAM score was 1 and 3 respectively which make hepatotoxicity is unlikely related haloperidol [15]. Possible reasons for hepatotoxicity in these patients are the underlying disease.

QTc prolongation for all patients in the haloperidol group with one patient had significant QTc prolongation ≥ 500msec after 48 h from haloperidol therapy, but this QTc prolongation was not associated with major cardiac events. That agrees with Tisdale et al., who assessed the effect of intravenous haloperidol on QT interval dispersion in critically ill patients who received intravenous haloperidol for delusional agitation. QTc intervals were measured, and QT interval dispersion was calculated.

Haloperidol prolonged QTc interval compared to pretreatment values in Torsades de Pointes patients by a greater magnitude than in patients who did not experience Torsades de Pointes. It was concluded that intravenous haloperidol prolongs QTc intervals in critically ill patients [19].

Also, there was a significant decrease in Hemoglobin levels and total leucocytic count after 48 h from haloperidol therapy. Possible reasons for that decrease of the total leucocytic count are that TLC was elevated on day 0 of drug initiation due to sepsis which decreased with starting broad-spectrum antibiotics & control of sepsis. Also, the possible reasons for decreased level of hemoglobin are dilutional effect of intravenous fluids, blood sampling for lab and bone marrow suppression from sepsis.

References


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