Optimal glycemic control in neurocritical care patients

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See related research by Kramer et al, http://ccforum.com/content/16/5/R203

Abstract

Currently, the major issue in glycemic control in neurocritical care patients is that tight glycemic control (target range of 80 to 110 mg/dL) using intensive insulin therapy is associated with higher rates of hypoglycemia without an improvement in survival rate. The review by Kramer and colleagues in this issue of Critical Care confirms these data but provides solid evidence about the relationship between hyperglycemia and worsened neurological outcome after acute brain injury. In accordance with the conclusions of Kramer and colleagues, we recommend that a glucose control goal in neurocritical care patients be in the ‘moderate’ target range (110 to 180 mg/dL). In addition, we recommend adequate nutrition before and during insulin infusion, avoidance of insulin as a bolus, and the use of continuous insulin infusion, beginning with low doses with titration to individual sensitivity. Careful and accurate glycemic monitoring is especially important when insulin is infused.

The review and meta-analysis focusing on optimal glycemic control in neurocritical care (NCC) patients by Kramer and colleagues [1] in this issue of Critical Care provide a contemporary and comprehensive overview of randomized controlled trials (RCTs) that apply different glycemic control strategies in this challenging intensive care unit (ICU) population. Optimal glycemic control in critical care (CC) patients, in general, and NCC patients, in particular, has evolved dramatically over the past 15 years and remains under active investigation and debate about the ideal target range and impact of dysglycemia (high, low, and variable glucose levels) on outcome in the heterogeneous ICU population [2]. Prior to 2001, clinicians frequently applied a somewhat ‘permissive’ glycemic control approach in CC and NCC patients since hyperglycemia was considered a physiological response to stress or insult. Starting about a decade ago, the target range for glycemic control was the achievement of euglycemia (80 to 110 mg/dL) by using intense insulin therapy as an infusion in the hopes of reducing ICU and hospital morbidity and mortality [3]. Subsequently, the latter approach was shown to result in an increased, but varying, incidence of hypoglycemia [4]. The incidence appears to vary by study, patient population, glucose goal, and intensity of insulin infused. However, hypoglycemia develops also in some critically ill patients in the absence of infused insulin. Whether there is an unfavorable cause-and-effect impact on outcome in critically ill patients who become hypoglycemic remains under scrutiny.

Currently, a key issue with tight glycemic control (target range of 80 to 110 mg/dL) achieved by using intensive insulin therapy in NCC patients is the report of higher rates of hypoglycemia without an improvement in survival rate. The review by Kramer and colleagues in this issue of Critical Care confirms these data but provides solid evidence about the relationship between hyperglycemia and worsened neurological outcome after acute brain injury. In accordance with the conclusions of Kramer and colleagues, we recommend that a glucose control goal in neurocritical care patients be in the ‘moderate’ target range (110 to 180 mg/dL). In addition, we recommend adequate nutrition before and during insulin infusion, avoidance of insulin as a bolus, and the use of continuous insulin infusion, beginning with low doses with titration to individual sensitivity. Careful and accurate glycemic monitoring is especially important when insulin is infused.
(26% versus 27%, relative risk (RR) 0.99, 95% confidence interval (CI) 0.83 to 1.17, \( P = 0.89 \)). However, poor neurological outcome was less frequent in patients who received intensive glycemic management (58% versus 68%, RR 0.91, 95% CI 0.84 to 1.00, \( P = 0.04 \)). Hypoglycemia was markedly greater among treated patients (30% versus 14%, RR 3.10, 95% CI 1.54 to 6.23, \( P = 0.002 \)). No effects were detected in the incidence of nosocomial pneumonia, and other nosocomial infections were infrequently reported, and thus a conclusive statistical analysis was not possible.

The results of this review and meta-analysis confirm that tight blood glucose control in NCC patients does not reduce mortality but increases the rate of hypoglycemia. The new message from this review is that, in patients treated with an ‘active’ glucose control strategy with insulin infusion to maintain a blood glucose concentration (BGC) of less than 180 mg/dL, in comparison with those in whom ‘loose’ glycemia control was allowed (BGC of greater than 200 mg/dL before the start of insulin infusion), there is an improvement in neurological outcome.

Of relevance, the authors reported that information on nutritional status and nutritional support was reported poorly in the original articles but that ‘in most of the cases, tube feeding appeared to have been initiated as soon as possible’ \[1\].

In conclusion, optimal glucose control in NCC patients should include an active therapeutic strategy. Tight blood glucose control (with BGC target range of 80 to 110 mg/dL) exposes NCC patients to an increased risk of iatrogenic hypoglycemia. Patients with hypoglycemia (BGC values of less than 80 mg/dL) have increased mortality and the potential for worsened long-term functional status. On the other hand, patients with acute brain damage and permissive hyperglycemia with BGC values of greater than 180 mg/dL have a worsened neurological outcome.

During the last decade, there have been dramatic changes in our attitude toward glycemic control in NCC patients. It is now clear that this physiological variable cannot be overlooked and deserves committed clinical attention that is based on what we have learned. We now know that the time frame for manipulation of BGC is not as strict as that for hemodynamic variables and that sudden changes in BGC are potentially as dangerous as extreme BGC values \[8\]. We know that intense insulin infusion used to control glucose must be monitored appropriately and administered along with adequate enteral or parenteral nutrition and that iatrogenic hypoglycemia must be avoided. We also know that currently available clinical experience and technology are often derived from chronic management of patients with diabetes mellitus but that the unique characteristics of CC and NCC patients require new understandings and applications of technology.

Research areas that require further investigation include the following:

1. Assessment of the impact of glucose control differentiated by NCC-specific pathology. This requires more information on the impact of glucose homeostasis and control in specific NCC patient subgroups: traumatic brain injury, ischemic/hemorrhagic stroke, neuro-oncologic pathology, subarachnoid hemorrhage, and brain injury severity \[9-11\]. Of note is the study by Schlenk and colleagues \[12\], who used brain microdialysis in patients with subarachnoid hemorrhage to demonstrate the association of a target BGC of less than 110 mg/dL with increased acute brain metabolic derangements.

2. Mathematical models to describe and tailor an individual patient’s glucose sensitivity, the changes in glucose sensitivity over time and the relationship between changes in insulin sensitivity and the insulin/nutrition infusion protocol (amount of calories, access used for nutrition, amount and timing of insulin infusion) used in the individual patient.

3. Dedicated engineering technology, including continuous glucose monitoring devices with the application of closed loop insulin/nutrition infusion systems \[13-15\].

4. Greater understanding of the relationship between peripherally measured glucose, glucose in the healthy brain, and glucose in the injured or ischemic brain.

In accordance with the conclusions of Kramer and colleagues, we recommend that insulin infusion for glucose control in NCC patients be aimed at a ‘moderate’ target range (110 to 180 mg/dL). In addition, we would recommend an adequate nutrition support of NCC patients before and during insulin infusion, the avoidance of insulin boluses, and the use of continuous insulin infusions initially at low dose, titrated to individual sensitivity with the application of a standardized and easily applied glycemic monitoring protocol.

**Abbreviations**

BGC, blood glucose concentration; CC, critical care; CI, confidence interval; ICU, intensive care unit; NCC, neurocritical care; RCT, randomized controlled trial; RR, relative risk.

**Competing interests**

The authors declare that they have no competing interests.

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