Construction of Effective and Safe Glycemic Control in the Elderly by Using Continuous Glucose Monitoring

Koki Kakuta\textsuperscript{a, b}, Hiroki Adachi\textsuperscript{a, b}, Hidekatsu Yanai\textsuperscript{a, c}

To the Editor

Severe hypoglycemia is significantly associated with increased risks of a range of adverse clinical outcomes \cite{1}. An increased risk for developing dementia by older patients with diabetes who have had episode of severe hypoglycemia has been suggested \cite{2}. However, excellent evidence on glycemic control in the elderly is lacking, and optimal treatment needs to be constructed collaboratively with patients, incorporating the likelihood of benefits and harms and patients preferences about treatment in older adults with type 2 diabetes \cite{3}.

To show the usefulness of continuous glucose monitoring (CGM) to determine safe and effective glycemic control in the elderly, we present our old patient who showed asymptomatic severe hypoglycemia during sleeping. An 83-year-old type 2 diabetic female patient (body height, 151.4 cm; body weight, 56.7 kg; BMI, 24.7 kg/m\textsuperscript{2}) was admitted to our hospital due to poor glycemic control. She has been treated by ingestion of alogliptin (25 mg) before breakfast, and repaglinide (1.0 mg) before every meal, and injection of insulin glargine U300 (10 U) before breakfast. On admission, her plasma glucose and HbA1c level were 334 mg/dL and 8.7%, respectively.

We changed insulin treatment from insulin glargine U300 (10 U) to insulin degludec (IDeg)/insulin aspart (IAsp) (14 U) which is a soluble conformation of two distinct insulin analogues in the ratio of 70% ultra-long-acting IDeg and 30% rapid-acting IAsp \cite{4}. We stopped ingestion of repaglinide before breakfast because of the addition of IAsp before breakfast. We also switched from repaglinide (1.0 mg before lunch and dinner) to α-glucosidase inhibitor (α-GI), voglibose (0.3 mg before lunch and dinner).

We present the result of CGM which showed the effect of switching from repaglinide to voglbose on glycemic control in Figure 1. CGM showed hypoglycemia early at morning (from 2 to 6 o’clock in the morning) (Fig. 1a, b) during the treatment using repaglinide. After the switching to voglbose, hypoglycemia early at morning was disappeared (Fig. 1c, d). In addition, maximal glucose level and amplitude of glucose excursion decreased, in spite of reduction of insulin dose (Fig. 1d).

Repaglinide is a short-acting, insulin-releasing agent, and has been suggested to lessen the risk of long-lasting hypoglycemia \cite{5}, however, which was observed in our patient. We should consider the development of severe hypoglycemia in the elderly whose drug clearance is diminished when we use glinides. Furthermore, our observation indicated that α-GI does not induce long-lasting hypoglycemia and also ameliorates glycemic control in combination with insulin as compared with glinides.

We think that benefits and harms about treatment for diabetes depend largely on individuals, and individual differences get larger by aging. Therefore, we strongly recommend the construction of safe and effective glycemic control in the elderly by using CGM.

Conflicts of Interest

The authors declare that they have no conflicts of interest concerning this article.

References

Figure 1. Continuous glucose monitoring showing the effect of switching from repaglinide to voglibose for the treatment for type 2 diabetes in old women.