



Glucose transporter inhibitor-conjugated insulin mitigates hypoglycemia

Jinqiang Wang^{a,b}, Jicheng Yu^c, Yuqi Zhang^c, Anna R. Kahkoska^d, Zejun Wang^{a,b}, Jun Fang^a, Julian P. Whitelegge^e, Song Li^{a,f}, John B. Buse^d, and Zhen Gu^{a,b,g,h,1}

^aDepartment of Bioengineering, University of California, Los Angeles, CA 90095; ^bCalifornia NanoSystems Institute, University of California, Los Angeles, CA 90095; ^cJoint Department of Biomedical Engineering, University of North Carolina at Chapel Hill and North Carolina State University, Raleigh, NC 27514; ^dDepartment of Medicine, University of North Carolina School of Medicine, Chapel Hill, NC 27599; ^eThe Pasarow Mass Spectrometry Laboratory, The Jane and Terry Semel Institute for Neuroscience and Human Behavior, David Geffen School of Medicine, University of California, Los Angeles, CA 90095; ^fDepartment of Medicine, University of California, Los Angeles, CA 90095; ^gJonsson Comprehensive Cancer Center, University of California, Los Angeles, CA 90024; and ^hCenter for Minimally Invasive Therapeutics, University of California, Los Angeles, CA 90095

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Insulin therapy in the setting of type 1 and advanced type 2 diabetes is complicated by increased risk of hypoglycemia. This potentially fatal complication could be mitigated by a glucose-responsive insulin analog. We report an insulin-facilitated glucose transporter (Glut) inhibitor conjugate, in which the insulin molecule is rendered glucose-responsive via conjugation to an inhibitor of Glut. The binding affinity of this insulin analog to endogenous Glut is modulated by plasma and tissue glucose levels. In hyperglycemic conditions (e.g., uncontrolled diabetes or the postprandial state), the in situ-generated insulin analog–Glut complex is driven to dissociate, freeing the insulin analog and glucose-accessible Glut to restore normoglycemia. Upon overdose, enhanced binding of insulin analog to Glut suppresses the glucose transport activity of Glut to attenuate further uptake of glucose. We demonstrate the ability of this insulin conjugate to regulate blood glucose levels within a normal range while mitigating the risk of hypoglycemia in a type 1 diabetic mouse model.

drug delivery | diabetes | glucose-responsive | insulin | insulin analog

Diabetes mellitus affects more than 400 million people across the world (1–3). The treatment for type 1 and advanced type 2 diabetes is multiple daily injections or continuous infusion of exogenous insulin (1, 2, 4). However, the benefits of insulin therapy are often not fully realized, due to the risk of hypoglycemia associated with insulin overdose, which can result in seizure, coma, and death (5). Therefore, tremendous efforts have been devoted to the development of smart insulin delivery systems that mimic the glucose-dependent dynamic insulin secretion of β -cells, thereby reducing hyperglycemic condition and mitigating the risk of insulin-induced hypoglycemia associated with a miscalculated exogenous insulin dose (6, 7). To this end, chemically driven synthetic closed-loop insulin delivery systems integrating phenylboronic acid (7–16), glucose-binding protein (17–20), and glucose oxidase (21–26) have been extensively studied. However, synthetic strategies to tightly regulate blood glucose levels with a low risk of hypoglycemia remain elusive in clinical practice (27).

Here, we propose to expand the safety margin of insulin by simply conjugating insulin with a reversible glucose transporter (Glut) inhibitor. Glut is a family of transmembrane proteins that facilitate the transport of glucose across plasma membranes (28). Various compounds are able to competitively inhibit the glucose transport activity of Glut (29–31). Due to the presence of the Glut inhibitor, the insulin analog can reversibly and dynamically bind to Glut on cell membranes with an affinity modulated by surrounding glucose concentration, rendering the insulin molecule glucose-responsive (Fig. 1). Upon s.c. injection, this insulin analog can bind to insulin receptors (IR) as well as endogenous Glut, establishing an in situ-generated reservoir of insulin analog–Glut complex (Fig. 1). Upon a glucose challenge, such as the glucose rise associated with a meal, the insulin analog–Glut complexes dissociate to liberate free Glut on plasma membranes as well as free insulin analog into interstitial fluids and plasma.

The free insulin analog can subsequently bind to IR to trigger the translocation of Glut4 to cell membranes and enhance glucose clearance into muscle and fat. Meanwhile, the Glut, which is previously inaccessible to glucose as part of the insulin analog–Glut complex, can enhance the blood glucose clearance. In excess doses, the insulin analog induces overexpression of Glut on plasma membranes and subsequently triggers the glucose uptake by cells, which could potentially induce hypoglycemia; however, the formation of the glucose-responsive insulin analog–Glut complexes can suppress the glucose transport efficiency of Glut, therefore reducing the hypoglycemic risk.

Results

Glut-i2, a reversible Glut inhibitor with a low dissociation constant and high affinity for Glut4 and Glut1 (32), was selected for our study. Glut-i2 was integrated with a single terminal amino group to give Glut-i2–NH₂ (SI Appendix, Scheme S1 and Figs. S1–S5) (32). Next, Glut-i2–NH₂ was conjugated to insulin via a bifunctional linker (17) succinimidyl 4-(*N*-maleimidomethyl)cyclohexane-1-carboxylate to give insulin–Glut-i2 conjugate (designated *i*-insulin), which was confirmed by measuring the molecular weight via the matrix-assisted laser desorption/ionization with a time-of-flight analyzer (SI Appendix, Fig. S6), while the modification was validated on A1 of insulin (SI Appendix, Fig. S7).

To test the binding ability of the *i*-insulin toward Glut, it was labeled with sulfo-Cyanine 5 (Cy5-*i*-insulin) and used to treat erythrocyte ghost, a widely used Glut carrier (SI Appendix, Fig. S8) (33, 34). After 30-min incubation with *i*-insulin at room temperature, the erythrocyte ghosts showed a high fluorescence

Significance

Glucose-responsive insulin analogs or delivery systems are desirable for enhancing health and improving quality of life of people with diabetes. We describe here a simple strategy to engineer a long-acting insulin analog, which can establish an endogenous Glut-associated delivery reservoir of insulin that can modulate glucose metabolism in a blood glucose-dependent manner. Importantly, after subcutaneous injection, in vivo blood glucose regulation was validated in a type 1 diabetic mouse model with negligible hypoglycemia.

Author contributions: J.W., J.Y., and Z.G. designed research; J.W., J.Y., Z.W., J.F., and J.P.W. performed research; J.W., J.Y., Y.Z., A.R.K., S.L., J.B.B., and Z.G. analyzed data; and J.W., J.Y., Y.Z., A.R.K., S.L., J.B.B., and Z.G. wrote the paper.

Conflict of interest statement: J.W. and Z.G. have applied for patents related to this study.

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¹To whom correspondence should be addressed. Email: guzhen@ucla.edu.

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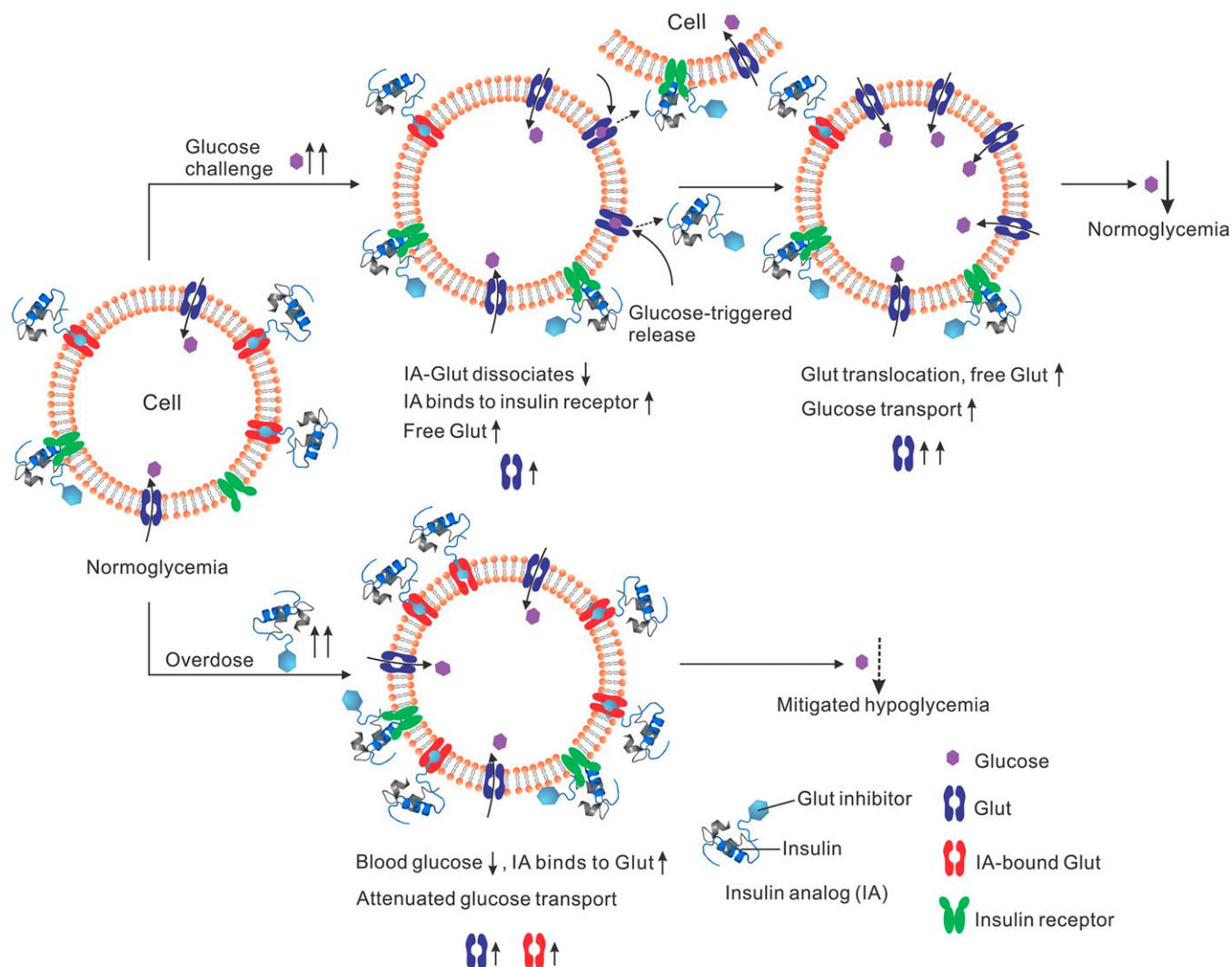


Fig. 1. Schematic of regulating the glucose-transport activity with insulin analog (IA; in this study, *i*-insulin serves as a model analog). Insulin analog can bind to Glut in a glucose-responsive manner. Upon injection and in normoglycemia, insulin analog achieves a regular blood glucose clearance rate, and an insulin analog–Glut complex reservoir is formed. Upon a glucose challenge, increased blood glucose levels result in the release of insulin analog from the insulin analog–Glut complex, which subsequently binds to IR to trigger the translocation of Gluts to cell membranes. With dissociation of insulin analog, glucose-inaccessible insulin analog-bound Glut becomes free Glut, enhancing excess blood glucose clearance. Upon an excess insulin analog injection (i.e., overdose), the formation of the insulin analog–Glut complex attenuates the glucose transport activity of Glut, therefore mitigating hypoglycemia risk.

intensity localized on the membranes, whereas the surrounding solvent showed a slightly weaker fluorescence (Fig. 2A). Control erythrocyte ghosts treated with Cy5-labeled native insulin only showed weak fluorescence comparable to the background (Fig. 2B). Moreover, the amount of Cy5-*i*-insulin bound to erythrocyte ghosts increased along with the increase of the concentration of free Cy5-*i*-insulin (SI Appendix, Fig. S9), and the K_d was measured as 13 nM (SI Appendix, Figs. S10 and S11 and Eqs. S1 and S2). The binding rate of *i*-insulin toward erythrocyte ghost was evaluated. Within 2 min of the addition of Cy5-*i*-insulin to the erythrocyte ghost solution, high fluorescence intensity localized on the erythrocyte membranes was observed (SI Appendix, Fig. S12A). A concurrent rapid decrease in the fluorescence intensity of the supernatant was also observed (SI Appendix, Fig. S13). Next, the in vitro release kinetics of *i*-insulin was investigated by diluting the Cy5-*i*-insulin–treated erythrocyte ghost solution and observing the fluorescent signal. A sharp decrease in fluorescence intensity was noted for all Cy5-*i*-insulin–treated erythrocyte ghosts within 2 min after a twofold, fourfold, and 10-fold dilution (SI Appendix, Fig. S12B).

Cy5-*i*-insulin–treated erythrocyte ghosts were further treated with glucose solutions at varying glucose concentrations of 0, 400, 800, and 1,600 mg/dL. The fluorescence intensity gradually decreased with increased glucose concentrations, consistent with the proposed dissociation of *i*-insulin–Glut complexes (Fig. 2C and D). The glucose-responsive dissociation of *i*-insulin–Glut complex was further tested at physiological-relevant concentrations (1 and 5 nM) of *i*-insulin (35). The concentration of *i*-insulin increased in the supernatant as the glucose concentration was increased (Fig. 2E and F). A 100% increase in the concentration of supernatant insulin was observed as the glucose concentration increased from 0 mg/dL to 400 mg/dL for both the 1- and 5-nM concentrations.

Next, the ability of *i*-insulin to regulate blood glucose levels was evaluated in the type 1 diabetic mice induced by streptozotocin (STZ). With s.c. injection, the *i*-insulin–treated mice sustained normoglycemia below 200 mg/dL for more than 10 h (Fig. 3A), whereas mice treated with native insulin showed less than 4 h of normoglycemia (Fig. 3A). Prolonged s.c. retention of *i*-insulin at the injection site compared with native insulin was observed (SI

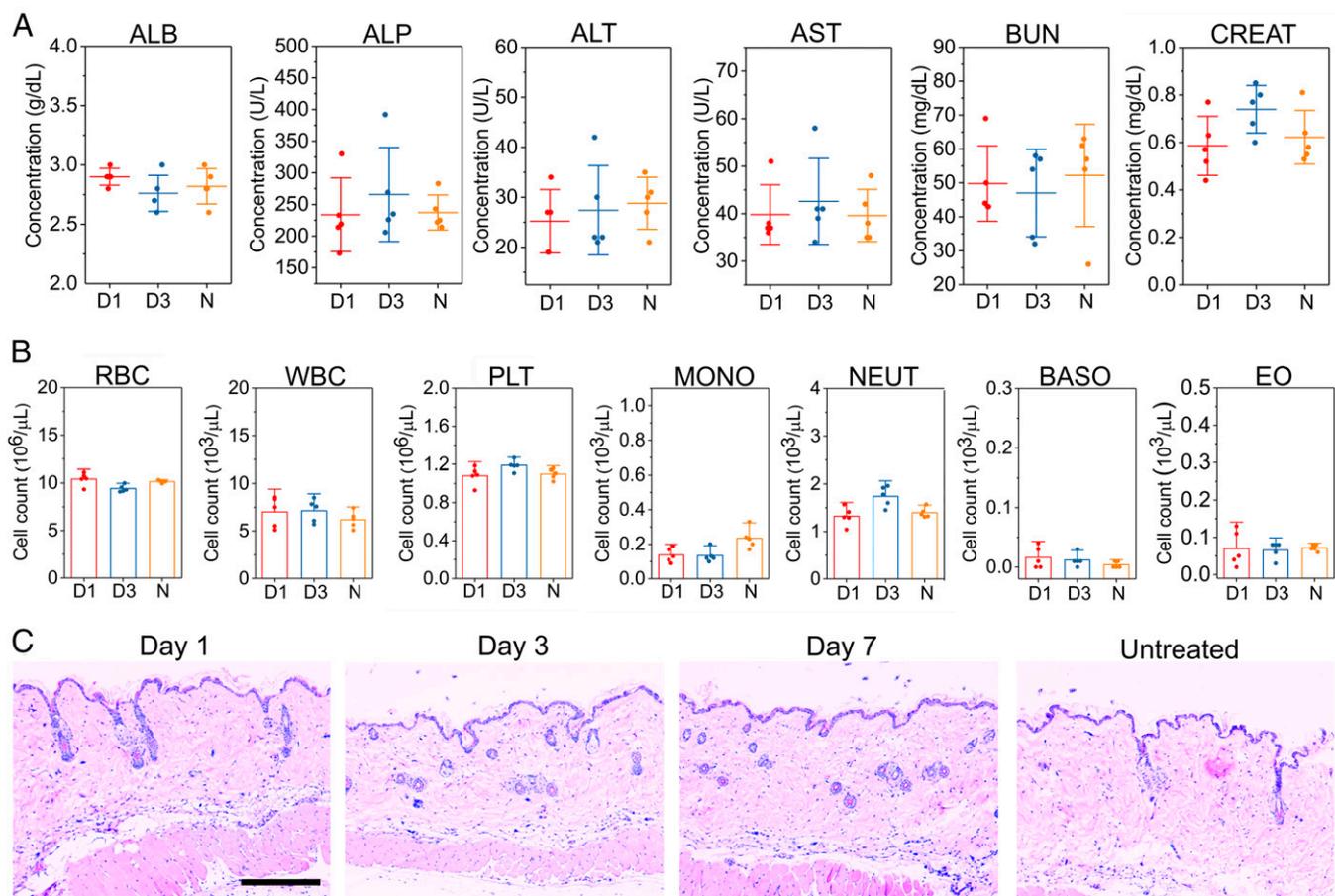


Fig. 4. Toxicity and biocompatibility evaluation of *i*-insulin in diabetic mice. (A) Serum ALB, ALP, ALT, AST, BUN, and CREAT concentration on day 1 or 3 posttreatment of *i*-insulin. Data are presented as mean ± SD ($n = 5$). (B) Red blood cell (RBC), white blood cell (WBC), platelet (PLT), monocyte (MONO), neutrophil (NEUT), basophil (BASO), and eosinophil (EO) were counted on day 1 or 3 posttreatment of *i*-insulin. D1, day 1 posttreatment; D3, day 3 posttreatment; N, no-treatment group. Data are presented as mean ± SD ($n = 5$). (C) H&E staining of the skin, where *i*-insulin was s.c. administered, from the diabetic mice. (Scale bar, 300 μ m.)

hypoglycemia (Fig. 3B). This hypoglycemia-mitigating effect of *i*-insulin was further evaluated on healthy mice. Following s.c. injection, native insulin induced severe hypoglycemia represented by blood glucose levels of ~60 mg/dL. In contrast, *i*-insulin-treated mice did not show glucose levels below 100 mg/dL (Fig. 3C).

An i.p. glucose tolerance test (IPGTT) was also performed at 3 h posttreatment. A spike in blood glucose levels was observed for all groups; however, only *i*-insulin-treated mice and healthy mice showed blood glucose levels back to normal range within 2 h (Fig. 3D). The enhanced ability of *i*-insulin in regulating blood glucose levels compared with native insulin was confirmed by the area under curve (AUC) analyses (Fig. 3E). Moreover, a peak of plasma insulin associated with IPGTT was observed without delay in the *i*-insulin-treated mice (Fig. 3F).

The toxicity of *i*-insulin to the liver and kidney was further evaluated via serum albumin (ALB), alkaline phosphatase (ALP), alanine aminotransferase (ALT), aspartate transaminase (AST), blood urea nitrogen (BUN), and creatinine (CREAT) levels, on days 1 and 3 after the s.c. injection of *i*-insulin (Fig. 4A). No significant differences across any biochemical measures were observed. In addition, there were no obvious changes in blood cell counts (Fig. 4B). Meanwhile, negligible neutrophil infiltration was observed at treated skin sites, as shown in the hematoxylin and eosin (H&E) stain results (Fig. 4C). Moreover, the total levels of IgM and IgG associated with both healthy mice and diabetic mice were insignificantly changed during the 2-wk treatment (SI Appendix, Fig.

S15) (36, 37). Systemic evaluation of the specific immune responses against *i*-insulin upon variation of treatment doses, frequencies, and routes are required for further translation (37, 38).

Discussion

Bioresponsive insulin-mediated treatment has the potential to revolutionize the current diabetes treatment. An insulin molecule with the properties of glucose responsiveness and hypoglycemia mitigation would offer a novel approach to regulate blood glucose levels with low risk for hypoglycemia. In this study, we engineered such a molecule via conjugation of insulin to the Glut competitive inhibitor Glut-i2 to allow *i*-insulin for reversible and glucose-responsive binding to endogenous Glut. In vitro, the *i*-insulin was able to rapidly bind to Glut on erythrocyte ghosts at low blood glucose concentrations, while releasing free and glucose-accessible Glut in response to hyperglycemia. Upon a glucose challenge, *i*-insulin was liberated from the *i*-insulin–Glut complex for subsequent binding to IR and rapid blood glucose clearance.

Upon s.c. injection in type 1 diabetic mice, *i*-insulin showed a significantly more durable normoglycemia effect with negligible hypoglycemia, even after a second injection. This result was further confirmed with studies showing that *i*-insulin only slightly lowered blood glucose of healthy mice, while native insulin induced severe hypoglycemia. Upon a glucose challenge, a portion of *i*-insulin was released from *i*-insulin–Glut complex directly to the interstitial fluid and plasma. Remarkably, the direct release

of *i*-insulin to the interstitial environment may help *i*-insulin rapidly reach IRs on target cells. In addition, the release of *i*-insulin from the *i*-insulin–Glut complex generates glucose-accessible free Glut to enhance the excess glucose clearance from blood.

This glucose transporter inhibitor-mediated insulin can be further optimized, regarding response kinetics, effective duration, and Glut specificity, through varying the component(s) of glucose transporter inhibitor, insulin, and spacer. Moreover, this glucose-responsive insulin can be further integrated with painless transdermal microneedle array patch to generate a new version of “smart insulin patch” (*SI Appendix*, Fig. S16) (39, 40) or oral delivery systems to form “smart insulin pills” (41).

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Materials and Methods

Experimental procedures for insulin analog synthesis and in vitro glucose-triggered insulin release, procedures for animal experiment, and additional control experiments are provided in *SI Appendix*. The animal study protocol was approved by the Institutional Animal Care and Use Committee at North Carolina State University and the University of California, Los Angeles.

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