Is Incretin-Based Therapy Ready for the Care of Hospitalized Patients With Type 2 Diabetes?
Insulin therapy has proven itself and is considered the mainstay of treatment

Guillermo Umpierrez, Emory University
Mary Korytkowski, University of Pittsburgh

Journal Title: Diabetes Care
Volume: Volume 36, Number 7
Publisher: American Diabetes Association | 2013-07, Pages 2112-2117
Type of Work: Article | Final Publisher PDF
Publisher DOI: 10.2337/dc12-2233
Permanent URL: http://pid.emory.edu/ark:/25593/gj8jm

Final published version: http://care.diabetesjournals.org/content/36/7/2112

Copyright information:
© 2013, American Diabetes Association
This is an Open Access article distributed under the terms of the Creative Commons Attribution 3.0 Unported License (http://creativecommons.org/licenses/by/3.0/), which permits distribution of derivative works, distribution, public display, and publicly performance, making multiple copies, provided the original work is properly cited. This license requires credit be given to copyright holder and/or author, copyright and license notices be kept intact.

Accessed January 7, 2018 8:22 PM EST
Is Incretin-Based Therapy Ready for the Care of Hospitalized Patients With Type 2 Diabetes?

Insulin therapy has proven itself and is considered the mainstay of treatment.

Significant data suggest that overt hyperglycemia, either observed with or without a prior diagnosis of diabetes, contributes to an increase in mortality and morbidity in hospitalized patients. In this regard, goal-directed insulin therapy has remained as the standard of care for achieving and maintaining glycemic control in hospitalized patients with critical and noncritical illness. As such, protocols to assist in management of hyperglycemia in the inpatient setting have become commonplace in hospital settings. Clearly, insulin is a known entity, has been in clinical use for almost a century, and is effective. However, there are limitations to its use. Based on the observed mechanisms of action and efficacy, there has been a great interest in using incretin-based therapy with glucagon-like peptide-1 (GLP-1) receptor agonists instead of, or complementary to, an insulin-based approach to improve glycemic control in hospitalized, severely ill diabetic patients. To provide an understanding of both sides of the argument, we provide a discussion of this topic as part of this two-part point-counterpoint narrative. In the point narrative preceding the counterpoint narrative below, Drs. Schwartz and DeFronzo provide an opinion that now is the time to consider GLP-1 receptor agonists as a logical consideration for inpatient glycemic control. In the counterpoint narrative provided below, Drs. Umpierrez and Korytkowski provide a defense of insulin in the inpatient setting as the unquestioned gold standard for glycemic management in hospitalized settings.

—William T. Cefalu, MD
Editor in Chief, Diabetes Care

Hypoglycemia is reported in one-third of general medicine and surgery patients with and without a known history of diabetes (1). The prevalence is even higher in intensive care unit (ICU) patients and following cardiac surgery, occurring in up to 80% of patients (2). While previously thought to be an epiphenomenon related to the acute underlying illness of the hospitalization itself, hypoglycemia is now recognized as a contributor to adverse outcomes in critical and noncritically ill patients, with higher mortality and disease-specific morbidity (3,4).

Protocols using insulin to maintain glycemia within a reasonable range reduces both mortality and morbidity (2,5,6). In critically ill patients in ICU settings, intravenously (IV) administered insulin is the preferred method of achieving recommended glycemic targets. The short half-life of IV insulin permits rapid dosing adjustments in response to alterations in insulin sensitivity observed during critical illness. For the majority of ICU patients, insulin infusion is started at a threshold of no higher than 10.0 mmol/L (180 mg/dL). Once IV insulin is started, glucose levels should be maintained between 6.1 and 10.0 mmol/L (110 and 180 mg/dL) (6). For patients in non-ICU settings, recent guidelines (6–8) recommend the use of subcutaneous (SC) insulin as the preferred therapy. Scheduled basal bolus SC insulin therapy consisting of long- or intermediate-acting preparations in combination with short- or rapid-acting analogs has been proven to be safe and effective for glycemic management and to reduce hospital complications including wound infections, pneumonia, bacteremia, and acute renal and respiratory failure when compared with the use of sliding-scale insulin alone in patients with type 2 diabetes (5).

Improved glycemic control with insulin therapy is associated with amelioration of the hormonal and proinflammatory aberrations associated with stress hyperglycemia (9,10). These include reductions in counterregulatory hormones and proinflammatory transcription factors, and potentially the formation of reactive oxygen species (9). Insulin therapy induces vasodilatation by stimulating nitric oxide release and inducing expression of endothelial nitric oxide synthase (11). Insulin-mediated inhibition of lipolysis reduces circulating free fatty acid levels with improved insulin sensitivity, while inhibition of platelet aggregation reduces thrombosis (9). The major concern raised with the use of insulin therapy in the hospital is hypoglycemia, which has been observed primarily in studies targeting near normal glucose ranges of 80–110 mg/dL (6,12). This concern has prompted a search for alternative methods of inpatient glycemic management that are not known to cause hypoglycemia.

Hospital use of native glucagon-like peptide-1 and glucagon-like peptide-1 receptor agonist in medical and surgical patients —In contrast to the solid data supporting the use of insulin in hospitalized patients, there are only a handful of studies investigating the use of native glucagon-like peptide-1 (GLP-1) or GLP-1 receptor agonist (GLP-1RA) infusions in the inpatient setting. Most of these studies are uncontrolled, open-label, and of short-duration with small numbers of subjects with or without a history of diabetes (13–20) (Table 1). The majority of these studies were performed with the primary objective of investigating the safety and efficacy of these agents, with the hypothesis that GLP-1 infusions will achieve the desired glycemic targets without the risk for hypoglycemia. To date, none of these benefits have been demonstrated, and in many GLP-1-treated patients, rescue therapy with insulin was required to achieve and maintain the desired glycemic targets with no difference in the frequency of hypoglycemia when compared with insulin therapy. In addition, there was added risk for gastrointestinal side effects (nausea, vomiting, constipation), which occurred in up to 66% of subjects (13,16,18).
Table 1—Hospital use of native GLP-1 in medical and surgical patients and GLP-1RAs in critical care

<table>
<thead>
<tr>
<th>Author</th>
<th>Design</th>
<th>Population, n</th>
<th>Intervention</th>
<th>Findings</th>
<th>Adverse events</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nikolaidis</td>
<td>Single-center, nonrandomized controlled pilot study</td>
<td>Acute MI and LV systolic dysfunction</td>
<td>72-h IV GLP-1 (1.5 pmol/kg/min) following successful angioplasty</td>
<td>GLP-1 improved LV function and global wall motion scores indices</td>
<td>GLP-1 group: nausea (n = 4), vomiting (n = 2), constipation (n = 2), reduced appetite (n = 3); Hypoglycemia: 2 events (52 and 58 mg/dL)</td>
</tr>
<tr>
<td>Meier 2004</td>
<td>Single-center, randomized, placebo-controlled trial</td>
<td>8 patients with T2D, 2–8 days postmajor surgery</td>
<td>8-h IV GLP-1 infusion (1.2 pmol/kg/min) vs. placebo</td>
<td>GLP-1 reduced BG levels, increased insulin, C-peptide (P &lt; 0.001), and suppressed glucagon (P = 0.041)</td>
<td>No recorded hypoglycemic events or other adverse reactions</td>
</tr>
<tr>
<td>Sokos 2006</td>
<td>Single-center, open-label, nonrandomized pilot study</td>
<td>CHF on stable medications</td>
<td>6-week SC GLP-1 infusion, started at 1.25 pmol/kg/min for 1 week, followed for 4 weeks at 2.5 pmol/kg/min</td>
<td>GLP-1 improved LVEF, 6-min walk test, and improvement in BG compared with control subjects</td>
<td>GLP-1 group: nausea and constipation in 5 patients and increase in HR (~5 bpm); Hypoglycemia: 9 episodes in 4 GLP-1 patients, 4 episodes in 2 control patients</td>
</tr>
<tr>
<td>Sokos 2007</td>
<td>Single-center, randomized, double-blind, placebo-controlled pilot study</td>
<td>Elective CABG, GLP-1, n = 10 (DM = 2)</td>
<td>48-h IV GLP-1 infusion started 12-h preoperative and continued for 48-h postoperative vs. standard IV insulin therapy</td>
<td>GLP-1 improved glycemic control pre- and intraoperatively, but no difference in postoperative period; No group differences in LVEF at baseline, 48 h, or at discharge</td>
<td>Rescue insulin therapy required in 5 GLP-1 subjects</td>
</tr>
<tr>
<td>Mussig 2008</td>
<td>Single-center, randomized, open-label, controlled trial</td>
<td>Insulin-naive patients with T2D after elective CABG, GLP1, n = 10 (DM = 10)</td>
<td>12-h IV GLP-1 infusion (3.6 pmol/kg/min) after transfer from the operating room to the ICU vs. standard IV insulin therapy</td>
<td>Glycemic control was comparable in the 2 groups</td>
<td>Rescue therapy with insulin required for several GLP-1 patients; No adverse events or hypoglycemia reported</td>
</tr>
</tbody>
</table>

Continued on p. 2114
### Table 1—Continued

#### Hospital use of native GLP-1 in medical and surgical patients

<table>
<thead>
<tr>
<th>Author</th>
<th>Design</th>
<th>Population, n</th>
<th>Intervention</th>
<th>Findings</th>
<th>Adverse events</th>
</tr>
</thead>
<tbody>
<tr>
<td>Halbirk</td>
<td>Single-center, double-blinded placebo-controlled crossover design</td>
<td>Ischemic heart failure in non-DM subjects</td>
<td>48-h IV GLP-1 (1.0 pmol/kg/min) or placebo in random orders 14 days apart</td>
<td>GLP-1 had no effect on LVEF, diastolic function, exercise capacity, or regional myocardial contractile function</td>
<td>Nausea and vomiting in half of GLP-1 group</td>
</tr>
<tr>
<td>2010 (18)</td>
<td></td>
<td>GLP-1, n = 10; Placebo, n = 10</td>
<td>Infusion rate reduced 0.7 pmol/kg/min because of high frequency of hypoglycemia</td>
<td>GLP-1 resulted in 9 episodes of hypoglycemia in 8 patients (nadir 40 mg/dL) vs. no hypoglycemia in placebo group</td>
<td></td>
</tr>
</tbody>
</table>

#### Hospital use of GLP-1RAs in critical care

<table>
<thead>
<tr>
<th>Author</th>
<th>Design</th>
<th>Population, n</th>
<th>Intervention</th>
<th>Findings</th>
<th>Adverse events</th>
</tr>
</thead>
<tbody>
<tr>
<td>Abuannadi</td>
<td>Single-center, nonrandomized open-label, pilot study</td>
<td>T2D in cardiac ICU</td>
<td>24–48-h IV exenatide infusion (0.025 μg/min)</td>
<td>No differences in mean steady-state BG between exenatide and IV insulin therapy</td>
<td>Nausea with exenatide; 6 patients (15%) excluded because of severe nausea; Hypoglycemia reported in 10% with exenatide and 15–20% during insulin infusion</td>
</tr>
<tr>
<td>2013 (20)</td>
<td></td>
<td>Exenatide, n = 40 (DM = 40); Historic control subjects treated with insulin infusion (n = 133)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

| Mecott    | Single-center, open-label, controlled study          | Severely burned pediatric patients in a burned unit | SC exenatide 5–10 μg every 12 h vs. IV or SC insulin therapy | No differences in mean steady-state BG or glycemic variability between exenatide and insulin; Lower insulin requirement in the exenatide group | Hypoglycemia 0.38 events/patient/month in each group                             |
| 2010 (19) |                                                       | SC exenatide, n = 6; insulin infusion (n = 18) | | | |

DM, diabetes; HR, heart rate; LV, left ventricular; LVEF, left ventricular ejection fraction; MI, myocardial infarction; T2D, type 2 diabetes.
Surgical patients

In one small study of 20 subjects (5 with type 2 diabetes) undergoing coronary artery bypass grafting, glycemic control and left ventricular function was compared between 10 subjects (2 with type 2 diabetes) randomized to GLP-1 infusions initiated 12-h preoperatively at a dose of 1.3 pmol/kg/min and continued for 48-h postoperatively, and 10 subjects (3 with type 2 diabetes) assigned to standard insulin therapy (16). Standard therapy was not clearly defined but included IV insulin infusion, which was given to 5 subjects in each group. GLP-1 resulted in better glycemic control in the pre- and perioperative periods compared with insulin therapy, but there were no differences in postoperative blood glucose (BG) levels between treatment groups. In addition, there were no differences in insulin levels, hemodynamic parameters, or the number of hypoglycemic events during the study period. Half of the GLP-1 subjects required exogenous insulin infusions to achieve the desired level of glycemic control during the postoperative period. In a different study, 20 insulin-naïve patients with type 2 diabetes were randomized to exenatide at doses of 5 or 10 μg every 12 h (19,20). One open-label study included 48-h infusions of GLP-1 or placebo in random order (18). GLP-1 infusions resulted in more hypoglycemia with 8 patients experiencing 9 episodes of hypoglycemia (glucose <3.5 mmol/L) compared with none with placebo. Five patients were unable to complete the study because of severe nausea and vomiting and were excluded from analysis. Contrary to the results in prior studies, there was no beneficial effect of GLP-1 on left ventricular ejection fraction, diastolic function, or myocardial contractile function.

There are only two small uncontrolled pilot studies investigating the use of GLP-1RA for glycemic management in critically ill patients (19,20). One open-label study compared the efficacy and safety of SC administration of exenatide at doses of 5–10 μg every 12 h (n = 6) with standard intensive insulin therapy (n = 18) in severely burned pediatric patients without diabetes. Similar levels of glycemic control were achieved in both groups (130 ± 28 vs. 138 ± 25 mg/dL) (19); however, the dose of administered insulin was significantly lower in the exenatide group (22 ± 14 vs. 76 ± 11 unit/patients/day; P = 0.01). Three patients receiving exenatide required rescue therapy with SC insulin to maintain glycemic control. The number of BG determinations was identical, as was the incidence hypoglycemia (0.38 events/patient/month). There were no reported gastrointestinal side effects.

In another pilot nonrandomized, uncontrolled, open-label study evaluating the safety and efficacy of IV exenatide in 40 cardiac ICU patients, 75% with type 2 diabetes (20), subjects received an initial 30-min bolus of 0.05 μg/min followed by 0.025 μg/min for 24–48 h. Exenatide infusions resulted in similar mean steady-state BG and hypoglycemic events when compared with historic control subjects treated with IV insulin infusions targeting BG 90–119 mg/dL (n = 94) or 100–140 mg/dL (n = 39). The mean steady-state BG in the group treated with exenatide (139 ± 41 mg/dL) was similar to that achieved with IV insulin therapy (115 ± 36 mg/dL and 147 ± 52 mg/dL, respectively). Hypoglycemia (BG <70 mg/dL) was reported in 10% of patients receiving exenatide compared with 21 and 15% in those treated with IV insulin (P = 0.27). A total of 8 patients (20%) experienced nausea because of exenatide, and 6 patients (15%) requested early termination because of severe nausea.

Safety concerns of GLP-1 and GLP-1RA therapies in the hospital setting—Treatment with GLP-1 and GLP-1RA is associated with a high incidence of gastrointestinal side effects, as was observed in the majority of reported studies (Table 1). Nausea and vomiting can be potentially dangerous in hospitalized patients with altered sensorium, who are maintained in a supine position, or who receive sedating medications, all of which increase the risk for aspiration pneumonia. In addition, the risk for pancreatitis, although rarely reported with the GLP-1 therapy, cautions against the use of these agents in patients with abdominal pain or postsurgical ileus.

The observed increase in heart rate of 2–5 bpm reported in clinical trials with GLP-1 was also reported in several of the inpatient studies (17,18). The mechanism underlying the increase in heart rate has not yet been clarified, but in at least in one report was attributed to possible undetected hypoglycemia (18). Although preliminary cardiovascular safety analyses of GLP-1RA demonstrate trends toward reduced cardiovascular events (21,22), long-term studies are needed to determine the clinical relevance of these chronotropic effects, particularly in critically ill patients.

Current practice guidelines recommend against inpatient use of oral antidiabetic drugs in part because of the absence of efficacy studies as well as safety concerns (6–8). A major limitation to inpatient use of oral antidiabetic agents relates to the delay in and unpredictable onset of action, which prevents rapid attainment of glycemic control or dose adjustments to meet the changing needs of the acutely ill patient. The low risk of hypoglycemia and good tolerability of dipeptidyl peptidase 4 (DPP-4) inhibitors however make them attractive considerations.
for use in hospitalized patients. At this time however no randomized clinical trial studies have reported on the use of these agents in the hospital setting. A concern with the use of DPP-4 inhibitors is the increase in frequency of upper respiratory infections observed in some preclinical trials (23). There are no reports of an increase in serious infections, but this observation raises concern for the use of these medications in inpatients with compromised immune systems, such as those undergoing solid organ transplantation.

Conclusions — Since approval of exenatide and sitagliptin in 2005–2006, the use of GLP-1RA and DPP-4 inhibitors has been widely incorporated into clinical practice and treatment guidelines from leading diabetes organizations for use in outpatient settings. The question is whether or not these agents are safe for use in the inpatient population. It is important to emphasize that these are new medications for which even the long-term safety of their outpatient use remains to be established. It is therefore premature to make recommendations for their routine use in the inpatient population.

Insulin therapy remains the standard of care for achieving and maintaining glycemic control in hospitalized patients with critical and noncritical illness. Insulin is a known entity that has now been in clinical use for almost a century. Hypoglycemia, the major complication of insulin therapy, is avoidable in the majority of cases with appropriate use of either intravenous or basal-bolus insulin regimens with regular monitoring of bedside BG levels and modification of insulin dosage in response to changes in clinical and nutritional status (1,7,8).

The proposed reasons for using GLP-1 therapy for the management of hyperglycemia in hospitalized patients include a theoretical improvement in glucose control with low frequency of hypoglycemia, and less nursing time to monitor BG levels and adjust insulin doses. In the small numbers of patients studied to date, glycemic control with GLP-1 therapies has been shown to be superior to placebo, but not to insulin therapy (Table 1). In fact, almost 50% of subjects in some studies required rescue therapy with insulin in order to achieve and maintain glycemic control. The frequency of hypoglycemia was not reduced with GLP-1 therapies compared with insulin. In addition, in the studies investigating the frequency of BG monitoring and nurse time, no differences were found between insulin and GLP-1 therapies.

Novel drugs for the management of patients with diabetes and hyperglycemia are welcome when they are proven to be safe in improving glycemic control and in reducing cardiovascular and other complications. The preliminary experience with native GLP-1 is promising and has the potential to improve cardiovascular function in patients with heart failure and acute ischemic cardiovascular events (15–17). However, this requires further study in a larger number of patients. It is possible that these favorable results may extend to the use of GLP-1RA and DPP-4 inhibitors; however, until the safety and efficacy are addressed with large randomized controlled clinical trials, the routine use of these agents for inpatient glycemic control cannot be recommended.

History has taught us that despite the potential promising effects of some drugs, long-term and widespread use unmask undesirable and in some cases life-threatening side effects resulting in the removal of these agents from clinical use. Recent examples from diabetes management occurred following the introduction of thiazolidinediones, a class of medications associated with effective glucose-lowering properties as well as potential cardiovascular benefits. However, troglitazone was removed from the market following reports of sometimes fatal idioopathic hepatic failure (24); and the use of rosiglitazone became highly restricted following reports of higher mortality because of ischemic cardiovascular events (25). More recent observational and clinical studies have shown an association between the use of thiazolidinediones and increased bone fractures (26). Until there are well-conducted published studies demonstrating both the efficacy and safety of using the potentially promising GLP-1RA or DPP-4 inhibitors in the inpatient setting, it is best to observe the sound principles of evidence-based medicine and adhere to the well-known saying from the 12th century: All that glitters is not gold.

Guillermo E. Umpierrez, MD
Mary Korytkowski, MD

From the 1Department of Medicine, Division of Endocrinology, Emory University, Atlanta, Georgia; and the 2Department of Medicine, Division of Endocrinology, University of Pittsburgh, Pittsburgh, Pennsylvania.

Corresponding author: Guillermo E. Umpierrez, gUMPie@emory.edu.
11. Aljada A, Ghanim H, Mohanty P, Kapur N, Dandona P. Insulin inhibits the proinflammatory transcription factor early growth response gene-1 (Egr)-1 expression in mononuclear cells (MNC) and reduces plasma tissue factor (TF) and plasminogen activator inhibitor-1 (PAI-1) concentrations. J Clin Endocrinol Metab 2002;87:1419–1422