



Advancing Therapy in Suboptimally Controlled Basal Insulin–Treated Type 2 Diabetes: Clinical Outcomes With iGlarLixi Versus Premix BIAsp 30 in the SoliMix Randomized Controlled Trial

Diabetes Care 2021;44:2361–2370 | <https://doi.org/10.2337/dc21-0393>

Julio Rosenstock,¹ Rifat Emral,² Leobardo Sauque-Reyna,³ Viswanathan Mohan,⁴ Carlos Trescolí,⁵ Saud Al Sifri,⁶ Nebojsa Lalic,⁷ Agustina Alvarez,⁸ Pascaline Picard,⁹ Mireille Bonnemaire,¹⁰ Nacima Demil,¹¹ and Rory J. McCrimmon,¹² on behalf of the SoliMix Trial Investigators

OBJECTIVE

To directly compare the efficacy and safety of a fixed-ratio combination, of insulin glargine 100 units/mL and the glucagon-like peptide 1 receptor agonist lixisenatide (iGlarLixi), with those of a premix insulin analog, biphasic aspart insulin 30 (30% insulin aspart and 70% insulin aspart protamine) (BIAsp 30) as treatment advancement in type 2 diabetes suboptimally controlled on basal insulin plus oral antihyperglycemic drugs (OADs).

RESEARCH DESIGN AND METHODS

In SoliMix, a 26-week, open-label, multicenter study, adults with suboptimally controlled basal insulin–treated type 2 diabetes ($HbA_{1c} \geq 7.5\%$ and $\leq 10\%$) were randomized to once-daily iGlarLixi or twice-daily BIAsp 30. Primary efficacy end points were noninferiority in HbA_{1c} reduction (margin 0.3%) or superiority in body weight change for iGlarLixi versus BIAsp 30.

RESULTS

Both primary efficacy end points were met: after 26 weeks, baseline HbA_{1c} (8.6%) was reduced by 1.3% with iGlarLixi and 1.1% with BIAsp 30, meeting noninferiority (least squares [LS] mean difference -0.2% [97.5% CI -0.4 , -0.1]; $P < 0.001$). iGlarLixi was also superior to BIAsp 30 for body weight change (LS mean difference -1.9 kg [95% CI -2.3 , -1.4]) and percentage of participants achieving $HbA_{1c} < 7\%$ without weight gain and $HbA_{1c} < 7\%$ without weight gain and without hypoglycemia (all $P < 0.001$). iGlarLixi was also superior versus BIAsp 30 for HbA_{1c} reduction ($P < 0.001$). Incidence and rates of American Diabetes Association level 1 and 2 hypoglycemia were lower with iGlarLixi versus BIAsp 30.

CONCLUSIONS

Once-daily iGlarLixi provided better glycemic control with weight benefit and less hypoglycemia than twice-daily premix BIAsp 30. iGlarLixi is a more efficacious, simpler, and well-tolerated alternative to premix BIAsp 30 in suboptimally controlled type 2 diabetes requiring treatment beyond basal insulin plus OAD therapy.

¹Dallas Diabetes Research Center at Medical City, Dallas, TX

²Department of Endocrinology and Metabolic Diseases, Ankara University Faculty of Medicine, Ankara, Turkey

³Instituto de Diabetes Obesidad y Nutrición S.C., Cuernavaca, Morelos, Mexico

⁴Dr. Mohan's Diabetes Specialities Centre & Madras Diabetes Research Foundation, IDF Centre of Excellence in Diabetes Care & ICMR Centre for Advanced Research on Diabetes, Chennai, India

⁵Hospital Universitario de La Ribera, Alzira, Spain

⁶Al Hada Military Hospital, Taif, Saudi Arabia

⁷Faculty of Medicine of the University of Belgrade, Clinic for Endocrinology, Diabetes and Metabolic Diseases, Clinical Center of Serbia, Belgrade, Serbia

⁸Sanofi, Buenos Aires, Argentina

⁹IviData LIFE SCIENCES, Levallois-Perret, France

¹⁰Sanofi, Paris, France

¹¹Diabetes Medical Operation Department, Sanofi, Chilly-Mazarin, France

¹²Division of Systems Medicine, School of Medicine, University of Dundee, Dundee, U.K.

Corresponding author: Julio Rosenstock, juliorosenstock@dallasdiabetes.com

Received 16 February 2021 and accepted 8 June 2021

Clinical trial reg. no. 2017-003370-13, <https://www.clinicaltrialsregister.eu/>

This article contains supplementary material online at <https://doi.org/10.2337/figshare.14754426>.

A brief summary of the SoliMix trial results can be found in the accompanying video abstract and infographic available with the online, full-text version of the article at <https://doi.org/10.2337/dc21-0393>.

© 2021 by the American Diabetes Association. Readers may use this article as long as the work is properly cited, the use is educational and not for profit, and the work is not altered. More information is available at <https://www.diabetesjournals.org/content/license>.

Clinical guidelines recommend a target HbA_{1c} of <7.0% (<53 mmol/mol) for most nonpregnant adults with type 2 diabetes (1,2), while recognizing the need to individualize glycemic targets based on patient preference, and treatment efficacy and safety profiles (1,2). Most current guidelines advocate a stepwise introduction of pharmacotherapy for people with type 2 diabetes not achieving their individualized glycemic targets. With this approach, advancing basal insulin therapy involves four options: 1) adding rapid-acting insulin progressively to an existing basal insulin regimen, 2) multiple daily premix insulin doses (basal and prandial insulin coformulation), 3) adding a daily or weekly glucagon-like peptide 1 receptor agonist (GLP-1 RA) to an existing basal insulin regimen, and 4) switching to a once-daily fixed-ratio combination (FRC) of basal insulin and GLP-1 RA (2,3).

Each aforementioned treatment option has been shown to improve glycemic control when used to advance therapy from basal insulin but is also associated with specific adverse effects (4–7). GLP-1 RA therapy can be associated with gastrointestinal (GI) adverse events (AEs) and resultant adherence issues (4). Basal plus rapid-acting insulin regimens and premix insulin regimens can increase the risks of hypoglycemia and weight gain, while requiring multiple daily injections and frequent glucose monitoring that increase treatment burden and may reduce adherence (5–7). Despite this, premix insulins are widely used globally, particularly in Asia, Africa, the Middle East, China, and some European Union countries (8–11).

Titratable FRCs of basal insulin and a GLP-1 RA can provide a novel alternative therapy advancement option to premix insulin, as tested for the first time in this randomized controlled trial (RCT). FRCs combine the complementary mechanisms of action of two individual components in one formulation; basal insulin primarily reduces fasting plasma glucose (FPG) while the GLP-1 RA targets postprandial glucose (PPG). Overall, GLP-1 RAs act through a glucose-dependent mechanism by which they stimulate insulin secretion while preventing glucagon increase (12). Short-acting GLP-1 RAs specifically target PPG with a predominant gastric emptying effect, while long-acting GLP-1 RAs exert their effect predominantly through

pancreatic functions, resulting in a lesser impact on PPG but larger overall reductions in FPG (12). Two once-daily titratable FRCs of basal insulin and GLP-1 RA are available that are approved for use in adults with type 2 diabetes. iGlarLixi is an FRC of basal insulin glargine 100 units/mL (iGlar) and the short-acting GLP-1 RA lixisenatide (Lixi) (13,14), while IDegLira is an FRC of the basal insulin degludec and the long-acting GLP-1 RA liraglutide (15,16). Both have been shown to provide improved glycemic control versus their individual components, along with weight benefits compared with basal insulin and fewer GI AEs compared with their GLP-1 RA component (17–22).

Here we report the results of the first randomized, head-to-head study directly comparing the efficacy and safety of an FRC (iGlarLixi) with a premix insulin (biphasic insulin aspart 30 [BIAsp 30]) in adults with type 2 diabetes advancing from basal insulin plus one or two oral antihyperglycemic drugs (OADs).

RESEARCH DESIGN AND METHODS

Detailed methods have previously been published (23). In brief, SoliMix was an open-label, multicenter, randomized, 26-week study undertaken to compare the efficacy and safety of iGlarLixi with BIAsp 30 in adults with suboptimally controlled type 2 diabetes (HbA_{1c} ≥7.5% [≥58.5 mmol/mol] and ≤10% [≤85.8 mmol/mol]) despite receiving stable doses of basal insulin plus OADs (metformin ± sodium–glucose cotransporter 2 [SGLT2] inhibitor) for 3 months. Exclusion criteria included individuals having type 1 diabetes, BMI of <20 and ≥40 kg/m², and basal insulin dose of <20 or >50 units at screening and use of any antihyperglycemic agent other than basal insulin, metformin, or SGLT2 inhibitors in the 3 months prior to screening.

Participants were randomized (1:1) to receive once-daily subcutaneous iGlarLixi (Suliqua, [SOLIUQUA]; Sanofi, Paris, France) or twice-daily subcutaneous BIAsp 30 (30% insulin aspart + 70% insulin aspart protamine, NovoMix 30; Novo Nordisk A/S, Bagsværd, Denmark). iGlarLixi was injected before a meal with a prefilled disposable SoloStar pen injector. BIAsp 30 was administered subcutaneously twice daily in the morning and before dinner. Participants were switched from

their prior basal insulins at randomization; OADs were continued without adjustment. Starting doses of iGlarLixi were based on prior basal insulin doses, according to labeling instructions. If the previous basal insulin dose at randomization was <30 units, the starting dose was 20 dose steps (20 units iGlar, 10 μg Lixi) administered with the 10–40 units pen (2 units:1 μg ratio); if basal insulin was ≥30 to ≤50 units, the starting dose was 30 dose steps (30 units iGlar, 10 μg Lixi) administered with the 30–60 units pen (3 units:1 μg ratio). Further details are shown in Supplementary Table 1. Starting total daily doses of BIAsp 30 were the same as the participants' previous basal insulin dose on a unit-to-unit basis and split into two daily doses. Doses of iGlarLixi and BIAsp 30 were recommended for weekly titration based on fasting or premeal self-measured plasma glucose, respectively, to a target of 80–110 mg/dL (4.4–6.1 mmol/L). The recommended dose adjustment algorithms for iGlarLixi and BIAsp 30 were indicated according to label recommendations and are shown in Supplementary Tables 2 and 3.

Rescue therapy use was recommended according to the investigator's clinical judgment for both arms to correct hyperglycemia persisting beyond prespecified thresholds (HbA_{1c} >8% or FPG >200 mg/dL from week 12). Rescue therapy was to be considered in the iGlarLixi group when the maximal dose of 60 dose steps was reached. The use of any additional antihyperglycemic treatment (basal insulin, rapid-acting insulin, third daily injection of premix, or OADs) administered to participants in either group with the objective of rescue was included in the analysis of the proportion of participants requiring rescue therapy.

This study is registered on the European Union Drug Regulating Authorities Clinical Trials Database (2017-003370-13) and was conducted in accordance with the ethics principles of the Declaration of Helsinki, the International Conference on Harmonization guidelines for good clinical practice, and all applicable laws, rules, and regulations.

Study End Points

The two primary objectives of this study were to demonstrate that, compared with BIAsp 30, iGlarLixi was noninferior

in terms of HbA_{1c} reduction or superior in terms of body weight change from baseline to week 26. Key secondary efficacy end points were assessed at week 26, including HbA_{1c} <7% without weight gain at week 26, HbA_{1c} <7% without weight gain at week 26 and without hypoglycemia (plasma glucose <70 mg/dL [<3.9 mmol/L]) during the treatment period, and the superiority of iGlarLixi versus BIAsp 30 in terms of HbA_{1c} reduction from baseline to week 26. Other secondary exploratory glycaemic end points included the proportion of patients reaching HbA_{1c} target <7% at week 26, HbA_{1c} target <7% without American Diabetes Association (ADA) level 2 hypoglycemia, HbA_{1c} <7% without weight gain of >1 kg, and HbA_{1c} <6.5%. Other secondary end points included change in total insulin dose and change in FPG, from baseline to week 26.

Safety end points were hypoglycemia, AEs, serious AEs (SAEs), AEs leading to treatment discontinuation, and AEs leading to death. Hypoglycemia was defined as current ADA level 1 (<70 mg/dL [<3.9 mmol/L] and ≥ 54 mg/dL [≥ 3.0 mmol/L]), level 2 (<54 mg/dL [<3.0 mmol/L]), or level 3 (severe hypoglycemia). Nocturnal hypoglycemia was also assessed post hoc using two definitions: between bedtime and waking, and between 0000 and 0600 h.

Statistical Analysis

A sample size of 864 randomized participants (432 randomized or 388 evaluable participants per treatment group) was calculated based on the primary efficacy variables of HbA_{1c} and weight change from baseline to week 26. Assuming a dropout rate of 10%, this sample size provides >95% power to demonstrate noninferiority (margin 0.3%) of iGlarLixi versus BIAsp 30 for HbA_{1c} reduction or superiority for weight reduction at week 26. The assumptions made for noninferiority of HbA_{1c} were an SD of 1.1%, a noninferiority margin of 0.3%, and a zero true difference in HbA_{1c} between treatment groups. Assumptions for superiority testing of iGlarLixi over BIAsp 30 in terms of weight gain included an expected difference of 1 kg between treatment groups and an SD of 3.46 kg for changes from baseline. The two-sided significance level of 0.025 was assumed for each of the above tests.

The primary efficacy end points were analyzed using a multiple imputation strategy and an ANCOVA model including screening HbA_{1c} value (<8.0% vs. $\geq 8.0%$, for the change in body weight primary end point only), basal insulin dose (<30 units, ≥ 30 units) and SGLT2 inhibitor use (yes, no), treatment group, and country as fixed categorical effects and fixed continuous covariates of baseline values for each primary end point (HbA_{1c} and body weight).

Continuous secondary efficacy end points (e.g., FPG and total daily insulin dose) were analyzed using the same approach as the primary end points including the baseline values for the end point in question as fixed covariates. Categorical secondary efficacy end points (e.g., the first two key secondary end points) were analyzed using a logistic regression model adjusting for treatment group randomization strata, and HbA_{1c} and weight baseline covariates.

A multiple testing procedure was prespecified for analysis of the primary and key secondary efficacy end points (Supplementary Fig. 1). Following the two primary end points, the three key secondary end points were assessed using a hierarchical order: superiority of iGlarLixi versus BIAsp 30 in achieving HbA_{1c} <7% without weight gain, then superiority of HbA_{1c} <7% without weight gain and without hypoglycemia, and then superiority of HbA_{1c} reduction. More detailed information pertaining to control for type I error has previously been published (23).

All efficacy analyses were performed on data from the intention-to-treat (ITT) population, defined as all randomized participants. The coronavirus disease 2019 (COVID-19) pandemic occurred during the last few weeks of the study in some countries, making it difficult for some participants to comply with the protocol. For assessment of the potential impact of this on the primary and key secondary efficacy end points, sensitivity analyses were performed on a subgroup of the ITT population who had no major or critical deviations related to the COVID-19 pandemic situation that could have affected the primary efficacy analysis. Participants who followed the study visits and assessments without being impacted by the COVID-19 pandemic and its consequences (e.g., lockdown, sites closed, postponed/incomplete

end-of-treatment visit) were defined as the “nonimpacted by COVID-19 population.” Further sensitivity analyses were performed for the noninferiority objective in the “per protocol population,” defined as all participants in the ITT population who completed 26 weeks of randomized treatment without any major protocol violations.

Safety analyses were based on data from the safety population, defined as all randomized participants who received at least one dose of study drug.

RESULTS

Participant Disposition and Baseline Characteristics

In total, 887 participants from 89 centers in 17 countries were randomized in the study, of whom 443 were allocated to iGlarLixi and 444 to BIAsp 30. Of the 887 participants in the ITT population, 403 in the iGlarLixi group and 404 in the BIAsp 30 group were included in the nonimpacted by COVID-19 population. No participants discontinued due to COVID-19. Overall, participants received treatment with iGlarLixi or BIAsp 30 for a mean duration of 184 or 181 days, respectively. In total, 844 (95.2%) participants completed the 26-week treatment period: 428 (96.6%) in the iGlarLixi group and 416 (93.7%) in the BIAsp 30 arm (Supplementary Fig. 2).

Demographics and baseline characteristics were similar across both treatment groups (Table 1) and have previously been reported (23). Briefly, the randomized population was primarily white (62.4%) with a mean \pm SD age of 59.8 \pm 10.2 years, BMI of 29.9 \pm 4.9 kg/m², and duration of type 2 diabetes of 13.0 \pm 7.2 years. Metformin was used at baseline in 99.8% of all participants: approximately one-quarter were also receiving SGLT2 inhibitors at baseline in both treatment groups. Basal insulins used at randomization were insulin glargine 100 units/mL (46%), insulin glargine 300 units/mL (22%), NPH insulin (21%), insulin detemir (7%), and insulin degludec (5%).

Efficacy End Points

The two primary efficacy end points and all three key secondary efficacy end points were met. Mean \pm SD baseline HbA_{1c} was 8.6 \pm 0.7% (71 \pm 7 mmol/mol) in the iGlarLixi group and 8.6 \pm 0.7% (70 \pm 7 mmol/mol) in the BIAsp 30

Table 1—Abbreviated baseline characteristics (randomized population)

Demographic/clinical characteristic	iGlarLixi (n = 443)	BIAsp 30 (n = 444)	All participants (N = 887)
Age (years)			
Mean ± SD	59.8 ± 10.3	59.8 ± 10.0	59.8 ± 10.2
Median	61	60	61
Q1, Q3	52, 67	54, 67	53, 67
Sex, n (%)			
Male	224 (50.6)	218 (49.1)	442 (49.8)
Female	219 (49.4)	226 (50.9)	445 (50.2)
BMI (kg/m²)			
Mean ± SD	29.7 ± 4.7	30.0 ± 5.1	29.9 ± 4.9
Median	29.1	29.2	29.1
Q1, Q3	26.2, 32.9	26.2, 34.2	26.2, 33.6
Duration of type 2 diabetes (years)			
Mean ± SD	13.0 ± 7.1	13.0 ± 7.4	13.0 ± 7.2
Median	12.0	12.0	12.0
Q1, Q3	7.6, 17.0	7.2, 17.0	7.5, 17.0
Prior basal insulin at baseline, n (%)*			
Insulin glargine 100 units/mL	188 (42.4)	219 (49.2)	407 (45.8)
Insulin glargine 300 units/mL	100 (22.6)	92 (20.7)	192 (21.6)
NPH	102 (23.0)	82 (18.4)	184 (20.7)
Insulin detemir	34 (7.7)	31 (7.0)	65 (7.3)
Insulin degludec	19 (4.3)	21 (4.7)	40 (4.5)
Average basal insulin daily dose (units)†			
Mean ± SD	33.8 ± 9.6	33.8 ± 9.9	33.8 ± 9.8
Median	34.0	34.0	34.0
Q1, Q3	25.0, 40.0	24.0, 42.0	25.0, 40.0
Average basal insulin daily dose (units/kg)†			
Mean ± SD	0.43 ± 0.15	0.43 ± 0.14	0.43 ± 0.14
Median	0.42	0.42	0.42
Q1, Q3	0.32, 0.52	0.32, 0.51	0.32, 0.52
Previous noninsulin antihyperglycemic treatment, n (%)*			
Metformin	443 (100.0)	442 (99.5)	885 (99.8)
SGLT2 inhibitor	104 (23.5)	102 (23.0)	206 (23.2)
Other	1 (0.2)	2 (0.5)	3 (0.3)
Daily metformin dose at baseline (mg)			
Mean ± SD	1,761 ± 542	1,722 ± 549	1,741 ± 546
Median	2,000	1,850	2,000
Q1, Q3	1,500, 2,000	1,500, 2,000	1,500, 2,000
Diabetes-related complications, n (%)			
Diabetic neuropathy	119 (26.9)	127 (28.6)	246 (27.7)
Diabetic retinopathy (incl. proliferative diabetic retinopathy)	67 (15.1)	67 (15.1)	134 (15.1)
Diabetic nephropathy	45 (10.2)	41 (9.2)	86 (9.7)
Heart failure	11 (2.5)	8 (1.8)	19 (2.1)
Peripheral artery disease	2 (0.5)	9 (2.0)	11 (1.2)
Ischemic stroke	2 (0.5)	0	2 (0.2)

Full baseline characteristics have previously been reported (23). incl., including; Q, quartile; SGLT2, sodium–glucose cotransporter 2. *A participant can be counted in more than one category. †Within the 3 days immediately before randomization.

group. At week 26, mean ± SD HbA_{1c} had improved to 7.3 ± 1.1% (56 ± 12 mmol/mol) in the iGlarLixi group and to 7.5 ± 1.0% (58 ± 11 mmol/mol) in the BIAsp 30 group (Fig. 1A and B). Statistical noninferiority (margin 0.3%) of iGlarLixi over BIAsp 30 was demonstrated for the change in HbA_{1c} from baseline to week 26 (LS mean difference vs. BIAsp 30, −0.2% [97.5% CI −0.4,

−0.1]; −2.6 mmol/mol [−4.5, −0.9]; *P* < 0.001). Additionally, statistical superiority in HbA_{1c} reduction from baseline to week 26 of iGlarLixi over BIAsp 30 was demonstrated as part of the key secondary end point analysis, on the basis of the hierarchical testing procedure (Fig. 1B).

At baseline, mean ± SD body weight was 80.7 ± 16.5 kg in the iGlarLixi group and 82.2 ± 18.5 kg in the BIAsp 30 group.

From baseline to week 26, mean ± SD body weight decreased to 80.2 ± 16.6 kg for iGlarLixi and increased to 83.4 ± 19.0 kg for BIAsp 30 (Fig. 1C). Statistical superiority of iGlarLixi over BIAsp 30 was demonstrated for the change in body weight from baseline to week 26 (LS mean difference vs. BIAsp 30, −1.9 kg [95% CI −2.3, −1.4]; *P* < 0.001).

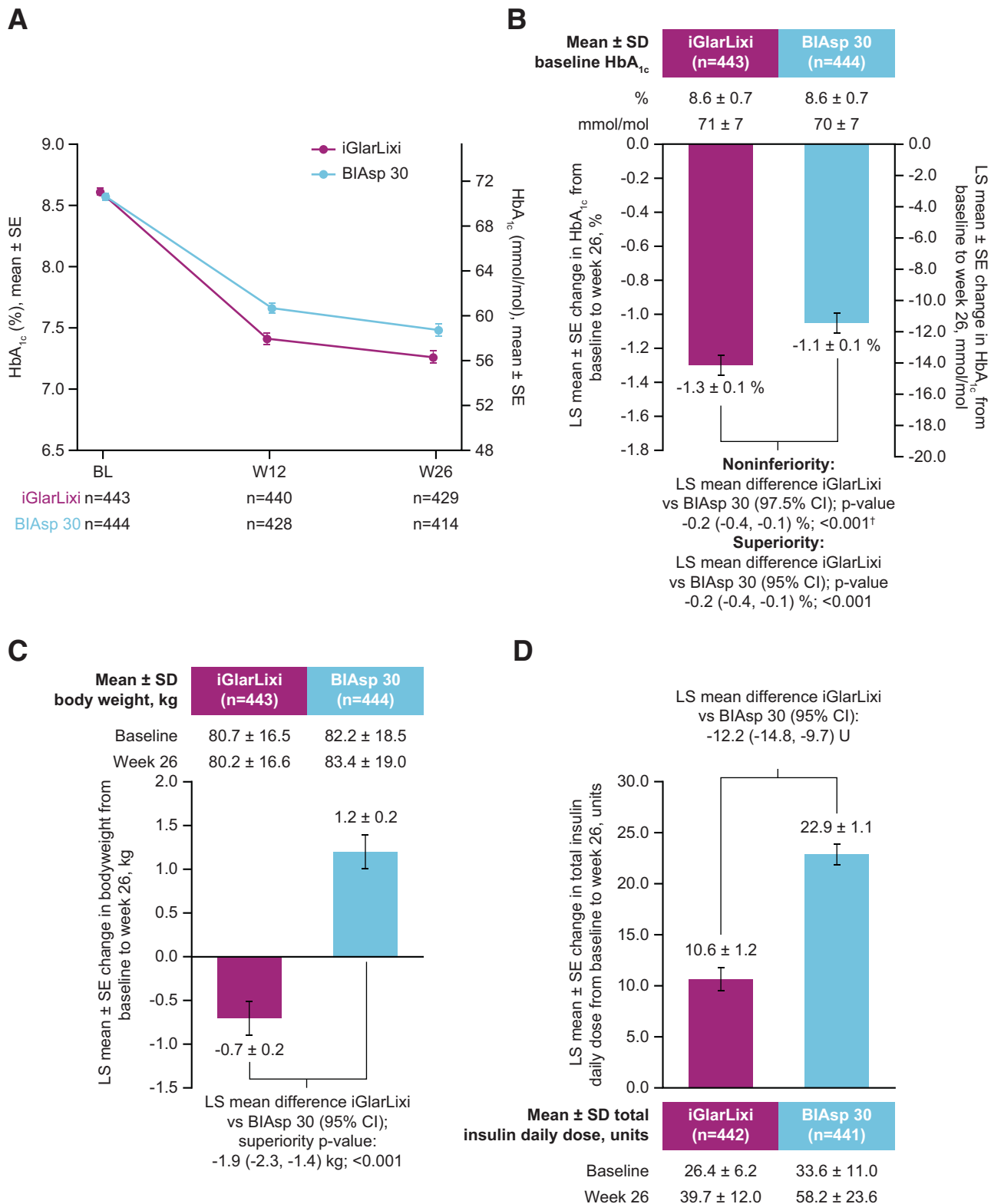


Figure 1—HbA_{1c} over 26 weeks (A) and change in HbA_{1c} (B), body weight (C), and total insulin daily dose (D) from baseline to week 26 (ITT population). Missing data in the primary HbA_{1c} and weight end points were imputed through a multiple imputation strategy under the missing not at random framework, with separate multiple imputation process function of treatment completeness. Missing values were imputed 1,000 times. †Noninferiority P value was calculated using a noninferiority margin of 0.3%. BL, baseline; W, week.

Key secondary efficacy end points showed that, compared with the BIAsp 30 group, a significantly greater proportion of participants in the iGlarLixi group reached HbA_{1c} <7% (<53 mmol/mol) without weight gain at week 26, and without weight gain at week 26 and without hypoglycemia (<70 mg/dL [<3.9 mmol/L]) during the treatment period (Fig. 2). The percentage of participants who reached HbA_{1c} target <7% was higher in the iGlarLixi group than in the BIAsp 30 group (Fig. 2) (exploratory end point). iGlarLixi was also associated with higher proportions of HbA_{1c} <7% target achievement without ADA level 2 hypoglycemia, HbA_{1c} <7% without weight gain of >1 kg, and HbA_{1c} <6.5% than BIAsp 30 (Fig. 2) (exploratory end points).

Mean \pm SD FPG at baseline was 151 \pm 44 mg/dL (8.4 \pm 2.4 mmol/L) in

the iGlarLixi group and 149 \pm 41 mg/dL (8.3 \pm 2.3 mmol/L) in the BIAsp 30 group. At week 26, mean \pm SD FPG was 130 \pm 44 mg/dL (7.2 \pm 2.4 mmol/L) in the iGlarLixi group and 146 \pm 51 mg/dL (8.1 \pm 2.8 mmol/L) in the BIAsp 30 group. The LS mean difference between groups in change from baseline to week 26 was -16 mg/dL (95% CI $-26, -6$) (-0.9 mmol/L [$-1.5, -0.3$]).

After 26 weeks, the increase in LS mean total daily insulin dose was smaller in the iGlarLixi group than in the BIAsp 30 (Fig. 1D). The percentage of participants who required rescue therapy was low and similar for iGlarLixi (1.8%) and BIAsp 30 (2.3%).

Detailed data for efficacy end points can be found in Supplementary Table 4. All key sensitivity analyses performed

on the two primary and key secondary end points demonstrated results similar to those observed in the ITT population (Supplementary Table 5).

Safety Profile

The proportion of participants with at least one hypoglycemic event was lower in the iGlarLixi group compared with the BIAsp 30 group (odds ratio [OR] 0.62 [95% CI 0.47, 0.81]) (Fig. 3). Lower incidence of hypoglycemia with iGlarLixi versus BIAsp 30 was also observed across level 1 and level 2 hypoglycemia categories (Fig. 3).

Rates of hypoglycemia followed the same pattern as incidence. There was an overall lower rate of any hypoglycemia with iGlarLixi compared with BIAsp 30, as well as lower rates of level 1 and level 2 hypoglycemia (Fig. 3).

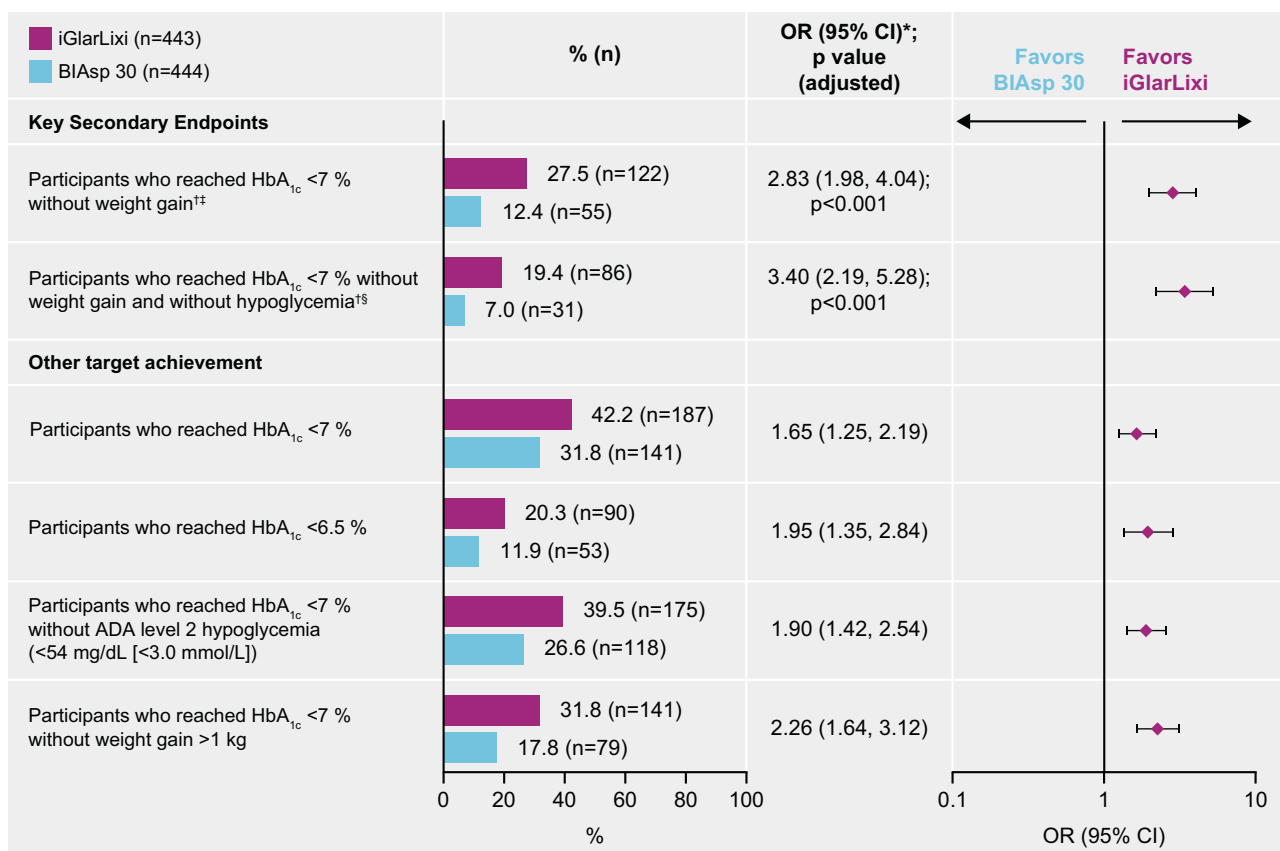
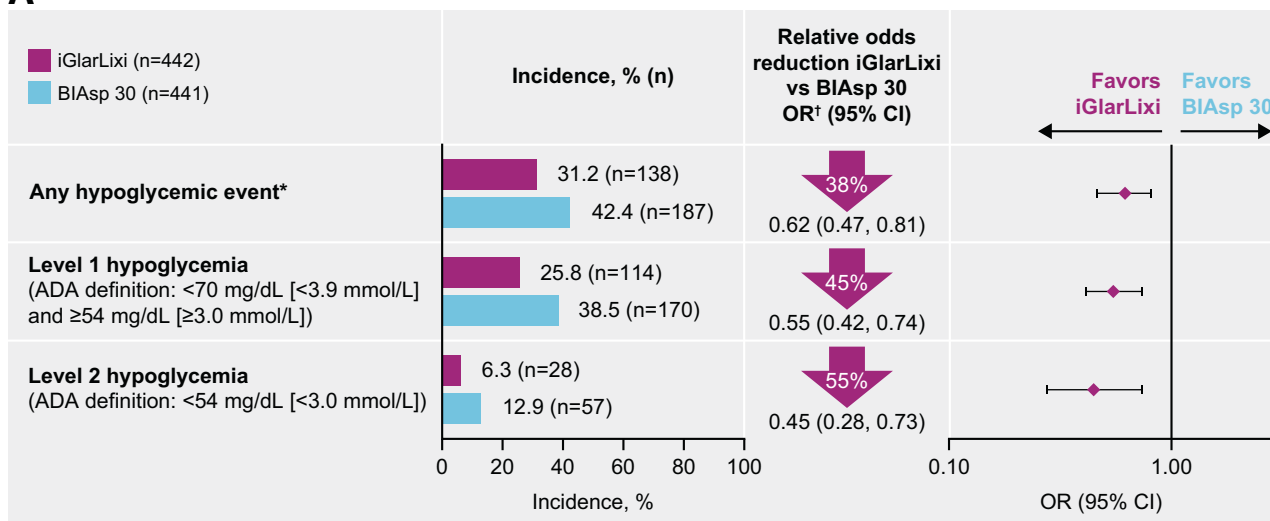


Figure 2—Glycemic target achievement and composite secondary efficacy end points (ITT population). *Adjusted OR of iGlarLixi vs. BIAsp 30 with associated two-sided CI (at the specified significance level that is passed from family 1 of the primary objectives), calculated by logistic regression model adjusted for fixed categorical effects of randomization strata (basal insulin dose at screening <30 units and \geq 30 units and SGLT2 inhibitor use [yes, no] at screening) and treatment group as well as fixed continuous covariates of baseline values for each of the primary end points (HbA_{1c} and body weight). †Imputed as not having reached HbA_{1c} target (failure, i.e., nonresponder) in the case of missing HbA_{1c} or weight values at week 26. ‡Weight gain defined as any increase >0 kg from baseline. §Hypoglycemia defined as plasma glucose <70 mg/dL (<3.9 mmol/L) occurring at any point within the 26-week open-label randomized treatment period. Assessments were done in hierarchical order, starting with the proportion of participants who reached HbA_{1c} <7% without weight gain. n, number of participants.

A



B

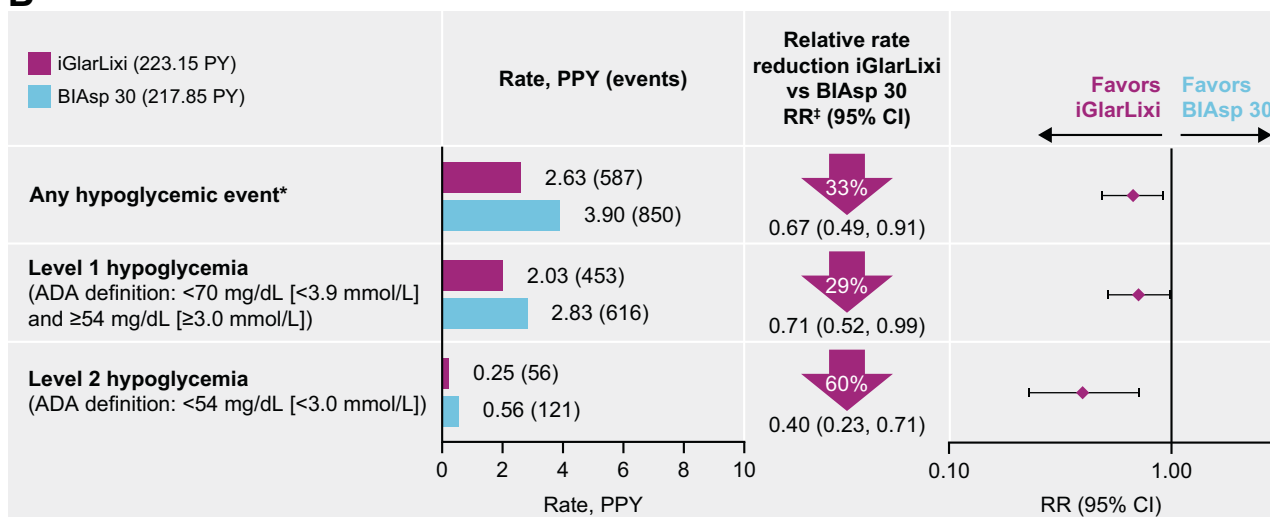


Figure 3—Incidence (A) and rates (B) of hypoglycemic events over the 26-week treatment period (safety population). *A participant can have more than one documented event. †OR for iGlarLixi vs. BIAsp 30 and 95% CI based on logistic regression with treatment group (iGlarLixi and BIAsp 30) and randomization strata (HbA_{1c} $<8.0\%$ and $\geq 8.0\%$, basal insulin dose at screening <30 units and ≥ 30 units, and SGLT2 inhibitor use [yes, no] at screening) as fixed effects. ‡RR for iGlarLixi vs. BIAsp 30 and 95% CI estimated from a negative binomial regression model with a log-link function and the log of the time period in which a hypoglycemia episode is considered treatment emergent as offset. The model included fixed effect terms for treatment group (iGlarLixi and BIAsp 30) and randomization strata (HbA_{1c} $<8.0\%$ and $\geq 8.0\%$, basal insulin dose at screening <30 units and ≥ 30 units, and SGLT2 inhibitor use [yes, no] at screening). *n*, number of participants; PY, participant-years; PPY, per participant-year.

Three severe hypoglycemic episodes (level 3) were reported: one occurred in the iGlarLixi group and two in the BIAsp 30 group. In addition, lower incidence (OR 0.37 [95% CI 0.16, 0.84]) and event rates (rate ratio [RR] 0.28 [95% CI 0.11, 0.71]) of level 2 nocturnal hypoglycemia (defined as occurring between bedtime and waking) were observed in the iGlarLixi group versus the BIAsp 30 group. Similar patterns were seen when using between 0000 and 0600 h to define the nocturnal interval: lower incidence (OR 0.32 [95% CI 0.12, 0.90]) and

event rates (RR 0.30 [95% CI 0.10, 0.88]) were seen with iGlarLixi versus BIAsp 30.

During the 26-week randomized treatment period, the percentage of participants who had at least one AE was slightly higher in the iGlarLixi group (32.6%) compared with the BIAsp 30 group (27.7%), the difference being mainly due to the higher incidence of GI events in the iGlarLixi group (10.4% vs. 2.3%). A large proportion of these GI events were reported in the first week of treatment (Supplementary Fig. 3).

The most commonly reported AE in the iGlarLixi group was nausea (7.7% vs. 0% for BIAsp 30), while nasopharyngitis was the most commonly reported AE in the BIAsp 30 group (2.7% vs. 3.2% for iGlarLixi). In both treatment groups, the majority of participants had AEs considered mild or moderate in severity. SAEs were reported by a similar proportion of participants in both treatment groups (2.7% iGlarLixi and 2.9% BIAsp 30). Overall, the rate of study discontinuation due to an AE was low and similar in both treatment groups (0.9%). There

pandemic occurred during the last few weeks of the study in some countries. Systems were put in place to ensure participant safety, retention, and data capture. Sensitivity analyses in a nonimpacted by COVID-19 ITT population showed that COVID-19 was unlikely to have influenced the results of any end points assessed.

In conclusion, the once-daily FRC, iGlarLixi, is an efficacious and well-tolerated regimen that is simpler for the patient, providing better glycemic control with weight benefit and less hypoglycemia compared with premix BIAsp 30 as an alternative for advancing therapy in people with type 2 diabetes previously suboptimally controlled with basal insulin plus OADs.

Acknowledgments. The authors thank the study participants, trial staff, and investigators for their participation. The authors thank Elisabeth Souhami (Sanofi) for a courtesy review from a medical and scientific perspective, Mathieu Couderc (Sanofi) for a courtesy review from a statistical perspective, and Ana Merino-Trigo (Sanofi) for coordinating manuscript development, facilitating author discussions, and review of this manuscript. Editorial assistance was provided by Fishawack Communications Ltd., part of Fishawack Health.

Duality of Interest. Sponsorship for this study was funded by Sanofi, Paris, France. Editorial assistance provided by Fishawack Communications Ltd. was funded by Sanofi. J.R. has served on advisory panels for Applied Therapeutics, Boehringer Ingelheim, Eli Lilly, Intarcia, Janssen, Novo Nordisk, Oramed, Sanofi, and Zealand and has received research support from Applied Therapeutics, AstraZeneca, Boehringer Ingelheim, Eli Lilly, Genentech, GlaxoSmithKline, Intarcia, Janssen, Lexicon, Merck, Novo Nordisk, Oramed, Pfizer, and Sanofi. R.E. has acted as an advisor and speaker for AstraZeneca, Boehringer Ingelheim, Lilly, Merck Sharp & Dohme (MSD), Novo Nordisk, and Sanofi. L.S.-R. has acted as an advisor and speaker for Novo Nordisk, Sanofi, and Boehringer Ingelheim. N.L. has acted as a speaker for Sanofi, AstraZeneca, Boehringer Ingelheim, Eli Lilly, Novo Nordisk, and Merck. V.M. has served on advisory panels and has acted as a speaker for Abbott, Boehringer Ingelheim, Eli Lilly, MSD, Novo Nordisk, Sanofi, and several Indian pharmaceutical companies. He has also received research grants from some of these companies. C.T. has been an investigator in clinical trials for Eli Lilly and Sanofi and has acted as a speaker for AstraZeneca, Boehringer Ingelheim, Eli Lilly, Novo Nordisk, and Sanofi. S.A.S. has acted as an advisor and speaker for AstraZeneca, Boehringer Ingelheim, Eli Lilly, MSD, Novartis, Novo Nordisk, and Sanofi. A.A., M.B., and N.D. are employees of Sanofi and may hold shares and/or stock options in the company. P.P. is an employee of IviData LIFE SCIENCES working as an external contractor on

behalf of Sanofi. R.J.M. has acted as an advisor and speaker for Sanofi and Novo Nordisk. No other potential conflicts of interest relevant to this article were reported.

Author Contributions. All named authors meet the International Committee of Medical Journal Editors criteria for authorship for this article. All authors participated in the interpretation of the data and the writing, reviewing, and editing of the manuscript, and had final responsibility for approving the published version. J.R. and P.P. are the guarantors of this work and, as such, had full access to all the data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis.

Prior Presentation. Parts of this study were presented in abstract form and as an oral presentation at the 81st Scientific Sessions of the American Diabetes Association, New Orleans, LA, 25–29 June 2021.

References

- American Diabetes Association. 6. Glycemic targets: *Standards of Care in Diabetes—2021*. *Diabetes Care* 2021;44(Suppl. 1):S73–S84
- Davies MJ, D'Alessio DA, Fradkin J, et al. Management of hyperglycaemia in type 2 diabetes, 2018. A consensus report by the American Diabetes Association (ADA) and the European Association for the Study of Diabetes (EASD). *Diabetologia* 2018;61:2461–2498
- American Diabetes Association. 9. Pharmacologic approaches to glycemic treatment: *Standards of Care in Diabetes—2021*. *Diabetes Care* 2021;44(Suppl. 1):S111–S124
- Levin PA, Nguyen H, Wittbrodt ET, Kim SC. Glucagon-like peptide-1 receptor agonists: a systematic review of comparative effectiveness research. *Diabetes Metab Syndr Obes* 2017; 10:123–139
- Meece J. Basal insulin intensification in patients with type 2 diabetes: a review. *Diabetes Ther* 2018;9:877–890
- Vijan S, Hayward RA, Ronis DL, Hofer TP. Brief report: the burden of diabetes therapy: implications for the design of effective patient-centered treatment regimens. *J Gen Intern Med* 2005;20:479–482
- Peyrot M, Barnett AH, Meneghini LF, Schumma-Draeger PM. Insulin adherence behaviours and barriers in the multinational Global Attitudes of Patients and Physicians in Insulin Therapy study. *Diabet Med* 2012;29:682–689
- Chang P. Datamonitor Healthcare – Diabetes type 2 disease analysis report. 2020
- Polinski JM, Kim SC, Jiang D, et al. Geographic patterns in patient demographics and insulin use in 18 countries, a global perspective from the multinational observational study assessing insulin use: understanding the challenges associated with progression of therapy (MOSAIC). *BMC Endocr Disord* 2015;15:46
- Aschner P, Gagliardino JJ, Ilkova H, et al. Persistent poor glycaemic control in individuals with type 2 diabetes in developing countries: 12 years of real-world evidence of the International Diabetes Management Practices Study (IDMPS). *Diabetologia* 2020;63:711–721
- Ji LN, Lu JM, Guo XH, et al. Glycemic control among patients in China with type 2 diabetes

mellitus receiving oral drugs or injectables. *BMC Public Health* 2013;13:602

- Blonde L, Anderson JE, Chava P, Dendy JA. Rationale for a titratable fixed-ratio co-formulation of a basal insulin analog and a glucagon-like peptide 1 receptor agonist in patients with type 2 diabetes. *Curr Med Res Opin* 2019;35:793–804
- Highlights of prescribing information. SOLIQUA 100/33 (insulin glargine and lixisenatide injection), for subcutaneous use, 2019. Accessed 30 June 2020. Available from <https://products.sanofi.us/soliqua100-33/soliqua100-33.pdf>
- Suliqua: EU summary of product characteristics, 2019. Accessed 31 May 2020. Available from <https://www.ema.europa.eu/en/medicines/human/EPAR/suliqua>
- Xultophy 100/3.6: Insulin degludec and liraglutide injection 100 units/mL and 3.6 mg/mL. Highlights of prescribing information, 2019. Accessed 30 April 2020. Available from https://www.accessdata.fda.gov/drugsatfda_docs/label/2019/208583s002s003lbl.pdf
- Xultophy: EU summary of product characteristics, 2020. Accessed 30 April 2020. Available from <https://www.novomedlink.com/content/dam/novonordisk/novomedlink/resources/generaldocuments/Xultophy%20Prescribing%20Information%20-%20FUF.PDF>
- Aroda VR, Rosenstock J, Wysham C, et al.; LixiLan-L Trial Investigators. Efficacy and safety of LixiLan, a titratable fixed-ratio combination of insulin glargine plus Lixisenatide in type 2 diabetes inadequately controlled on basal insulin and metformin: the LixiLan-L randomized trial. *Diabetes Care* 2016;39:1972–1980
- Rosenstock J, Aronson R, Grunberger G, et al.; LixiLan-O Trial Investigators. Benefits of LixiLan, a titratable fixed-ratio combination of insulin glargine plus lixisenatide, versus insulin glargine and lixisenatide monocomponents in type 2 diabetes inadequately controlled on oral agents: the LixiLan-O randomized trial. *Diabetes Care* 2016;39:2026–2035
- Blonde L, Rosenstock J, Del Prato S, et al. Switching to iGlarLixi versus continuing daily or weekly GLP-1 RA in type 2 diabetes inadequately controlled by GLP-1 RA and oral antihyperglycemic therapy: the LixiLan-G randomized clinical trial. *Diabetes Care* 2019;42:2108–2116
- Gough SC, Bode B, Woo V, et al.; NN9068-3697 (DUAL-I) Trial Investigators. Efficacy and safety of a fixed-ratio combination of insulin degludec and liraglutide (IDegLira) compared with its components given alone: results of a phase 3, open-label, randomised, 26-week, treat-to-target trial in insulin-naïve patients with type 2 diabetes. *Lancet Diabetes Endocrinol* 2014;2:885–893
- Buse JB, Vilsbøll T, Thurman J, et al.; NN9068-3912 (DUAL-II) Trial Investigators. Contribution of liraglutide in the fixed-ratio combination of insulin degludec and liraglutide (IDegLira). *Diabetes Care* 2014;37:2926–2933
- Linjawi S, Bode BW, Chaykin LB, et al. The efficacy of IDegLira (insulin degludec/liraglutide combination) in adults with type 2 diabetes inadequately controlled with a GLP-1 receptor agonist and oral therapy: DUAL III randomized clinical trial. *Diabetes Ther* 2017;8:101–114
- McCrimmon RJ, Al Sifri S, Emral R, et al. Advancing therapy with iGlarLixi versus premix BIAsp 30 in basal insulin-treated type 2 diabetes: design and baseline characteristics of the Soli-

Mix randomized controlled trial. *Diabetes Obes Metab* 2021;23:1221–1231

24. Home P, Blonde L, Kalra S, et al. Insulin glargine/lixisenatide fixed-ratio combination (iGlarLixi) compared with premix or addition of meal-time insulin to basal insulin in people with type 2 diabetes: a systematic review and Bayesian network meta-analysis. *Diabetes Obes Metab* 2020;22:2179–2188
25. Liebl A, Prager R, Binz K, Kaiser M, Bergenstal R; PREFER Study Group. Comparison of insulin analogue regimens in people with type 2 diabetes mellitus in the PREFER Study: a randomized controlled trial. *Diabetes Obes Metab* 2009;11:45–52
26. Jin SM, Kim JH, Min KW, et al. Basal-prandial versus premixed insulin in patients with type 2 diabetes requiring insulin intensification after basal insulin optimization: a 24-week randomized non-inferiority trial. *J Diabetes* 2016;8:405–413
27. Ligthelm RJ, Gylvin T, DeLuzio T, Raskin P. A comparison of twice-daily biphasic insulin aspart 70/30 and once-daily insulin glargine in persons with type 2 diabetes mellitus inadequately controlled on basal insulin and oral therapy: a randomized, open-label study. *Endocr Pract* 2011;17:41–50
28. Lingvay I, Pérez Manghi F, García-Hernández P, et al.; DUAL V Investigators. Effect of insulin glargine up-titration vs insulin degludec/liraglutide on glycated hemoglobin levels in patients with uncontrolled type 2 diabetes: the DUAL V randomized clinical trial. *JAMA* 2016;315:898–907
29. LaSalle JR, Berria R. Insulin therapy in type 2 diabetes mellitus: a practical approach for primary care physicians and other health care professionals. *J Am Osteopath Assoc* 2013;113:152–162
30. Tibaldi JM. Intensifying insulin therapy in type 2 diabetes mellitus: dosing options for insulin analogue premixes. *Clin Ther* 2011;33:1630–1642