

Supplementary Material

Impact of Participant Characteristics on Clinical Outcomes with iGlarLixi in Type 2

Diabetes: Post Hoc Analysis of SPARTA Japan

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Supplementary Table S1 Baseline demographics and characteristics by dose increase/dose not increased in the not-achieved group

Characteristic	Insulin naïve			Insulin experienced		
	Achieved	Not achieved		Achieved	Not achieved	
	(n=49)	Dose not	Dose increased	(n=76)	Dose not	Dose increased
		increased (n=36)	(n=50)		increased (n=101)	(n=78)
Age, years	66.1 ± 14.9	61.9 ± 11.1	54.4 ± 11.7***†	66.5 ± 11.9	62.1 ± 13.0*	60.0 ± 10.9**
Age category, n (%)			***†		*	***
<65 years	19 (38.8)	19 (52.8)	40 (80.0)	130 (63.7)	56 (55.5)	53 (68.0)
≥65 to <75 years	15 (30.6)	16 (44.4)	8 (16.0)	25 (32.9)	30 (29.7)	18 (23.1)
≥75 years	15 (30.6)	1 (2.8)	2 (4.0)	23 (30.3)	15 (14.9)	7 (9.0)
Sex, n (%)						
Male	30 (61.2)	24 (66.7)	31 (62.0)	55 (72.4)	58 (57.4)	50 (64.1)
Female	19 (38.8)	12 (33.33)	19 (38.0)	21 (27.6)	43 (42.6)	28 (35.9)
Body weight, kg	67.8 ± 16.3	71.4 ± 19.0	75.8 ± 14.0*	70.4 ± 18.3	68.2 ± 13.9	74.9 ± 14.5†
BMI, kg/m ²	4.7 ± 5.2	26.4 ± 5.6	27.9 ± 3.9	26.4 ± 5.0	25.9 ± 4.1	27.8 ± 4.4†
BMI category, n (%)						†
<25 kg/m ²	21 (42.9)	15 (41.7)	10 (20.0)	32 (42.1)	38 (37.6)	19 (24.4)

≥25 to <30 kg/m ²	19 (38.8)	11 (30.6)	26 (52.0)	94 (46.1)	46 (45.5)	33 (42.3)
≥30 kg/m ²	8 (16.3)	10 (27.8)	13 (26.0)	14 (18.4)	15 (14.9)	22 (28.2)
Duration of T2D, years	11.5 ± 9.3	14.3 ± 9.0	10.1 ± 9.1	13.5 ± 12.1	14.4 ± 11.0	15.4 ± 8.3
Duration of T2D category, n (%)			†			** ††
<10 years	25 (51.0)	11 (30.6)	31 (62.0)	35 (46.1)	46 (45.5)	18 (23.1)
≥10 years	24 (49.0)	25 (69.4)	19 (38.0)	41 (54.0)	55 (54.5)	60 (76.9)
HbA1c, %	8.91 ± 1.55	9.17 ± 1.27	9.86 ± 1.71**	8.21 ± 2.06	8.88 ± 1.48*	8.98 ± 1.37*
HbA1c category, n (%)			*		***	***
<7%	0	0	0	6 (2.9)	3 (3.0)	3 (3.9)
≥7 to <8%	12 (24.5)	4 (11.1)	4 (8.0)	32 (42.1)	23 (22.8)	16 (20.5)
≥8 to <9%	19 (38.8)	14 (38.9)	15 (30.0)	11 (14.5)	39 (38.6)	21 (26.9)
≥8%	18 (36.7)	18 (50.0)	31 (62.0)	16 (21.1)	36 (35.6)	38 (48.7)
Prior treatments, n (%)						
Diet/exercise	7 (14.3)	2 (5.5)	6 (12.0)	0	0	0
OAD	35 (71.4)	24 (66.7)	23 (46.0)*	0	0	0
GLP-1 RA ± OAD	7 (14.3)	10 (27.8)	21 (42.0)**	0	0	0
BOT	0	0	0	18 (23.7)	34 (33.7)	26 (33.3)

GLP-1 RA + BI	0	0	0	15 (19.7)	20 (19.8)	17 (21.8)
Intensive insulin therapy	0	0	0	26 (34.2)	19 (18.8)	13 (16.7)*
Other insulin therapies	0	0	0	17 (22.4)	28 (27.7)	22 (28.2)
History of DPP-4 inhibitor use, n (%)	19 (38.8)	9 (25.0)	12 (24.0)	23 (30.3)	28 (27.7)	14 (18.0)
Type of concomitant OADs, n (%)						
Biguanide	28 (57.1)	27 (75.0)	36 (72.0)	35 (46.1)	61 (60.4)	60 (76.9)***
SGLT2 inhibitors	12 (24.5)	22 (61.1)**	32 (64.0)***	30 (39.5)	70 (69.3)***	49 (62.8)*
Sulfonylureas	15 (30.6)	7 (19.4)	8 (16.0)	5 (6.6)	12 (11.9)	11 (14.1)
Glinides	9 (18.4)	10 (27.8)	9 (18.0)	14 (18.4)	26 (25.7)	19 (24.4)
α-Glucosidase inhibitors	11 (22.5)	5 (13.9)	7 (14.0)	11 (14.5)	15 (14.9)	12 (15.4)
Thiazolidinediones	1 (2.0)	3 (8.3)	2 (4.0)	6 (7.9)	5 (5.0)	2 (2.6)
Number of concomitant OADs, n (%)					**	***
None	9 (18.4)	4 (11.1)	5 (10.0)	17 (22.4)	13 (12.9)	12 (15.4)
1	19 (38.8)	9 (25.0)	9 (18.0)	27 (35.5)	20 (19.8)	11 (14.1)
2	9 (18.4)	10 (27.8)	26 (52.0)	26 (34.2)	45 (44.6)	33 (42.3)
≥3	12 (24.5)	13 (36.1)	10 (20.0)	6 (7.9)	23 (22.8)	22 (28.2)

Data are presented as mean ± SD or n (%). Age-defined glycaemic targets were HbA1c <7.0% for participants aged <65 years, HbA1c <7.5% for participants aged ≥65 to <75 years and HbA1c <8.0% for participants aged ≥75 years

BI basal insulin, *BMI* body mass index, *BOT* basal insulin-supported oral therapy, *GLP-1 RA* glucagon-like peptide 1 receptor agonist, *HbA1c* glycated haemoglobin, *OAD* oral antidiabetic agent, *SD* standard deviation, *SGLT2* sodium-glucose transport protein 2, *T2D* type 2 diabetes

* $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ vs achieved group

† $p < 0.05$, †† $p < 0.01$ vs dose-not-increased group

The Tukey test was used for pair-wise comparisons of continuous variables and Steel-Dwass test for multiple comparisons of categorical variables