

Willingness to pay for diabetes drug therapy based on meta-analysis results

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Objectives

- This study aimed to investigate, in people with type 2 diabetes, the relative willingness to pay (WTP) for different diabetes drug therapies based on outcomes of clinical trials with liraglutide.

Methods

- WTP for diabetes drug therapy in people with type 2 diabetes was assessed by combining results from a meta-analysis of liraglutide compared with other diabetes drug therapies in the liraglutide clinical development programme (Liraglutide Effect and Action in Diabetes; LEAD) and a survey on WTP for important aspects of diabetes medication in people with type 2 diabetes.
- A meta-analysis of six randomised trials with 3967 subjects in the LEAD programme compared the efficacy and safety of liraglutide, a once-daily human glucagon-like peptide-1 analogue, with rosiglitazone, glimepiride, insulin glargine and exenatide (Table 1).
- The WTP survey had 461 participants with type 2 diabetes from Sweden and used a discrete choice experiment methodology to evaluate convenience and clinical effects of treatments in type 2 diabetes.¹ Results, shown in Figure 1, were converted from SEK to EUR (EUR 1 = SEK 10.14).

Results

- Combining meta-analysis and WTP results revealed that people with type 2 diabetes preferred liraglutide to all comparators.
- They were willing to pay an extra EUR 2.49/day for liraglutide 1.2 mg compared with rosiglitazone, EUR 1.82/day compared to glimepiride, EUR 3.17/day compared to insulin glargine and EUR 0.74/day compared to exenatide (Table 2; Figure 2).
- For the comparisons with rosiglitazone, glimepiride and insulin glargine, the largest component was based on the additional weight improvements with liraglutide.
- Compared to exenatide, the largest component of preference was administration of the drug.

Table 1 Main results from the meta-analysis.

Parameters	Liraglutide 1.2 mg	Rosiglitazone	Glimepiride	Insulin glargine	Exenatide
Change in HbA _{1c} , % (26 weeks)	-1.01	-0.35	-0.71	-0.98	-0.82
Change in systolic blood pressure, mmHg (26 weeks)	-2.57	-0.35	0.41	1.64	-3.89
Weight change, kg (26 weeks)	-1.52	1.94	1.04	1.57	-2.29
Hypoglycaemic event rate	0.284	0.134	1.365	1.403	2.669
Blood glucose measure, tests/day ²	0.77	0.77	0.77	1.63	0.77
Nausea, % of patients	4.1	0.2	0.8	0.1	12.2

Table 2 WTP per day for liraglutide 1.2 mg compared to other standard therapies (EUR per day).

Results compared to liraglutide 1.2 mg/day	Rosiglitazone	Glimepiride	Insulin glargine	Exenatide
Change in HbA _{1c} (26 weeks)	0.89	0.40	0.04	0.26
Change in SBP (26 weeks)	0.32	0.43	0.61	-0.19
Change in body weight (26 weeks)	2.53	1.76	2.21	-0.43
Minor hypo event rate (minor + major per patient per year)	0.00	0.03	0.03	0.07
Administration	-1.22	-0.77	0.00	0.98
Blood glucose measure	0.00	0.00	0.31	0.00
Nausea	-0.04	-0.03	-0.04	0.08
Total	2.49	1.82	3.17	0.74

Figure 1 Comparison of diabetes treatment attributes based on WTP calculations. Maroon bars represent WTP for beneficial attributes, grey bars represent amount needed to accept the change.

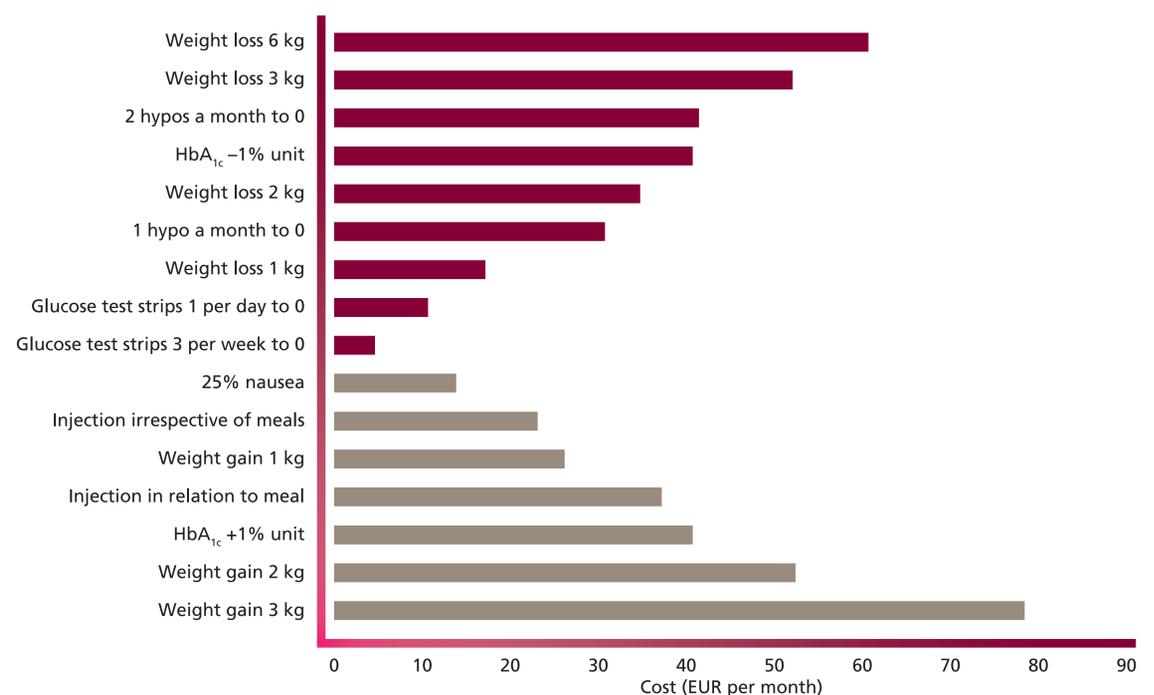
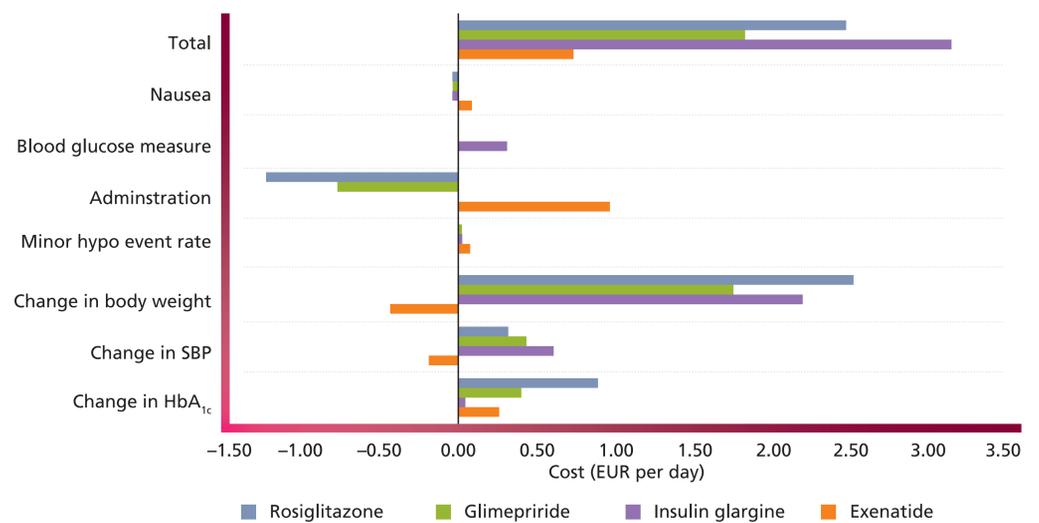


Figure 2 WTP per day for liraglutide 1.2 mg compared to other standard therapies (EUR per day).



References.

- Jendle et al. *Curr Med Res Opin* 2010;26:917-23.
- Gfk (2006): Diabetes Patient Market study – Roper Global Diabetes Programme.

Conclusions

- WTP for liraglutide by people with type 2 diabetes was noticeably higher compared to other standard therapies based on the clinical results from the meta-analysis.
- Primary drivers were weight decrease (compared to rosiglitazone, glimepiride and insulin glargine) and administration (compared to exenatide).
- In total, people were willing to pay up to EUR 3.17/day more to use liraglutide 1.2 mg than other glucose-lowering treatments.