

# Exploring Drivers for Treatment Preferences in Canadian Patients with Type 2 Diabetes:

## A Qualitative Interview Study to Inform a Discrete Choice Experiment

Jones AM<sup>1</sup>, Tatlock S<sup>1</sup>, de Laguiche E<sup>2</sup>, Jensen M S<sup>2</sup>, Kendal H<sup>1</sup>, Wallace S<sup>1</sup>, Mottershead C<sup>3</sup>, Mau G<sup>4</sup>, Besner A<sup>5</sup>

Presenting Author: Elisabeth de Laguiche

<sup>1</sup>Adelphi Values Patient-Centered Outcomes, Cheshire, UK; <sup>2</sup>Novo Nordisk A/S, Søborg, Denmark; <sup>3</sup>Adelphi Research, Cheshire, UK; <sup>4</sup>Novo Nordisk Canada Inc., Mississauga, Canada; <sup>5</sup>Diabetes Canada, Toronto, Canada



**OBJECTIVE:** To explore drivers in people living with type 2 diabetes (T2D) preferences for basal insulin treatment attributes to inform a discrete choice experiment (DCE).



**KEY FINDINGS:** Seven treatment attributes identified through a literature review were relevant to people living with T2D and had sufficient differentiation across insulin treatment profiles. Six of the attributes were reported to be important and influential to treatment decisions by interview participants.



**INTERPRETATION:** There are a wide range of factors people with T2D consider when making treatment decisions. Mode of administration, frequency of administration, dose timing and monitoring and risk of severe hypoglycaemia were identified as potentially important treatment differentiators to people living with T2D and were taken forward into the attributes and levels (A&L) grid incorporated into a DCE.



### Background

- T2D presents a significant challenge to healthcare systems worldwide.<sup>1</sup>
- Understanding preferences of people living with T2D for insulin treatment is critical to optimize treatment strategies, enhance satisfaction and improve overall clinical outcomes.<sup>2</sup>
- This study explored drivers in people living with T2D preferences for basal insulin treatment attributes to inform a DCE.



### Methods

- This study applied best practice guidelines<sup>3,4</sup> for patient preference studies in a three-phase study design, with an advisory panel of clinical experts and representatives of patient advocacy groups engaged at key points throughout the study (Figure 1).
- This poster presents the findings of the phase 1 (targeted literature review) and phase 2 (qualitative interviews). Phase 3 DCE findings are presented in a separate poster – PCR211.

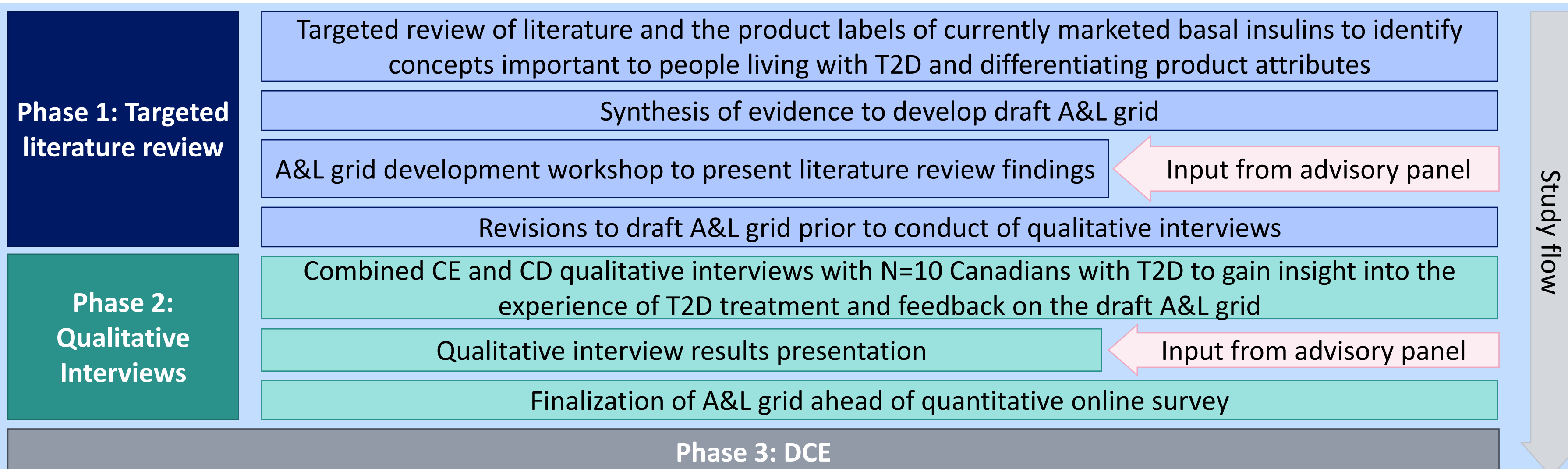


Figure 1. Study design  
A&L = Attributes and levels, CE = Concept elicitation, CD = Cognitive debriefing, T2D = Type 2 diabetes, DCE = Discrete choice experiment



### Results: Literature review

#### Patient-focused literature

- Of the 1192 abstracts identified from searches of bibliographic databases, 10 eligible publications were reviewed.
- Identified concepts were categorized into themes across treatment concepts and health-related quality of life (HRQoL) impacts (Figure 2).

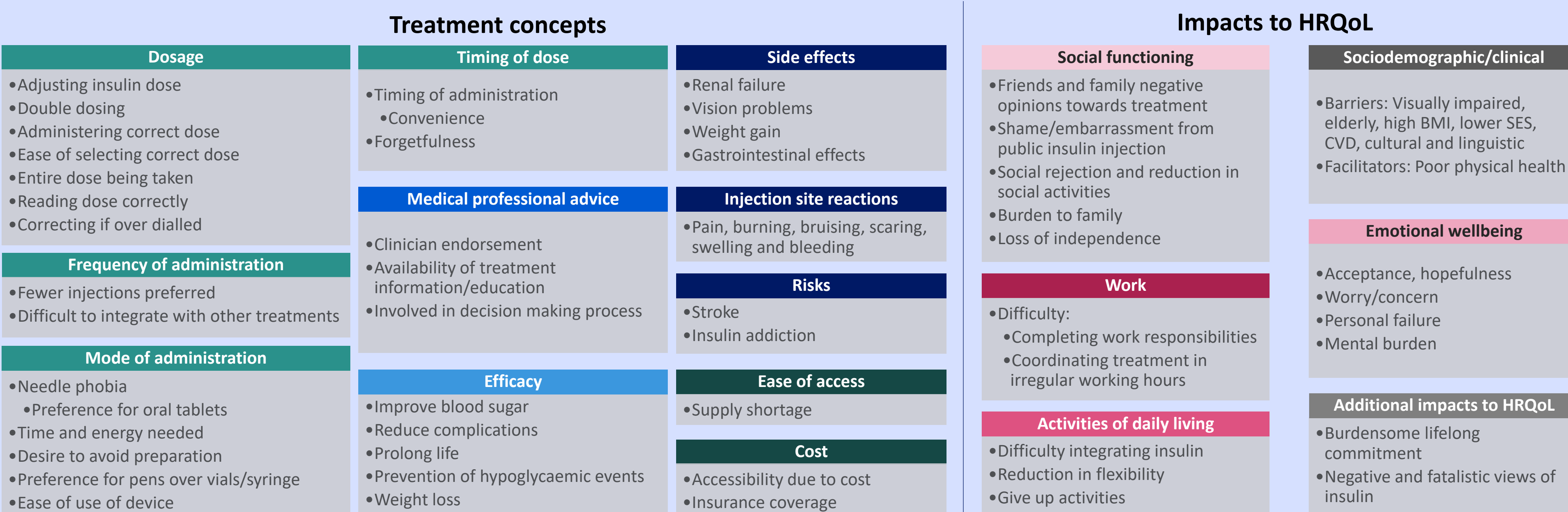


Figure 2. Broad concepts and impacts identified in the targeted literature review of patient-focused literature  
BMI = Body mass index, SES = Socioeconomic status, CVD = Cardiovascular disease

#### Clinical literature and product label review

- Data from six basal insulin products were assessed for differentiating attributes.
- Findings from the patient-focused literature and clinical review were compared to identify concepts which were both important to people living with T2D and differentiated across products (Figure 3) to inform the draft A&L grid.

Differentiating product attributes	Patient-relevant concepts								
	Mode of administration	Frequency of administration	Timing of administration	Dosing	HbA1c reduction	Risk of severe hypoglycemic event	Injection hold time	Onset/half-life/steady state	Storage
Mode of administration	Green								
Frequency of administration		Green							
Timing of administration			Green						
Dosing				Green					
HbA1c reduction					Green				
Risk of severe hypoglycemic event						Green			
Injection site reactions							Green		
Side effects								Green	

Figure 3. Matrix of relevant concepts to people living with T2D and differentiating product attributes  
Green = both important to people living with T2D and differentiated across products



### Results: Qualitative interviews

#### Study sample

- N=10 participants with T2D (aged 22-73; 50:50 female:male; 4.6-18 years since T2D diagnosis) were interviewed.
- A range of demographic and clinical characteristics were represented, available by scanning the QR code.

#### Concept elicitation

- Ten attributes of T2D treatments were reported by participants during the interviews (Figure 4).

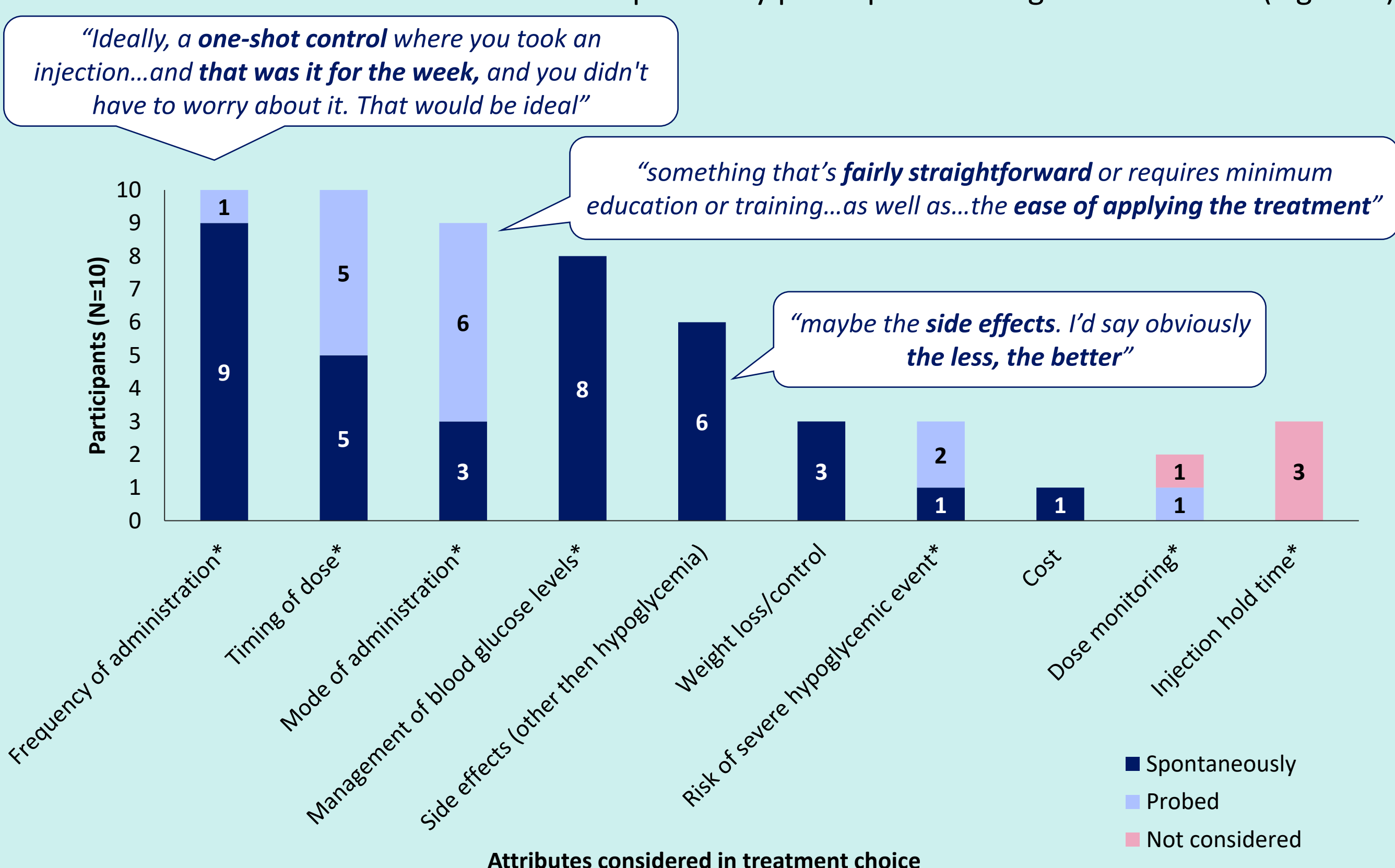


Figure 4. Treatment attributes discussed spontaneously and when probed during qualitative interviews  
Note. Attributes marked with an asterisk (\*) were included in the draft A&L grid and debriefed during the CD section of the interviews.

#### Cognitive debriefing

- Attributes included in the draft A&L grid were understood, important and influential to treatment decisions (scan QR code), except for:
  - Injection hold time which was not influential to the majority of participants;
  - Risk of a severe hypoglycemic event level wording which was not understood by most participants.

#### Final A&L grid

- Edits were made to the A&L grid following the qualitative interviews:
  - Removed HbA1c attributes due to the lack of understanding of the levels and inflated importance compared to other attributes. Instead, HbA1c (as well as cost) were held as constant during the DCE, due to the likelihood they would have dominated treatment preferences;
  - Removed injection hold time due to it not being influential to people living with T2D;
  - Edited the wording to improve participant understanding.
- The final A&L grid implemented in the DCE is shown in Figure 5.

Attributes	Level 1	Level 2	Level 3
Mode of administration	Injection pen that can provide several doses of insulin and is thrown away after 28 days	Injection pen that can provide several doses of insulin and can be refilled with insulin cartridges. The pen is thrown away after 5 years.	
Frequency of administration	Twice daily	Once daily	Once weekly
Timing of dose	Same time for each dose (give or take an hour)	Same time for each dose (give or take three hours)	Anytime for each dose (within 24 hours)
Dose monitoring	You manually record the dose you administered on paper	You manually record the dose you administered into an app	The dose you administered is automatically recorded in an app
Risk of a severe hypoglycemic event (insulin naive rates)	2 out of 100 insulin naive patients who took this insulin for a year experienced a severe hypoglycemic event	1 out of 100 insulin naive patients who took this insulin for a year experienced a severe hypoglycemic event	No insulin naive patients who took this insulin for a year experienced a severe hypoglycemic event
Risk of a severe hypoglycemic event (insulin experienced rates)	6 out of 100 patients who are on insulin took this insulin for a year experienced a severe hypoglycemic event	3 out of 100 patients who are on insulin took this insulin for a year experienced a severe hypoglycemic event	1 out of 100 patients who are on insulin took this insulin for a year experienced a severe hypoglycemic event

Figure 5. Final A&L grid implemented into the DCE



### References

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Supplementary materials