

# Do people prefer to participate in a clinical trial from home?

A discrete choice experiment in persons living with type 2 diabetes mellitus

Julia Kopanz<sup>1</sup>, Bart Lagerwaard<sup>1</sup>, Jorien Veldwijk<sup>2</sup>, Julia K. Mader<sup>3</sup>, Dietrich Tews<sup>4</sup>, Diederick E. Grobbee<sup>1</sup>, Mira G.P. Zuidgeest<sup>1</sup>, on behalf of the Trials@Home Consortium<sup>5</sup>

<sup>1</sup> Julius Center for Health Sciences and Primary Care, University Medical Center Utrecht, The Netherlands

<sup>2</sup> Erasmus School of Health Policy & Management, Erasmus University Rotterdam, The Netherlands

<sup>3</sup> Division of Endocrinology and Diabetology, Medical University of Graz, Austria

<sup>4</sup> MVZ Diabeteszentrum Dr. Tews & Partner, Gelnhausen, Germany

<sup>5</sup> Trialsathome.com



## Purpose

To determine the preferences and trade-offs for participation in clinical trials with different decentralisation levels in persons with type 2 diabetes mellitus.

## Background

- Decentralised clinical trials (DCTs) move trial activities to the participants' direct surroundings and promise to overcome some of the challenges faced by conventional clinical trials.
- There is no evidence available on what potential trial participants prefer regarding trial designs.
- This knowledge is important to design future trials more patient centric, less burdensome and ultimately improve trial conduct.

## Methods

- A discrete choice experiment (DCE) was conducted in three countries:



N = 276



N = ~265



N = ~244

- Participants were asked to complete a survey with 16 DCE choice tasks and background questions.
- Each choice task comprised of three options: two trial options described by six attributes and an opt-out option.
- Panel Mixed Multinomial Logit models were used in the interim data analyses.

## Results

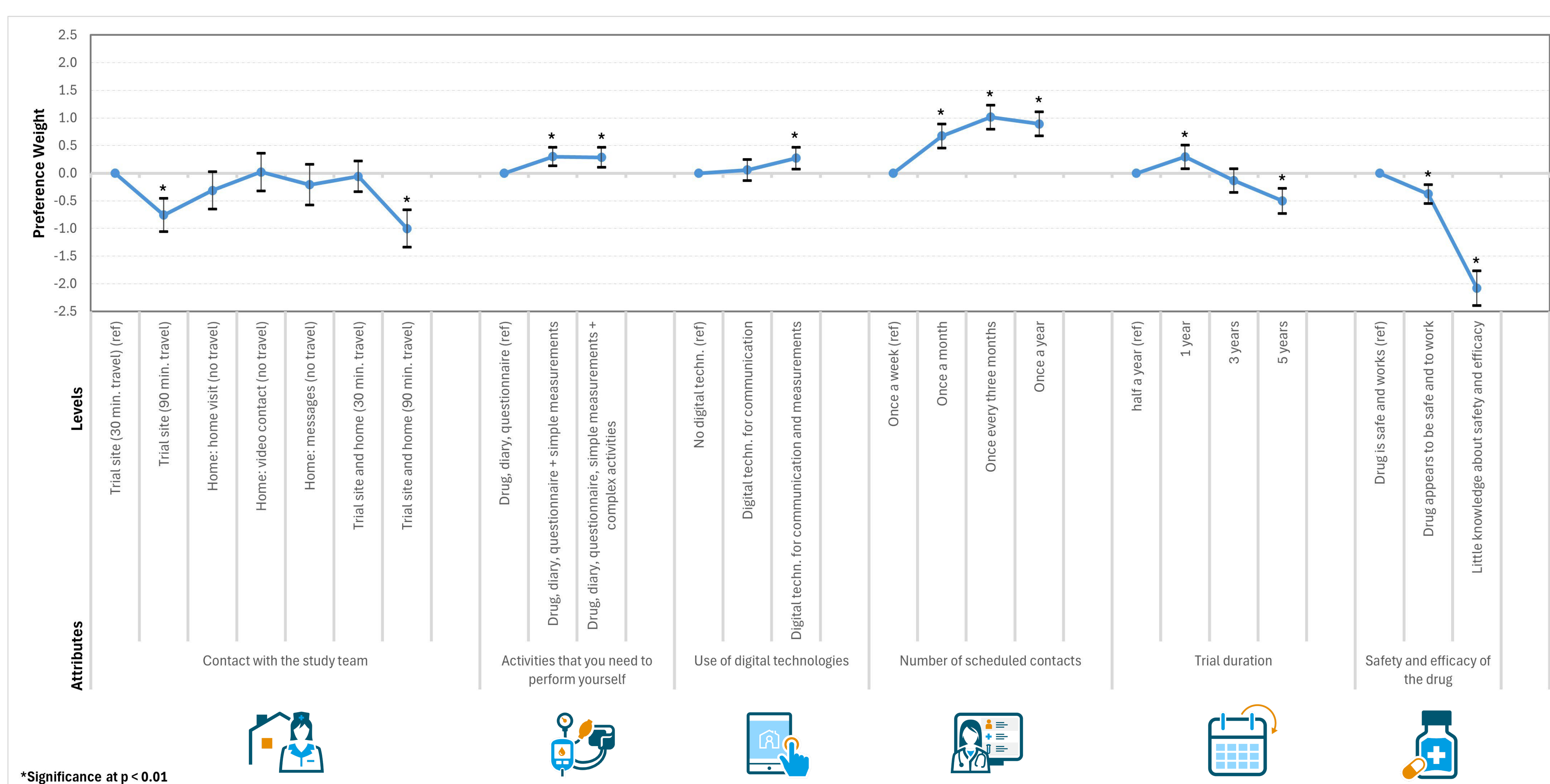


Figure 1: Attribute preference weights

## Interim results on the Dutch population

- 39% female
- median age: 66 years [IQR: 58-71]
- 19% had trial experience
- Trial participation was preferred over opting out (mean coefficient: -0.354) (Figure 1)
- Relative attribute importance is given in Figure 2
- Preference heterogeneity was significant for all attributes

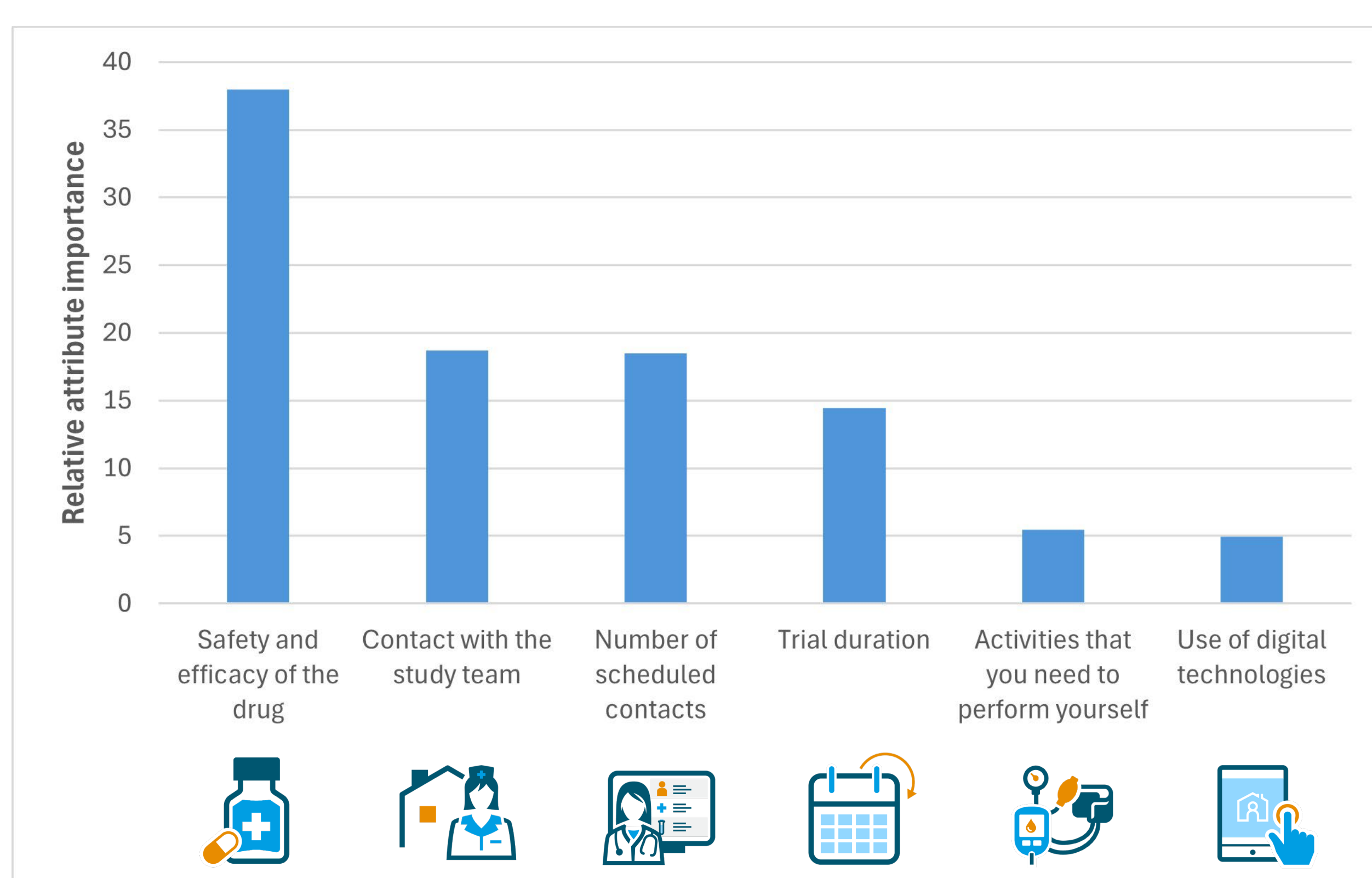


Figure 2: Relative attribute importance

## Conclusion

- ✓ Safety and efficacy of a drug was the most important attribute for Dutch respondents when deciding whether to take part in a trial relative to the other attributes.
- ✓ Further analyses will be performed to investigate preferences in certain subgroups and to predict the uptake of more decentralised trial scenarios.
- ✓ Next steps: finalise analyses of DCE data for all three countries.

University Medical Center Utrecht, Corresponding author: Julia Kopanz, Email: J.Kopanz@umcutrecht.nl