

# OneLook: Non invasive blood glucose measurement (pilot study)

<b>Submission date</b> 29/04/2010	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 29/04/2010	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
<b>Last Edited</b> 05/08/2021	<b>Condition category</b> Nutritional, Metabolic, Endocrine	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

**Plain English summary of protocol**  
Not provided at time of registration

## Contact information

**Type(s)**  
Scientific

**Contact name**  
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## Additional identifiers

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

**Secondary identifying numbers**  
5296

## Study information

**Scientific Title**

A non-randomised interventional clinical laboratory study to assess whether Lein's novel non-invasive glucose measurement technique offers a viable alternative to the current invasive capillary blood glucose solution

## **Acronym**

DRN208

## **Study objectives**

To assess whether Lein's novel non-invasive glucose measurement technique offers a viable alternative to the current invasive capillary blood glucose solution.

The purpose of this trial was to check for correlation between measured results taken from Lein's meter, the current leading capillary glucose meters and a gold standard venous blood laboratory measurement.

## **Ethics approval required**

Old ethics approval format

## **Ethics approval(s)**

National Research Ethics Service - Berkshire Research Ethics Committee approved on the 3rd June 2008 (ref: 08/H0505/70)

## **Study design**

Non-randomised interventional and observational process of care clinical laboratory study

## **Primary study design**

Observational

## **Secondary study design**

Cohort study

## **Study setting(s)**

GP practice

## **Study type(s)**

Diagnostic

## **Participant information sheet**

Not available in web format, please use the contact details below to request a patient information sheet

## **Health condition(s) or problem(s) studied**

Topic: Diabetes Research Network; Subtopic: Both; Disease: Device studies

## **Interventions**

Volunteers were tested over a period of 4 hours with blood glucose readings taken with conventional finger stick meters and a gold standard venous blood meter every 15 minutes. Eye data was also collected every 15 minutes in order to enable the investigating team to compare the readings. The volunteers were provided with a sandwich lunch part way through the trial in order to produce a change in their blood glucose levels.

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome measure**

Measurements were performed at specific points in time that compared various meters. The results were analysed using multi-level and mixed models in the statistical software package "R".

**Secondary outcome measures**

No secondary outcome measures

**Overall study start date**

11/08/2008

**Completion date**

23/03/2009

**Eligibility****Key inclusion criteria**

1. Age 18 to 75 years, male only
2. Type 2 diabetes for at least 3 months
3. Able and willing to do fingerprick blood glucose testing

**Participant type(s)**

Patient

**Age group**

Adult

**Lower age limit**

18 Years

**Sex**

Male

**Target number of participants**

Planned Sample Size: 30; UK Sample Size: 30

**Key exclusion criteria**

1. Age less than 18 and greater than 75 years
2. Significant renal impairment defined as a serum creatinine above 200 µg/ml
3. History of severe depression or mental instability
4. People with type 1 diabetes
5. Coexistent other serious illness
6. Pregnancy
7. Epilepsy
8. Known human immunodeficiency virus (HIV), hepatitis B, C or other blood borne infection

- 9. Previous laser refractive surgery or cataract surgery
- 10. An ocular refractive error that is not between -5 and +5 dioptres
- 11. Astigmatism over 1 dioptre
- 12. Glaucoma
- 13. Cataract
- 14. Colour blindness

**Date of first enrolment**

11/08/2008

**Date of final enrolment**

23/03/2009

## **Locations**

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

**London Road**

Reading

United Kingdom

RG1 5AQ

## **Sponsor information**

**Organisation**

Lein Applied Diagnostics Ltd (UK)

**Sponsor details**

London Road

Reading

United Kingdom

RG1 5AQ

**Sponsor type**

Industry

**Website**

<http://www.lein-ad.com/>

**ROR**

<https://ror.org/02nwjvq32>

# Funder(s)

## Funder type

Government

## Funder Name

NHS Innovations (UK)

# Results and Publications

## Publication and dissemination plan

Not provided at time of registration

Results presented as a poster entitled "OneLook Non-invasive Blood Glucose Measurement" at the International Diabetes Federation conference in Montreal in October 2009.

## Intention to publish date

## Individual participant data (IPD) sharing plan

## IPD sharing plan summary

Not provided at time of registration