

DESMOND (Diabetes Education and Self Management for Ongoing and Newly Diagnosed) Ongoing Study

Submission date 16/11/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 03/03/2011	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 04/02/2019	Condition category Nutritional, Metabolic, Endocrine	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
UNOLE 0106

Study information

Scientific Title

To test an integrated approach for promoting effective self management in people with established Type 2 Diabetes Mellitus (T2DM): a cluster randomised controlled trial of the DESMOND Ongoing education programme

Study objectives

An approach integrating ongoing self-management education with clinical management in those with established Type 2 Diabetes Mellitus (T2DM) within a primary care setting will improve glycaemic control, other biomedical parameters and patient level outcomes

Ethics approval required

Old ethics approval format

Ethics approval(s)

Leicestershire Northampton and Rutland Committee 1, 07/09/2010, ref:10/H0406/54

Study design

Multicentre cluster randomised controlled trial with randomisation at the level of the GP Practice

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Type 2 diabetes mellitus (T2DM)

Interventions

Patients in the intervention group will receive the following:

1. DESMOND Foundation. An existing DESMOND 6-hour group education course which provides introductory information for managing type-2 diabetes
2. Six group sessions of DESMOND-Ongoing (4 in the 1st year and 2 in the 2nd year) delivered by specially trained DESMOND Educators which concentrate on developing self-management skills in people with established type-2 diabetes and covers specific topics related to type-2 diabetes including diet, exercise, managing hypo- and hyper-glycaemia
3. Two Care Planning sessions (delivered at yearly intervals) delivered at the level of the GP Practice by trained GPs and/or Practice Nurses. Care Planning focuses on 'looking with rather than at' someone with diabetes to plan their care and areas for improvement over the forthcoming year. Patients are helped to identify goals and make action plans to achieve these. The healthcare professionals delivering the Care Planning sessions will first be given in-depth training in Care Planning by DESMOND Educators within the study group.

The intervention is a rolling intervention. In the real world it will comprise a six hour introductory session (DESMOND Foundation), a Care Planning session, and then four rolling sessions of DESMOND ongoing delivered at 6 monthly intervals (meaning that the complete material is delivered in 2 years. However, for the trial we are condensing the intervention during the first years as a proof of concept and then will change to the frequency that it would be delivered in the real world during the 2nd year. Therefore intervention patients will receive the six sessions mentioned above during the first year and then a further Care Planning session and 2 sessions of Going Forward with Diabetes during the 2nd year.

Intervention Type

Behavioural

Primary outcome(s)

Reduction in HbA1c at 24 months

Key secondary outcome(s)

1. Reduction in the following biomedical measures measured at 12 and 24 months

- 1.1. HbA1c (at 12 months)
- 1.2. Cholesterol (total, HDL and LDL)
- 1.3. Triglyceride
- 1.4. Blood pressure (BP) Systolic
- 1.5. BP Diastolic
- 1.6. Weight
- 1.7. Waist circumference
- 1.8. Hip circumference

2. Improvement in the following scores at 12 and 24 months

- 2.1. EQ5D
- 2.2. Illness Perceptions Questionnaire revised
- 2.3. Diabetes Illness Representations Questionnaire
- 2.4. Summary of Diabetes Self Care Activities questionnaire
- 2.5. Hospital Anxiety and Depression Score
- 2.6. Problem Areas in Diabetes Score
- 2.7. International Physical Activity Questionnaire

Completion date

30/09/2013

Eligibility**Key inclusion criteria**

Current participant inclusion criteria as of 04/02/2019:

1. Individuals with established Type 2 Diabetes Mellitus
2. Diabetes duration of 12 months or more
3. HbA1c between 7.8 and 11%

Previous participant inclusion criteria:

1. Individuals with established Type 2 Diabetes Mellitus
2. Diabetes duration of 12 months or more
3. HbA1c between 7.0 and 11%

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Receiving insulin therapy
2. Have existing comorbidities that could interfere with participation in the study over the 24 months
3. Unable to give informed consent
4. Severe and enduring mental health problems
5. Patients who are not primarily responsible for their own care
6. Unable to participate in a group programme (e.g. housebound)
7. Insufficient understanding of English to participate in the group education
8. Visually impaired or hard of hearing to the level that they can not participate in the group education
9. Aged under 18 years

Date of first enrolment

01/01/2011

Date of final enrolment

30/09/2013

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Leicester Diabetes Centre (Broadleaf)

Leicester

United Kingdom

LE5 4PW

Sponsor information**Organisation**

University of Leicester (UK)

ROR

<https://ror.org/04h699437>

Funder(s)

Funder type

Government

Funder Name

National Institute for Health Research (NIHR) (UK) - Collaboration for Leadership in Applied Health Research and Care (CLAHRC)

Funder Name

Leicestershire, Northamptonshire and Rutland CLARHC and CLAHRC for South Yorkshire (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Basic results		04/02/2019	04/02/2019	No	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes