

The MILESTONE study: improving transition from child to adult mental health care

Submission date 22/07/2015	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 23/07/2015	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 02/12/2025	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

The MILESTONE study focuses on the period when young people attending a children and adolescents mental health service (CAMHS) need to move on, or “transition”, to an adult mental health service (AMHS), if they still require care or treatment. It is known from other research that this transition is not always properly managed and that improving the transition process can have a positive impact on the health and well-being of young people in this position. The aim of this study is to assess the impact of the different transition experiences on young people’s health and well-being, and whether the process of Managed Transition has any benefits as compared to usual care.

Who can participate?

Young people aged under the regular care of a CAMHS and are reaching transition age (usually 16-18 years old)

What does the study involve?

The MILESTONE study is run in eight European countries (UK, Ireland, Belgium, Holland, France, Germany, Italy and Croatia). CAMHS in the study regions are selected to provide the young people in their care that are reaching transition age either usual care or a novel service called “Managed Transition”, which includes the use of a new decision support tool, the Transition Readiness and Appropriateness Measure (TRAM). This should help with decision making and enable better transitions by identifying cases for whom transition from CAMHS to AMHS is advisable and appropriate, or who can be safely discharged or referred to a community based service. CAMHS are randomly assigned to provide the intervention of Managed Transition or usual care. The health and well-being of the young people attending these services is assessed at the start of the study and then followed-up for 24 months to see whether they transition to AMHS or are discharged or referred to some other service.

What are the possible benefits and risks of participating?

There are no benefits or risks associated with taking part in the study.

Where is the study run from?

University of Warwick (UK)

When is the study starting and how long is it expected to run for?
September 2015 to January 2019

Who is funding the study?
Seventh Framework Programme (Belgium)

Who is the main contact?
Dr Helena Tuomainen

Contact information

Type(s)
Public

Contact name
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Additional identifiers

ClinicalTrials.gov (NCT)
NCT03013595

Protocol serial number
19173

Study information

Scientific Title
The effectiveness of managed transition in improving the health and social outcomes for young people transitioning from child to adult mental health care: the MILESTONE study

Acronym
MILESTONE (ManagIng the Link and Strengthening TransitiON from child to adult mental health carE)

Study objectives
Do young people reaching the transition boundary of their child and adolescent mental health service benefit from the implementation of a model of managed transition regarding their health and social outcomes, and transition to adult roles, as compared to usual care?

Ethics approval required
Old ethics approval format

Ethics approval(s)

NRES Committee West Midlands – South Birmingham, 24/02/2015, ref: 15/WM/0052

Study design

Both; Interventional and Observational; Design type: Process of Care, Cohort study

Primary study design

Interventional

Study type(s)

Quality of life

Health condition(s) or problem(s) studied

Topic: Children, Mental Health; Subtopic: All Diagnoses, Psychosis; Disease: Psychosis, All Diseases

Interventions

Intervention arm: The model of managed transition includes:

1. The completion of the "Transition Readiness and Appropriateness Measure" (TRAM, a standardized structured assessment) prior to the transition boundary by the child and adolescent mental health service (CAMHS) clinician, young person and parent/carer.
2. Feedback of TRAM findings to the CAMHS clinician to support decisions made regarding transition, communication with stakeholders and the transition process. Clinicians will be expected to communicate the findings to the young person and parent/carer, and, if a referral is made, to send the TRAM feedback to the adult clinician along with the referral letter.
3. The clinicians will also receive information prior to recruitment begin on the use of TRAM and the way in which it fits in with optimal transition.

The control arm: Patients, parent/carers and clinicians in the control arm will complete the TRAM prior to the transition boundary, but the clinicians won't receive any feedback from it nor any information on the benefits of using the decision support tool.

Intervention Type

Behavioural

Primary outcome(s)

Current primary outcome measures as of 23/01/2017:

Mental health status (need for care), measured by the use of the clinician's version of the Health of the Nation Outcome Scale for Children and Adolescents (HoNOSCA) at 15 months. The measure will be completed by a trained MILESTONE research assistant taking into account all available sources of information (including the young person, parent/carer, relevant clinician and the medical records) to ensure accuracy of data.

Previous primary outcome measures:

Mental health status (need for care), measured by the use of the clinician's version of the Health of the Nation Outcome Scale for Children and Adolescents (HoNOSCA) at baseline, 9, 15, and 24 months. The measure will be completed by a trained MILESTONE research assistant taking into account all available sources of information (including the young person, parent/carer, relevant clinician and the medical records) to ensure accuracy of data.

Key secondary outcome(s)

Current secondary outcome measures as of 20/06/2017:

1. Self-rated (i.e. completed by young person) mental health status (need for care), on Health of the Nation Outcome Scale for Children and Adolescents (HONOSCA) at baseline, 9, 15, and 24 months
 2. Transition outcome, assessed by the Transition Related Outcome Measure (TROM) (young person, parent/carer, clinician) at 9, 15, and 24 months
 3. Self-reported and parent/carer reported psychopathology, assessed by The Development and Well-being Assessment (DAWBA) (Young person, parent/carer) at baseline and 24 months and clinical diagnosis (clinician) at baseline, 9, 15 and 24 months
 4. Emotional and behavioural problems, assessed by the ASEBA Youth Self Report (YSR)/Adult Self Report (ASR) questionnaires (young person) and CBCL/ABCL questionnaires (parent/carer) at baseline, 9, 15, and 24 months
 5. Illness severity, assessed by Clinical Global Impression Severity scale (CGIseverity) (clinician) at baseline, 9, 15, and 24 months
 6. Quality of life assessed by WHOQOLBREF (young person) at baseline, 15 and 24 months
 7. Independent behaviour, assessed by Independent Behaviour During Consultation Scale (IBDCS) (young person) at baseline, 9, 15, and 24 months, if young person is a current service user
 8. Illness perception, assessed by the Brief Illness Perception Questionnaire (BIPQ) (young person) at baseline and 24 months
 9. Barriers to care, assessed by Barriers to Care (BtC) checklist (young person) at 9, 15, and 24 months, if young person is no longer a service user
 10. Transition experience and readiness assessed by OYOF-TES – Transition Experience Scale (young person, parent/carer) completed at 9, 15 or 24 months, completed only once at the first assessment after transition
 11. Adult functioning, assessed by Specific Levels of Functioning Scale (SLOF) (parent/carer) at baseline and 24 months.
 12. Cost-effectiveness, assessed by EQ-5D-5L (young person) at baseline, 9, 15, and 24 months
 13. Service use assessed by a MILESTONE specific Client Service Receipt Inventory (CSRI) at baseline, 9, 15, and 24 months (young person) and 9 and 24 months (parent/carer)
 14. Mental health status (need for care) on clinician's version of the Health of the Nation Outcome Scale for Children and Adolescents (HoNOSCA) at baseline, 9 and 24 months. The measure will be completed by a trained MILESTONE research assistant taking into account all available sources of information (including the young person, parent/carer, relevant clinician and the medical records) to ensure accuracy of data
- All measures are completed by online survey

Previous secondary outcome measures from 23/01/2017 to 20/06/2017:

1. Self-rated (i.e. completed by young person) mental health status (need for care), on Health of the Nation Outcome Scale for Children and Adolescents (HONOSCA) at baseline, 9, 15, and 24 months
2. Transition outcome, assessed by the Transition Outcome Measure (TROM) (young person, parent/carer, clinician) at 9, 15, and 24 months
3. Self-reported and parent/carer reported psychopathology, assessed by The Development and Well-being Assessment (DAWBA) (Young person, parent/carer) and clinical diagnosis (clinician) at baseline and 24 months
4. Emotional and behavioural problems, assessed by the ASEBA Youth Self Report (YSR)/Adult Self Report (ASR) questionnaires (young person) and CBCL/ABCL questionnaires (parent/carer) at baseline, 9, 15, and 24 months
5. Illness severity, assessed by Clinical Global Impression Severity scale (CGIseverity) (clinician) at baseline, 9, 15, and 24 months
6. Quality of life assessed by WHOQOLBREF (young person) at baseline, 15 and 24 months
7. Independent behaviour, assessed by Independent Behaviour During Consultation Scale

(IBDCS) (young person) at baseline, 9, 15, and 24 months, if young person is a current service user

8. Illness perception, assessed by the Brief Illness Perception Questionnaire (BIPQ) (young person) at baseline and 24 months
9. Barriers to care, assessed by Barriers to Care (BtC) checklist (young person) at 9, 15, and 24 months, if young person is no longer a service user
10. Transition experience and readiness assessed by OYOF-TES – Transition Experience Scale (young person, parent/carer) at 9 or 15 months, completed only once at the first assessment after transition
11. Adult functioning, assessed by Specific Levels of Functioning Scale (SLOF) (parent/carer) at baseline and 24 months.
12. Cost-effectiveness, assessed by EQ-5D-5L (young person) at baseline, 9, 15, and 24 months
13. Service use assessed by a MILESTONE specific Client Service Receipt Inventory (CSRI) at baseline, 9, 15, and 24 months (young person) and 9 and 24 months (parent/carer)
14. Mental health status (need for care) on clinician's version of the Health of the Nation Outcome Scale for Children and Adolescents (HoNOSCA) at baseline, 9 and 24 months. The measure will be completed by a trained MILESTONE research assistant taking into account all available sources of information (including the young person, parent/carer, relevant clinician and the medical records) to ensure accuracy of data

All measures are completed by online survey

Original secondary outcome measures:

1. Self-rated (i.e. completed by young person) mental health status (need for care), on Health of the Nation Outcome Scale for Children and Adolescents (HONOSCA) at baseline, 9, 15, and 24 months
2. Transition outcome, assessed by the Transition Outcome Measure (TROM) (young person, parent/carer, clinician) at 9, 15, and 24 months
3. Self-reported and parent/carer reported psychopathology, assessed by The Development and Well-being Assessment (DAWBA) (Young person, parent/carer) and clinical diagnosis (clinician) at baseline and 24 months
4. Emotional and behavioural problems, assessed by the ASEBA Youth Self Report (YSR)/Adult Self Report (ASR) questionnaires (young person) and CBCL/ABCL questionnaires (parent/carer) at baseline, 9, 15, and 24 months
5. Illness severity, assessed by Clinical Global Impression Severity scale (CGIseverity) (clinician) at baseline, 9, 15, and 24 months
6. Quality of life, assessed by WHOQOLBREF (young person) at baseline and 24 months
7. Independent behaviour, assessed by Independent Behaviour During Consultation Scale (IBDCS) (young person) at baseline, 9, 15, and 24 months, if young person is a current service user
8. Illness perception, assessed by the Brief Illness Perception Questionnaire (BIPQ) (young person) at baseline and 24 months
9. Barriers to care, assessed by Barriers to Care (BtC) checklist (young person) at 9, 15, and 24 months, if young person is no longer a service user
10. Transition experience and readiness, assessed by OYOF-TES – Transition Experience Scale (young person, parent/carer) at 9 or 18 months, completed only once at the first assessment after transition
11. Adult functioning, assessed by Specific Levels of Functioning Scale (SLOF) (parent/carer) at baseline and 24 months.
12. Cost-effectiveness, assessed by EQ-5D-5L (young person) at baseline, 9, 15, and 24 months
13. Service use, assessed by a MILESTONE specific Client Service Receipt Inventory (CSRI) at baseline, 9, 15, and 27 months (young person) and 9 and 24 months (parent/carer)

All measures are completed by online survey

Completion date

Eligibility

Key inclusion criteria

Young person inclusion:

1. Provides valid written informed consent, or assent, if below the legal age of consent
2. If age is within 1 year of reaching the transition boundary of their CAMHS during the trial recruitment period and (added 23/01/2017) in exceptional cases, not more than 3 months older than the transition boundary, if a decision about transition has not yet been made
3. Has a mental disorder defined by DSM-IV-TR, DSM-5 or ICD 10/11, or is under the regular care of CAMHS (if not yet diagnosed)
4. Has an IQ ≥ 70 as ascertained by previous standardised assessment or diagnosed by clinician, or no indication of intellectual impairment

Parent/carer inclusion:

1. Only one parent/carer per young person can be recruited into the study, the aim being to engage the same individual throughout the study. If the latter is not possible, then the aim is to involve another parent/carer
2. If the young person doesn't live with his/her biological parent/s, then his/her carer will be involved. A carer may be the legal guardian or a partner or an older adult sibling, or another individual living with and/or providing regular support to the young person
3. For young people under the legal age of consent, the parent/carer has to be the legal guardian of the young person
4. Young person consents to parent/carer participation
5. Parent/carer provides a valid written informed consent

Clinician/care provider inclusion:

1. Is responsible for the main care for young person at CAMHS (and AMHS or other relevant service provider, if referral is made)
2. Provides a valid written informed consent

Participant type(s)

Carer, Patient

Healthy volunteers allowed

No

Age group

Mixed

Sex

All

Total final enrolment

844

Key exclusion criteria

Young person exclusion:

1. Does not provide valid written informed consent, or assent, if below the legal age of consent
2. Is younger than a year before the transition boundary of their CAMHS

3. Has intellectual impairment (IQ <70) as ascertained by previous standardised assessment or diagnosed by clinician (if no data on intellectual functioning are available [because it has never been assessed] then care coordinators will be asked to make a clinical judgement on intellectual impairment before baseline assessment takes place)
4. If not able to (or expected not to be able to) complete the questionnaires due to severe physical disabilities or language problems, even with assistance from family members or research assistant
5. Service user in a secure forensic institution

Parent or carer refusal or inability to participate him/herself in assessments may prevent the participation of the service user, yet this will depend on the individual country's legal and ethical situation. In the UK, young people over the age of 16 will be able to take part even if their parent /carer does not take part.

Parent/carers exclusion:

1. A parent/carers who does not live with and/or provide regular support to young person
2. Young person does not provide consent for parent/carers participation
3. Parent/carers does not provide a valid written consent
4. If not able to (or expected not to be able to) complete the questionnaires due to severe physical disabilities or language problems, even with assistance from family members or research assistant

Clinician/care provider exclusion:

1. Does not provide care for young person, or only provides intermittent care at CAMHS (and AMHS or other relevant service provider, if referral is made)
2. Does not provide a valid written informed consent

Date of first enrolment

01/09/2015

Date of final enrolment

31/01/2017

Locations

Countries of recruitment

United Kingdom

England

Belgium

Croatia

France

Germany

Ireland

Italy

Netherlands

Study participating centre

University of Warwick

Gibbet Hill Road

Coventry

England

CV4 7AL

Sponsor information

Organisation

University of Warwick

ROR

<https://ror.org/01a77tt86>

Funder(s)

Funder type

Government

Funder Name

Seventh Framework Programme

Alternative Name(s)

EC Seventh Framework Programme, European Commission Seventh Framework Programme, EU Seventh Framework Programme, European Union Seventh Framework Programme, FP7

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

Results and Publications

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are/will be available upon request from Dr Helena Tuomainen (helena.tuomainen@warwick.ac.uk)

Added 12/06/2023:

The type of data that will be shared: Baseline and follow-up data

Whether consent from participants was required and obtained: Yes

Comments on data anonymization: All data is anonymous

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article		29/10/2021	12/06/2023	Yes	No
Results article	Demographic, clinical, and service-use characteristics results	10/02/2022	03/09/2024	Yes	No
Results article	Mental health trajectories of adolescents treated with psychotropic medications: insights from the european milestone study	17/10/2025	21/10/2025	Yes	No
Results article	Bridging Perspectives: Clinician-Adolescent Agreement on Psychopathological Severity in the European MILESTONE Cohort	30/11/2025	02/12/2025	Yes	No
Protocol article	TRAM and TROM development protocol	16/04/2020	29/01/2021	Yes	No
Protocol article	protocol	16/10/2017	29/01/2021	Yes	No
HRA research summary			28/06/2023	No	No
Interim results article	demographic and clinical characteristics	16/12/2021	20/06/2022	Yes	No
Other publications	background, rationale and methodology	04/06/2018		Yes	No
Other publications	TRAM validation	23/06/2020	29/01/2021	Yes	No
Study website	Study website	11/11/2025	11/11/2025	No	Yes