Ready for SDM - evaluation of a multidisciplinary training module in Shared Decision Making

Submission date	Recruitment status No longer recruiting	Prospectively registered		
17/04/2018		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
09/05/2018	Completed	[X] Results		
Last Edited	Condition category	Individual participant data		
10/06/2024	Other			

Plain English summary of protocol

Background and study aims

In response to an obvious lack of professional training about shared decision making (SDM) in Norway, a draft of the "Ready for SDM" program was developed based on a proven effective German module (doktormitSDM). Two applications (Norw. "Klar for samvalg") (M1 / M2) have recently been tested in a pilot study, indicating a need for better adaptation to various health professions and inter-professional learning. This study aims at testing the effectiveness of the revised M2 regarding improvement of SDM-related evaluation competences.

Who can participate?

6-10 District Psychiatric Centers recruited from the South – Eastern Norway Regional Health Authority and the Western Norway Regional Health Authority

What does the study involve?

Participating District Psychiatric Centers are randomly allocated to either an intervention or a waiting control group. After a demographic questionnaire, the intervention group starts with the introduction module A, followed by B: SDM in-depth, and C: interactive training. After completion of modules A-C the intervention group participants are assessed, and the control group participants are assessed as long as the group is untrained (has not yet received the training module). After the study, the training module is also conducted for participants in the control group.

What are the possible benefits and risks of participating?

Participants taking part in this study may benefit by getting increased knowledge and skills in SDM, furthermore it may have future benefits, as the results of the study are likely to influence the implementation of SDM in Norway. There are no notable risks to participants in this study, and both groups receive the same training.

Where is the study run from? University Hospital of North Norway

When is the study starting and how long is it expected to run for? March 2016 to June 2019

Who is funding the study? Northern Norway Regional Health Authority

Who is the main contact? Simone Kienlin simone.kienlin@helse-sorost.no

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

Protocol serial number 15154

Study information

Scientific Title

Evaluation of a multidisciplinary training module in Shared Decision Making (Ready for SDM): a cluster randomized controlled trial

Acronym

Ready for SDM

Study objectives

The specialist health service in Norway includes the district Psychiatric Outpatient Services (DPS), where both patients with acute problems and those who need long-term follow-up can receive help. The intervention provided in this study addresses patient involvement in making medical decisions. The overarching aim of this approach is to enhance patient involvement by provision of evidence-based patient information, alongside with the patients' encouragement to participate in the decision process and thereby to increase the frequency of patient led

informed choices. Patient involvement in this study is approached to via training of health professionals' communication skills. The psychiatric domain has been chosen as the medical context for the study, to demonstrate efficacy of the intervention.

Hypothesis: A 2-hour group training module can improve health professionals' communication competencies in Shared Decision Making (SDM), operationalized as accuracy of observer judgements regarding realized SDM made by use of the MAPPIN'SDM observer instrument.

Ethics approval required

Old ethics approval format

Ethics approval(s)

- 1. The local ethics committee at the University Hospital in Northern Norway (UNN), ref: 2017 /1461
- 2. South-eastern regional ethics committee, ref: 2017/82 C

Study design

Cluster randomized waiting control group trial

Primary study design

Interventional

Study type(s)

Other

Health condition(s) or problem(s) studied

Shared decision making

Interventions

Participants are randomized to either an intervention or a waiting control group:

Intervention: an SDM training module addressing health professionals, which includes the following:

- 1. 30-min didactics, introduction to SDM
- 2. 30-min deepening in SDM
- 3. Interactive SDM observer training using a video-recording of a psychiatric consultation and a set of evaluation criteria

After administering a demographic questionnaire, the intervention group starts with the introduction module A, followed by B: SDM in-depth, and C: interactive training. After completion of modules A-C, primary and secondary endpoints are assessed.

Control: waiting/no intervention

Primary and secondary endpoints will be assessed as long as the group is untrained (has not yet received the training module). After the study, the training module will also be conducted for participants in the control group.

Intervention Type

Behavioural

Primary outcome(s)

Accuracy of observation-based judgements made on SDM performance in a given medical consultation, which is provided as a video-recorded consultation. Judgements will be made in terms of the dyadic MAPPIN´SDM immediately after provision of the SDM training module or the waiting condition, respectively. Accuracy is calculated using weighted T coefficients calculated pairwise between participants' judgements and an expert standard rating.

Key secondary outcome(s))

SDM related knowledge assessed by using a five-item multiple choice knowledge test, previously used in a similar study. The level of achieved knowledge will be assessed immediately after provision of the SDM training module or the waiting condition, respectively.

Completion date

16/06/2019

Eligibility

Key inclusion criteria

- 1. Health professional teams from District Psychiatric Centers
- 2. Actively involved in psychiatric patient care

Participant type(s)

Health professional

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Total final enrolment

357

Key exclusion criteria

- 1. Non-health professionals and professionals from other settings than the pre-defined institutions
- 2. Health professionals working in administrative functions

Date of first enrolment

16/04/2018

Date of final enrolment

16/06/2018

Locations

Countries of recruitment

Norway

Study participating centre Betanien District Psychiatric Center Bergen Norway 5145

Study participating centre
Bjørgvin District Psychiatric Center
Bergen
Norway
5113

Study participating centre Øyane District Psychiatric Center Straume Norway 5353

Study participating centre Kronstad District Psychiatric Center Bergen Norway 5054

Study participating centre Solli District Psychiatric Center Bergen Norway 5228

Study participating centre Voss District Psychiatric Center Voss Norway 5705 Study participating centre Gjøvik District Psychiatric Center Gjøvik Norway 2819

Sponsor information

Organisation

University Hospital of North Norway

ROR

https://ror.org/030v5kp38

Funder(s)

Funder type

Government

Funder Name

Helse Nord RHF

Alternative Name(s)

Northern Norway Regional Health Authority

Funding Body Type

Government organisation

Funding Body Subtype

Local government

Location

Norway

Results and Publications

Individual participant data (IPD) sharing plan

The data sharing plans for the current study are unknown and will be made available at a later date.

IPD sharing plan summary

Data sharing statement to be made available at a later date

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/04/2020	04/06/2020	Yes	No
Results article		30/04/2021	04/10/2022	Yes	No
Results article		17/03/2022			No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes