

Information in English

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More information available at <http://www.norkog.no>



The Norwegian registry of persons assessed for cognitive symptoms

Participation in a national registry

Patients admitted to hospital outpatient clinics for assessment of cognitive symptoms, will be invited to participate in The Norwegian registry of persons assessed for cognitive symptoms (NorCog)

The aim of NorCog

- ❖ To improve the quality of assessment and treatment of cognitive symptoms and dementia
- ❖ To improve and develop assessment methods
- ❖ To ensure equal assessment, by using internationally approved and standardized assessment methods and cognitive tests.
- ❖ To stimulate research regarding cognitive symptoms and dementia
- ❖ To contribute data to decision makers, to improve planning of health care service for persons with cognitive symptoms and dementia

Participation is voluntary

It is voluntary to participate in the registry. Patients can withdraw their consent at any time, without giving any reason. Withdrawing consent will not have consequences for the participant's further treatment.

If you consent to participate, data from a standard assessment will be registered in the NorCog database.

If the patient is not able to give informed consent, the next of kin is invited to consent, on behalf of the patient

Use of data

Data from the registry will be used to generate new knowledge and to improve the quality of assessment, treatment and the care for persons with cognitive impairment and dementia.

On the website www.norkog.no, you will find information about how to withdraw from research projects.

Results

Results from NorCog are presented on the registry website: www.norkog.no