

# Evaluation of the parenting support group intervention "Everyday life and parenting" for parents of preschool children with neurodevelopmental difficulties

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|--|---|---|
| <b>Submission date</b><br>16/08/2024   | <b>Recruitment status</b><br>No longer recruiting             | <input type="checkbox"/> Prospectively registered               |
| <b>Registration date</b><br>08/10/2024 | <b>Overall study status</b><br>Ongoing                        | <input type="checkbox"/> Protocol                               |
| <b>Last Edited</b><br>06/03/2025       | <b>Condition category</b><br>Mental and Behavioural Disorders | <input type="checkbox"/> Statistical analysis plan              |
|  |   | <input type="checkbox"/> Results                                |
|  |   | <input type="checkbox"/> Individual participant data            |
|  |   | <input checked="" type="checkbox"/> Record updated in last year |

## Plain English summary of protocol

### Background and study aims

Young children with subclinical neurodevelopmental disorders (NDDs) and concurrent emotional and behavioral problems (EBP) are at significant risk of negative consequences in the short and long term. Early parenting support interventions are recommended and requested. However, there is a lack of interventions specifically designed for this group and adapted to a Swedish context. Based on this gap, a parenting support intervention for parents with children 2 - 6 years of age with subclinical NDD and EBP has been co-created with clinicians and parents. The current project aims to evaluate the clinical- and cost-effectiveness of this parenting support group intervention.

### Who can participate?

Parents of children aged 2 - 6 years old with subclinical levels of NDD and behavioral problems. The children have been referred to child health psychologists at the Child Pediatric Outpatient Clinic in Uppsala Region. Children aged 4 – 6 years old whose parents consented to participate in the project, will be invited to participate themselves.

### What does the study involve?

Parents offered the parenting support group intervention Everyday life and parenting will be included in the baseline (4, 7 or 10 weeks) and they will thus start the intervention after the baseline period. They will answer 25 questions every week during the baseline, the intervention and four weeks after the intervention. The questions are about their child's behavior, their own self-efficacy as parents and stress. They will also answer questionnaires at the start of baseline, when the intervention is finished and three months after the intervention. Children aged 4 - 6 years old will answer 5 questions weekly during the same time period as their parent.

### What are the possible benefits and risks of participating?

The potential benefit of participating in the research project is that parents are given the

opportunity to reflect on their parenting and thus gain a greater awareness of the family's challenges. Based on this, it can be easier for parents to recognise what support they and their children need. It can also be meaningful to share your own experiences and perspectives and through this contribute to knowledge that can be used to help other parents and children in similar situations.

Children may enjoy answering a questionnaire that asks about their experiences in the family. This can make them feel that their opinions are important. It is rare for children of pre-school age to be involved in similar studies and this project gives them the opportunity to make their voices heard and thus fulfil a human right.

Where is the study run from?

The Child Health and Parenting (CHAP) research group at Uppsala University (Sweden) together with the Child Pediatric Outpatient Clinic in Uppsala Region.

When is the study starting and how long is it expected to run for?

January 2024 to December 2026

Who is funding the study?

Swedish Research Council for Health, Working Life and Welfare  
Foundation Sunnerdahls Disability Fund (Sweden)

Who is the main contact?

Dr Karin Fångström, [karin.fangstrom@uu.se](mailto:karin.fangstrom@uu.se)

## Contact information

### Type(s)

Scientific, Principal investigator

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Public

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## Additional identifiers

### Clinical Trials Information System (CTIS)

Nil known

### ClinicalTrials.gov (NCT)

Nil known

### Protocol serial number

Nil known

## Study information

### Scientific Title

The clinical- and cost-effectiveness of a parenting support group intervention for parents of preschool children with subclinical neurodevelopmental disorders and mental health problems

### Study objectives

Children's behavioral problems, reported by parents, will decrease as an effect of the intervention, that parent's self-efficacy will increase and their stress will decrease

### Ethics approval required

Ethics approval required

### Ethics approval(s)

approved 05/06/2024, The Swedish Ethical Review Authority (Box 2110, Uppsala, 750 02, Sweden; +46 10-475 08 00; [registrator@etikprovning.se](mailto:registrator@etikprovning.se)), ref: 2024-03064-01

### Study design

Multiple-baseline single-case experimental design (SCED) across participants with pre- and post-measures and a 3-month follow-up

### Primary study design

Interventional

### Study type(s)

Treatment

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

Subclinical neurodevelopmental disorders and mental health problems

### Interventions

Current interventions as of 08/11/2024:

The parenting support group intervention "Everyday life and parenting" consist of four modules. The intervention is theoretically based in social-learning theory and it uses methods from applied behavior analysis, e.g., functional behavior assessment, as well as structured teaching. Three of the intervention modules includes key components from the most effective parenting programs targeting behavior problems in children. Parenting techniques taught during the intervention includes positive parenting skills such as positive reinforcement techniques, relationship enhancement techniques and reframing unhelpful cognitive perceptions about the child. The fourth intervention module is based on theories and knowledge from treatment provided to children with autism and/or intellectual disabilities as well as the ICF framework. Important elements in this module are teaching parents how to organize the physical environment, create activity schedules, visual schedules and/or cues, establish routines with flexibility, and use visually structured activities. These elements create clarity and predictability in everyday life. Furthermore, the module includes teaching parents general analytical and learning principles for teaching new skills or dealing with problem behaviors. In addition, parents are thought how to adapt activities and demands on the individual child based on his/her cognitive and adaptive abilities.

The delivery format of our intervention is also based on well researched components and includes providing psychoeducation; discussions; homework assignments and peer support. Each module last 3 hours and the intervention is given every other week for 8 weeks.

Previous interventions:

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The delivery format of our intervention is also based on well researched components and includes providing psychoeducation; discussions; homework assignments and peer support. Each module last 3 hours and the intervention is given every other week for 8 weeks.

A computer-generated randomization sequence will be used to assign the participants to a 4-, 7- or 10-week baseline in a 1:1:1 ratio. Block randomization of block sizes 3 or 6 will be generated in a computerized randomization scheme.

### **Intervention Type**

Behavioural

### **Primary outcome(s)**

Current primary outcome measure as of 08/11/2024:

Questionnaire distributed to parents every week for 15, 18 or 21 weeks, i.e. during baseline phase, intervention and four weeks post intervention:

1. Child behavior problems, questions based on the Eyberg Child Behavior Inventory
2. Parental self-efficacy, the Me As a Parent questionnaire
3. Parents' perceived stress, the Parental Stress Scale

Previous primary outcome measure:

Questionnaire distributed to parents every week for 16, 19 or 22 weeks, i.e. during baseline phase, intervention and four weeks post intervention:

1. Child behavior problems, questions based on the Eyberg Child Behavior Inventory
2. Parental self-efficacy, the Me As a Parent questionnaire
3. Parents' perceived stress, the Parental Stress Scale

### **Key secondary outcome(s)**

Current secondary outcome measure as of 08/11/2024:

Measured at pre-, post and 3-month follow-up, parent's ratings:

1. Child mental health problems, using Strength and Difficulties Questionnaire
2. Parental self-efficacy, using Me As a Parent
3. Parental stress using the Parental Stress Scale
4. Parent's mental health using General health questionnaire, GHQ-12
5. Quality of Life, using Assessing Quality of Life 8 Dimensions (AQOL-8D)
6. Costs related to consumption of societal resources, Trimbos/iMTA questionnaire (TIC-P)
7. Costs related to healthcare consumption using data from child medical records on the number of healthcare visits, prescribed drugs, referrals etc.

Measured weekly during 15, 18 or 21 weeks:

8. Children's positive and negative experiences of family life using 5 questions developed for this study

Previous secondary outcome measure:

Measured at pre-, post and 3-month follow-up, parent's ratings:

1. Child mental health problems, using Strength and Difficulties Questionnaire
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Measured weekly during 16, 19 or 22 weeks:

8. Children's positive and negative experiences of family life using 5 questions developed for this study

### **Completion date**

31/12/2026

## **Eligibility**

### **Key inclusion criteria**

1. Parents of children aged 2 - 6 years old with subclinical levels of NDD and behavioral problems (as assessed by a clinical psychologist) who have been offered and accepted to participate in the parenting intervention Everyday Life and Parenting
2. Parents need to be able to answer questionnaires in Swedish
3. Only one parent from each family will be included (i.e., the father as fathers often are underrepresented and therefore will contribute to a gender balance in the sample)
4. Children aged 4 - 6 years old, who's parent/parents participate in Everyday life and parenting

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Mixed

**Lower age limit**

4 years

**Upper age limit**

99 years

**Sex**

All

**Key exclusion criteria**

1. Parents who cannot answer the questionnaires in Swedish
2. Parents who don't understand the study information or informed consent
3. Children whose parents are excluded based on the above criteria
4. Children below 4 years of age

**Date of first enrolment**

16/08/2024

**Date of final enrolment**

31/12/2025

**Locations****Countries of recruitment**

Sweden

**Study participating centre**

**Child Pediatric Outpatient Clinic in Uppsala Region**

Islandsgatan 2

Uppsala

Sweden

75308

# Sponsor information

## Organisation

Uppsala University

## ROR

<https://ror.org/048a87296>

## Organisation

Akademiska Children's Hospital/Region Uppsala

# Funder(s)

## Funder type

Research council

## Funder Name

Forskningsrådet om Hälsa, Arbetsliv och Välfärd

## Alternative Name(s)

Swedish Research Council for Health, Working Life and Welfare, Forskningsrådet om Hälsa, Arbetsliv och Välfärd, FORTE

## Funding Body Type

Government organisation

## Funding Body Subtype

Local government

## Location

Sweden

## Funder Name

Foundation Sunnerdahls Disability Fund

# 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 Karin Fångström (karin.fangstrom@uu.se).

### **IPD sharing plan summary**

Available on request