Norm data for the Infant Motor Profile (IMP) and the Standardized Infant NeuroDevelopmental Assessment (SINDA)

No registrations found.

Ethical review	Not applicable
Status	Pending
Health condition type	-
Study type	-

Summary

ID

NL-OMON22603

Source NTR

Brief title

Health condition

The project studies the general population to create norms. IMP and SINDA are used to detect at early age children with developmental disorders such as cerebral palsy. Het huidige onderzoek betreft de verzameling van normgegevens in de algemene populatie. IMP en SINDA richten zich op de vroegdectectie van ontwikkelingsstoornissen zoals cerebrale parese ('spasticiteit')

Sponsors and support

Primary sponsor: University Medical Center Groningen **Source(s) of monetary or material Support:** Cornelia Stichting Stichting Ontwikkelingsneurofysiologie Groningen

Intervention

Outcome measures

Primary outcome

Infant Motor Profile Total Score

Secondary outcome

Infant Motor Profile Domain scores, i.e., score of Variation, Adaptatility, Symmetry, Fluency and Performance

SINDA neuromotor score

SINDA developmental score

Study description

Background summary

Recently two neurodevelopmental assessments have been developed: the Infant Motor Profile (IMP) and the Standardized Infant NeuroDevelopmental Assessment (SINDA). The IMP assesses infant motor behaviour and assists in high risk infants tailor-made physical therapy guidance and prediction of developmental outcome. SINDA assists general paediatricians to detect at early age infants at risk for neurodevelopmental disorders.

Objective: To obtain norm data for IMP and SINDA

Study design: Cross-sectional study

Study population: Representative sample of the general population of infants, 2–18 months (n=1700; 100 per month)

Main study parameters/endpoints: IMP scores and SINDA scores

Study objective

descriptive study: data collection for norms of two novel infant assessments, i.e., the Infant Motor Profile (IMP) and the Standardized Infant NeuroDevelopmental Assessment (SINDA)

Study design

Infants are assessed once.

Intervention

none

Contacts

Public

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Eligibility criteria

Inclusion criteria

The child lives in one of the three northern provinces of The Netherlands

Age at inclusion between 2 and 18 months

Parents or legal representatives will provide written informed consent

Exclusion criteria

Severe illness precluding the assessment, such as complex congenital heart disorders with insufficient oxygen saturation

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Parents have insufficient understanding of the Dutch or English language to be able to give informed consent

Study design

Design

Intervention model: Other Control: N/A , unknown

Recruitment

NL	
Recruitment status:	Pending
Start date (anticipated):	01-01-2017
Enrollment:	1700
Туре:	Anticipated

Ethics review

Not applicable	
Application type:	Not applicable

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register	ID
NTR-new	NL6039
NTR-old	NTR6170
Other	NL 58069.042.16 : METc 2016/294

Study results

Summary results

not yet