

Studie naar het werkgeheugen bij adolescenten met autisme in Nederland.

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In our view a proper functioning working memory (WM) is not only important for cognitive processes as language, memory etc. (i.e. 'cold' cognitive information), but is also essential for successfully navigating in the social world (i.e. 'hot'...

Ethische beoordeling	Positief advies
Status	Werving nog niet gestart
Type aandoening	-
Onderzoekstype	Observationeel onderzoek, zonder invasieve metingen

Samenvatting

ID

NL-OMON23771

Bron

NTR

Verkorte titel

SWAAN

Aandoening

Neuropsychology
Adolescents
High Functioning Autism
Working Memory

Neuropsychologie
Adolescenten
Hoog Functionerend Autism
Werk geheugen

Ondersteuning

Primaire sponsor: Epilepsy Centre Kempenhaeghe, Heeze

Overige ondersteuning: Epilepsy Centre Kempenhaeghe, Heeze

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

The neuropsychological WM parameters and behavioural WM parameters as obtained by the neuropsychological tasks and the questionnaires. The endpoints for the neuropsychological tasks and behavioural data as obtained by the questionnaires are standardized (z-) scores.

This all will be processed in terms of:

1. Correlation between neuropsychological test results, and behavioural questionnaires data results;

2. Differences between adolescents with HFA and normal controls.

Toelichting onderzoek

Achtergrond van het onderzoek

Autism spectrum disorder (ASD) is a heterogeneous neurodevelopmental syndrome. The aetiology of ASD is still unknown, and in spite of many studies the main cognitive theories of ASD - Theory of Mind deficit hypotheses, Weak Central Coherence account, and the Executive dysfunction theory - fail to explain the broad spectrum of symptoms found in ASD.

In the Netherlands a growing number of high average intelligent (HAVO/VWO) adolescents with autism is in the need for special secondary education due to their social, executive function (EF) and specific learning problems. Although these adolescents experience many problems in their everyday life, it is very difficult to objectify these problems in clinical or laboratory settings due to the compensation strategies these adolescents apply in these structured settings. This makes the need for finding new ways to objectify their problems and to target (educational) intervention specified to their needs, especially high.

Although there have been many studies that looked at working memory (WM) (a core domain of EF) in adolescents with high functioning autism (HFA), the results remain inconclusive. In these studies it is assumed that WM only processes abstract information. However our hypothesis is that WM plays a more central role in processing both 'cold' and 'hot' cognitive information. WM capacity problems therefore may play a leading role in the multiple symptoms displayed by adolescents with HFA.

This study will be a case-control study.

Study population: Adolescents with HFA aged 12-16 years, and a normal control group. The study will be performed at Epilepsy Centre Kempenhaeghe Heeze, Special

Education School de Berkenschutse Heeze and at the Radboud University Nijmegen.

Doel van het onderzoek

In our view a proper functioning working memory (WM) is not only important for cognitive processes as language, memory etc. (i.e. 'cold' cognitive information), but is also essential for successfully navigating in the social world (i.e. 'hot' cognitive information).

Therefore our hypothesis is that WM plays a more central role in remembering, uploading and selecting social cues from the environment, hence in processing both 'cold' and 'hot' cognitive information. WM capacity problems therefore may play a leading role in the multiple symptoms (both 'cold' and 'hot' cognitive) displayed by intelligent adolescents with high functioning autism (HFA).

Onderzoeksopzet

Adolescent, one of their parent(s)/ caregiver(s), and their teacher/mentor are asked to fill in at home two questionnaires. The neuropsychological assessment in the HFA-group will proximally take 4 hours. The neuropsychological assessment in the normal control group will proximally take 3 hours. One appointment will be made for the neuropsychological assessment.

Onderzoeksproduct en/of interventie

This study will be a case-control study.

Contactpersonen

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Wetenschappelijk

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

For adolescents with HFA:

1. Age of 12 to 16 years;
2. Autism disorder or Asperger disorder diagnosis made conform the diagnosis criteria as formulated in the DSM-IV;
3. A signed informed consent (IC) from the adolescent and the parent/ caregiver;
4. A high average intelligence conform a HAVO/VWO education level.

For normal control adolescents:

1. Age of 12 to 16 years;
2. A signed IC from the adolescent and the parent/ caregiver;
3. A high average intelligence conform a HAVO/VWO education level.

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

For adolescents with HFA:

1. A diagnosis for other (co-morbid) psychological disorders or psychiatric diseases as formulated in the DSM-IV, such as: Attention-deficit and disruptive behaviour disorders, anxiety disorders and mood disorders;
2. Appearance of additional variables that can influence cognitive functioning such as pathology of the Central Nervous System (CNS), significant hearing impairment, and medicinal treatment for epilepsy;

3. Use of Methylphenidate. In consultation with the responsible doctor, parent/caregiver and adolescent are asked if they are willing to not take the medicine on the day of the neuropsychological assessment. If all agree, the adolescent may participate in the study;
4. Inability to speak/understand the Dutch language;
5. Vision less than +4.5D or - 4.5D.

For normal control adolescents:

1. A diagnosis for a psychological disorder or psychiatric disease as formulated in the DSM-IV, such as: Pervasive developmental disorders, attention-deficit and disruptive behaviour disorders, separation anxiety disorders, selective mutism, reactive attachment disorder of infancy or early childhood, anxiety disorders and mood disorders;
2. Appearance of additional variables that can influence cognitive functioning such as pathology of the CNS, significant hearing impairment, and medicinal treatment for epilepsy;
3. Use of Methylphenidate. In consultation with the responsible doctor, parent/caregiver and adolescent are asked if they are willing not to take the medicine on the day of the neuropsychological assessment. If all agree, the adolescent may participate in the study;
4. Inability to speak/understand the Dutch language;
5. Vision less than +4.5D or - 4.5D.

Onderzoeksopzet

Opzet

Type:	Observationeel onderzoek, zonder invasieve metingen
Onderzoeksmodel:	Parallel
Toewijzing:	Niet-gerandomiseerd
Blinding:	Open / niet geblindeerd
Controle:	N.v.t. / onbekend

Deelname

Nederland	
Status:	Werving nog niet gestart

(Verwachte) startdatum: 01-11-2010
Aantal proefpersonen: 108
Type: Verwachte startdatum

Ethische beoordeling

Positief advies
Datum: 16-09-2010
Soort: Eerste indiening

Registraties

Opgevolgd door onderstaande (mogelijk meer actuele) registratie

Geen registraties gevonden.

Andere (mogelijk minder actuele) registraties in dit register

Geen registraties gevonden.

In overige registers

Register	ID
NTR-new	NL2411
NTR-old	NTR2519
Ander register	METC / CCMO : 10-3-055 / NL-32788.068.10
ISRCTN	ISRCTN wordt niet meer aangevraagd.

Resultaten

Samenvatting resultaten

N/A