New ultrasound techniques in the detection of testicular adrenal rest tumours in children with congenital adrenal hyperplasia

Published: 05-04-2011 Last updated: 06-05-2024

To study the incidence of TART in children with congenital adrenal hyperplasia with new imaging techniques such as high frequency ultrasound and elastography

Ethical review Approved WMO **Status** Recruiting

Health condition type Endocrine disorders of gonadal function

Study type Observational non invasive

Summary

ID

NL-OMON34364

Source

ToetsingOnline

Brief title

Ultrasound in the detection of testicular adrenal rest tumours (TART)

Condition

Endocrine disorders of gonadal function

Synonym

testicualr adrenal rest tumors

Research involving

Human

Sponsors and support

Primary sponsor: Universitair Medisch Centrum Sint Radboud

Source(s) of monetary or material Support: Ministerie van OC&W

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Intervention

Keyword: congenital adrenal hyperplasia, Elastography, High frequency ultrasound, testicular adrenal rest tumours

Outcome measures

Primary outcome

the incidence of TART in children

Secondary outcome

no

Study description

Background summary

In adult male patients with congenital adrenal hyperplasia benign testicular adrenal rest tumors (TART) is a common cause of infertility. The prevalence is high up to 94%. It is thought that the tumors are already present in childhood or even in utero. However, the presence of TART in children is not described in detail in young children.

Study objective

To study the incidence of TART in children with congenital adrenal hyperplasia with new imaging techniques such as high frequency ultrasound and elastography

Study design

cross sectional with a healthy control group

Study burden and risks

Ultrasound is non invasive so there are no risks for the patient.

Contacts

Public

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Scientific

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Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adolescents (12-15 years) Adolescents (16-17 years) Children (2-11 years)

Inclusion criteria

All male children (0 - 18 years) with congenital adrenal hyperplasia; Control group: male children without a condition of adrenal or gonadal pathology

Exclusion criteria

geen

Study design

Design

Study type: Observational non invasive

Intervention model: Other

Allocation: Non-randomized controlled trial

Masking: Open (masking not used)

Primary purpose: Diagnostic

Recruitment

NL

Recruitment status: Recruiting
Start date (anticipated): 01-05-2011

Enrollment: 35

Type: Actual

Ethics review

Approved WMO

Date: 05-04-2011

Application type: First submission

Review commission: CMO regio Arnhem-Nijmegen (Nijmegen)

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 CCMO

ID

NL32839.091.10