The Effect of Thiazide, Amiloride and Hypertonic Saline on Urinary Biomarkers in Healthy Subjects (THAM)

This study has been completed.

Sponsor:
Regional Hospital Holstebro

Information provided by (Responsible Party):
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ClinicalTrials.gov Identifier:
NCT01635231

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Last verified: March 2014
History of Changes

Purpose

Urinary biomarkers (u-NKCC2, u-ENaC-gamma and u-AQP2) reflects the activity of the sodium- and water channels in the human kidney. Changes in the sodium-and water channel activity can be induced by blocking the sodium channels with diuretics in healthy subjects

Condition	Intervention
Nephropathy	Other: hypertonic saline

Study Type: Interventional
Study Design: Allocation: Randomized
Intervention Model: Crossover Assignment
Masking: Double Blind (Subject, Caregiver, Investigator)
Primary Purpose: Diagnostic

Official Title: **Effect** of **Thiazide**, **Amiloride** and Hypertonic Saline on Sodium- and Water Channel Activity in the Nephron in Healthy Subjects Estimated by Urinary Biomarkers

Resource links provided by NLM:

MedlinePlus related topics: Drinking Water

Drug Information available for: Amiloride Amiloride hydrochloride

U.S. FDA Resources

Further study details as provided by Regional Hospital Holstebro:

Primary Outcome Measures:

- urinary biomarkers [Time Frame: baseline, 0 hours and 1.5 hours after intervention] [Designated as safety issue: No]
  Urinary excretion of epithelial sodium channels (ENaC), Sodium-potassium-2chloride transporters (NKCC2) and aquaporin2 channels (AQP2) before, during and after fluid infusion

Secondary Outcome Measures:

- vaso active hormones [Time Frame: baseline, 0 hours and 1.5 hours after intervention] [Designated as safety issue: No]
  plasma concentrations of: renin, Angiotensin II, aldosterone, Vasopressin, ANP and BNP
- central blood pressure [Time Frame: baseline, 0 hours and 1.5 hours after intervention] [Designated as safety issue: No]
  measured by applanation-tonometry
- intracellular (ICV)- and extracellular volume (ECV) [Time Frame: baseline, 0 hours and 1.5 hours after intervention] [Designated as safety issue: No]