Acetylsalicylic acid as an adjuvant therapy for schizophrenia.

No registrations found.

Ethical review Positive opinion

Status Recruitment stopped

Health condition type -

Study type Interventional

Summary

ID

NL-OMON22886

Source

NTR

Brief title

Aspirine Trial

Intervention

Outcome measures

Primary outcome

3-month change in positive and negative symptoms on the total PANSS score.

Secondary outcome

3-month change in the PANSS subscales, cognitive symptoms, immunological parameters (ginterferon, IL-4, IL-6 and IL-12).

Study description

Background summary

N/A

Study objective

Findings from both epidemiological and basic research point to the possibility that NSAIDS impede the deterioration in schizophrenia.

Intervention

Acetylsalicylacid 1000mg vs Placebo for 3 months (All daily pantoprazol 40 mg).

Contacts

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Eligibility criteria

Inclusion criteria

- 1. Schizophrenia
- 2. Schizo-affective disorder
- 3. Schizofreniform disorder according to DSM-IV (for max 10 years)
- 4. Age 18 55 years

5. Stabile, min. 60 on PANSS, min 2x a score of min 4 on PANSS.

Exclusion criteria

- 1. No contraindication for acetylsalicylic acid
- 2. No hypersensitivity to acetylsalicylic acid or pantoprazole
- 3. No significant somatic illness
- 4. No chronic use of a nonsteroidal anti-inflammatory drug (NSAID)
- 5. No use of corticosteroids
- 5. Not pregnant
- 6. No drugs dependency
- 7. Informed consent obtained.

Study design

Design

Study type: Interventional

Intervention model: Parallel

Masking: Double blinded (masking used)

Control: Placebo

Recruitment

NI

Recruitment status: Recruitment stopped

Start date (anticipated): 01-01-2004

Enrollment: 80

Type: Actual

Ethics review

Positive opinion

Date: 22-12-2004

Application type: First submission

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

RegisterIDNTR-newNL5NTR-oldNTR29Other: N/A

ISRCTN ISRCTN27745631

Study results

Summary results

N/A