Dyspepsia - Dopamine.

No registrations found.

Ethical review	Not applicable
Status	Suspended
Health condition type	-
Study type	Interventional

Summary

ID

NL-OMON22541

Source
Nationaal Trial Register

Brief title
N/A

Intervention

Study description

Background summary

Male patients with FD will undergo a [123I]IBZM SPECT scan to assess the number of striatal dopaminergic D2 receptors. Two hours later a brainscan (lasting 50 minutes) will be made.

Study objective

N/A

Intervention

Single dose of 185 MBq [123I]IBZM intravenously. As controlgroup an existing file of healthy male volunteers will be used.

Contacts

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Eligibility criteria

Inclusion criteria

1. Nepean Dyspepsia Index (NDI) score >= 25;

2. >18 years of age.

Exclusion criteria

1. Use of drugs that affect gastrointestinal motility (antidepressants, opiates, NSAID’s, calcium-blockers, nitrates);

2. use of drugs that can affect binding of the radioligand to dopamine D2 receptors (neuroleptics, dopamine D2 agonists, anti-vertigo-drugs metoclopramide, alizapride);

3. IBS;

4. previous abdominal surgery;

5. history of myocardial infarct or CVA;
6. epilepsy;

7. severe liver or renal insufficience;
8. depression;

9. claustrophobia.

Study design

Design

Study type:	Interventional
Intervention model:	Other

Control: N/A , unknown

Recruitment

NL

Recruitment status:	Suspended
Start date (anticipated):	01-08-2004
Enrollment:	15
Type:	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	NL28
NTR-old	NTR49
Other	: N/A
ISRCTN	ISRCTN wordt niet meer aangevraagd

Study results

Summary results
N/A