Endoscopic sclerotherapy compared with no specific treatment for the primary prevention of bleeding from oesophageal varices: a randomised controlled multicentre trial

Submission date	Recruitment status No longer recruiting	Prospectively registered		
06/08/2003		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
13/08/2003	Completed	[X] Results		
Last Edited	Condition category	[] Individual participant data		
18/07/2008	Digestive System			

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number 90/006

Study information

Scientific Title

Study objectives

To compare endoscopic sclerotherapy (ES) with no specific treatment for the primary prevention of bleeding from oesophageal varices in patients with cirrhosis.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Liver cirrhosis

Interventions

Prophylactic sclerotherapy versus no specific treatment.

For more information, please contact the College voor Zorgverzekeringen (Health Care Insurance Board) on infoSBA@cvz.nl.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s))

Not provided at time of registration

Completion date

31/05/1993

Eligibility

Key inclusion criteria

- 1. Adults with endoscopically documented grade II, III or IV oesophageal varices
- 2. Absence of prior bleeding from varices
- 3. Evidence of active and/or progressive liver disease (e.g. as indicated by repeatedly elevated serum transaminases or increasing serum bilirubin levels or development of ascites within the past year)
- 4. Documented increase in the size of oesophageal varices
- 5. All patients gave informed consent

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Αll

Key exclusion criteria

- 1. Use of beta-blockers and nitrates
- 2. Malignancy
- 3. Haemophilia
- 4. Age older than 70 75 years
- 5. No opportunity for follow-up visits

Date of first enrolment

01/05/1986

Date of final enrolment

31/05/1993

Locations

Countries of recruitment

Netherlands

Study participating centre

Erasmus MC

Rotterdam Netherlands 2040 CA

Sponsor information

Organisation

Dutch Health Care Insurance Board (College voor Zorgverzekeringen) (The Netherlands)

Funder(s)

Funder type

Government

Funder Name

Dutch Health Care Insurance Board (College Voor Zorgverzekeringen) (The Netherlands) (ref: OG 90/006)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results	15/08/2003		Yes	No