Duodenum-preserving head resection versus pancreatico-duodenectomy for chronic pancreatitis of the head

Submission date Recruitment status [X] Prospectively registered 21/04/2009 No longer recruiting [X] Protocol Statistical analysis plan Registration date Overall study status 30/04/2009 Completed [X] Results Individual participant data **Last Edited** Condition category 28/08/2018 Digestive System

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

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Additional identifiers

Protocol serial number

SDGC 01/2009

Study information

Scientific Title

Duodenum-preserving head resection versus pancreatico-duodenectomy for chronic pancreatitis of the head: a randomised controlled multicentre trial

Acronym

ChroPac

Study objectives

The two-sided null-hypothesis states that both surgical interventions lead to the same average quality of life (QoL) scores during 24 months after surgery. The two-sided alternative-hypothesis states that the two interventions perform differently in terms of the primary efficacy endpoint.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics Committee of the University of Heidelberg, approved on 20/04/2009 (ref: S-029/2009)

Study design

Prospective randomised controlled observer and patient blinded multicentre surgical trial with two parallel study groups

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Chronic pancreatitis

Interventions

Experimental intervention:

Any surgical technique that removes inflamed pancreatic tissue of the head without resection of the duodenum (e.g. Beger, Frey or Berne procedure).

Control intervention:

Pylorus preserving/classic Whipple procedure.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Average QoL during 24 months after surgery, measured 6, 12 and 24 months after surgery by the European Organisation for Research and Treatment of Cancer Quality of Life Questionnaire for Cancer patients (EORTC QLQ-C30) scale "physical functioning". The primary efficacy analysis will be conducted for the intention-to-treat population. An analysis of covariance (ANCOVA) model will be applied for the intervention group comparison adjusting for age, centre and EORTC QLQ-C30 scale "physical functioning" before surgery. The level of significance is set at 5% (two-sided) and sample size is determined to assure a power of 1-beta = 90%.

Key secondary outcome(s))

Exploratory analyses

Completion date

30/06/2013

Eligibility

Key inclusion criteria

- 1. Patients (no age or gender restrictions) with chronic pancreatitis of the head and pain, eligible for elective surgical resection
- 2. Ability of subject to understand character and individual consequences of the clinical trial
- 3. Written informed consent

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Other

Sex

All

Key exclusion criteria

Participation in another intervention-trial with interference of intervention and outcome of this study

Date of first enrolment

18/05/2009

Date of final enrolment

30/06/2013

Locations

Countries of recruitment

United Kingdom

Germany

Netherlands

Slovenia

Study participating centre

Im Neuenheimer Feld 110

Heidelberg Germany 69120

Sponsor information

Organisation

Study Centre of the German Surgical Society (SDGC) (Germany)

ROR

https://ror.org/00ew91p29

Funder(s)

Funder type

Research council

Funder Name

German Research Council (Deutsche Forschungsgemeinschaft [DFG]) (Germany) (ref: ChroPac SE 1682/2-1)

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	09/09/2017	Yes	No
<u>Protocol article</u>	protocol	29/04/2010	Yes	No
Participant information sheet	Participant information sheet	11/11/2025 11/11/2025	No	Yes