

G-COACH: Feasibility of the geriatric co-management for cardiology patients in the hospital programme

Submission date 21/05/2020	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 23/05/2020	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 26/10/2022	Condition category Circulatory System	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Older patients admitted to an cardiac care unit often experience functional decline and sustained disability. Geriatric co-management has demonstrated significant improvements on outcomes in older patients with a hip fracture, but has never been evaluated in the cardiac care setting.

This study introduced a new geriatric co-management programme on the cardiac care units of the University Hospitals Leuven. The aim was to evaluate whether this programme was acceptable and feasible to perform.

Who can participate?

Patients aged 75 years or older admitted for acute cardiovascular disease or transcatheter aortic valve implantation to the cardiac care units of the University Hospitals Leuven and healthcare professionals working on the acute cardiac care units of the University Hospitals Leuven.

What does the study involve?

On admission to the hospital, a comprehensive geriatric assessment was performed, care needs were determined, and an interdisciplinary care plan was drafted. A nurse from the geriatrics department provided daily follow-up until hospital discharge.

What are the possible benefits and risks of participating?

The programme was developed to benefit patients, including a better functional status and less complications. There are no direct risks.

Where is the study run from?

University Hospitals Leuven (Belgium)

When is the study starting and how long is it expected to run for?

January 2015 to May 2018

Who is funding the study?
Onderzoeksraad, KU Leuven (Research Council, KU Leuven), Belgium

Who is the main contact?
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Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

Protocol serial number

REF 22/15/028

Study information

Scientific Title

Implementation of the Geriatric CO-mAnagement for Cardiology patients in the Hospital (G-COACH) programme: a feasibility study

Acronym

G-COACH

Study objectives

Is a geriatric co-management programme implemented on the acute cardiac care units of the University Hospitals Leuven considered acceptable and feasible to perform?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 03/08/2017, The Medical Ethics Committee of the University Hospitals Leuven (Ethische commissie onderzoek UZ Leuven, Herestraat 49, 3000 Leuven, Belgium; +32 16 34 86 00; ec@uzleuven.be), ref: s59543

Study design

One-group experimental single centre feasibility study

Primary study design

Interventional

Study type(s)

Other

Health condition(s) or problem(s) studied

Older patients with geriatric profile admitted to cardiac care units

Interventions

Nurse-led geriatric co-management programme, including a comprehensive geriatric assessment on hospital admission, risk stratification, daily follow-up by geriatrics team until hospital discharge, implementation of standardised and evidence-based protocols for geriatric syndromes, early rehabilitation, individual exercise programme, and early discharge planning.

Informed consent is obtained within three days of admission to the cardiac care unit (15 minutes). A nurse enrolls the patient in the geriatric co-management programme (30 minutes). A researcher performs an interview and asks about the experiences with the programme (15 minutes). A researcher observes daily health status and whether there are any complications (10 minutes) until discharge from the cardiac care unit. The participation ends when discharged from the hospital.

Intervention Type

Other

Primary outcome(s)

Experiences with the programme, including self-perceived acceptability and self-perceived feasibility, measured using structured interviews with questionnaires during hospitalisation.

Key secondary outcome(s)

Measured during hospitalisation on the cardiac care unit. This is case dependent (some patients have a long length of stay, others short) and there is no one fixed timepoint:

1. Reach of the programme, measured using registrations in the electronic patient record
2. Fidelity, measured using registrations in the electronic patient record
3. Dose, measured using registrations in the electronic patient record
4. Determinants for implementation, measured using a questionnaire and interviews

Completion date

31/05/2018

Eligibility

Key inclusion criteria

1. Patients aged 75 years or older admitted for acute cardiovascular disease or transcatheter aortic valve implantation to the cardiac care units of the University Hospitals Leuven
2. Healthcare professionals working on the acute cardiac care units of the University Hospitals Leuven

Participant type(s)

Mixed

Healthy volunteers allowed

No

Age group

Mixed

Sex

All

Total final enrolment

199

Key exclusion criteria

1. Length of stay < 3 days
2. No baseline assessment (included transfers from other units or hospitals)
3. Unable to complete assessment
4. No inform consent
5. Receiving palliative care

Date of first enrolment

07/11/2017

Date of final enrolment

02/05/2018

Locations**Countries of recruitment**

Belgium

Study participating centre

University Hospitals Leuven

Herestraat 49

Leuven

Belgium

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Sponsor information**Organisation**

KU Leuven

ROR

<https://ror.org/05f950310>

Funder(s)**Funder type**

Research council

Funder Name

Onderzoeksraad, KU Leuven

Alternative Name(s)

Research Council, KU Leuven

Funding Body Type

Private sector organisation

Funding Body Subtype

Universities (academic only)

Location

Belgium

Results and Publications

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are/will be available upon request from Koen Milisen (koen.milisen@kuleuven.be). Each request will be considered on a case basis. Reasonable requests include use of baseline data for epidemiological purposes.

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article		02/05/2022	26/10/2022	Yes	No
Protocol article	protocol	01/10/2018	22/05/2020	Yes	No
Study website	Study website	11/11/2025	11/11/2025	No	Yes