

Hospital at home – a study to reduce rehospitalizations

Submission date 28/01/2023	Recruitment status Recruiting	<input checked="" type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 04/04/2023	Overall study status Ongoing	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 23/03/2026	Condition category Other	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

The aim of the Hospital@Home study is to reduce the rehospitalization rate in in-hospital patients on a medical ward at high risk of rehospitalization. The intervention consists of a targeted multidisciplinary intervention over 5 days after hospital discharge and aims to ensure an optimal transition from hospital to ambulatory care. The primary objective is to reduce the rate of unplanned rehospitalisation in the aforementioned patient group by 25%.

Who can participate?

Hospital inpatients at high risk of rehospitalization scheduled for discharge to their home

What does the study involve?

In addition to the standard-of-care counseling prior to hospital discharge, patients in the intervention group will be contacted by telephone daily for 5 days, starting on the day after hospital discharge. They will be asked about their general well-being, need for support in their daily care, and their adherence to prescribed medications. If a home visit is deemed necessary before discharge or on one of the daily phone calls, the visit will be carried out by a member of the study team. Members of the Hospital@Home study team will coordinate the discharge management prior to discharge, such as ensuring that all the necessary paperwork is present, organizing and coordinating post-discharge outpatient care, organizing any necessary material and medication and follow-up appointments with primary care physicians.

What are the possible benefits and risks of participating?

Participants could potentially receive a direct benefit from participation if they receive additional care from the Hospital@Home team. The goal of the intervention is to improve patient care in the home environment by coordinating the various inpatient and outpatient services involved, thus reducing unplanned readmissions and increasing patient satisfaction with the discharge procedure. No risks are expected.

Where is the study run from?

Cantonal Hospital of Baden, Canton Aargau (Switzerland)

When is the study starting and how long is it expected to run for?
November 2021 to January 2026

Who is funding the study?
1. Department of Health, Canton of Aargau (Switzerland)
2. Stiftung Kardio (Switzerland)

Who is the main contact?
Prof. Dr. med. M. Wertli, maria.wertli@ksb.ch

Contact information

Type(s)
Principal investigator

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Additional identifiers

Clinical Trials Information System (CTIS)
Nil known

Protocol serial number
SNCTP000005155, BASEC2022-01040

Study information

Scientific Title
Hospital@Home: Improving discharge management and reducing the risk of rehospitalizations in multimorbid Patients

Acronym
H@H

Study objectives
Current study hypothesis as of 29/10/2024:

We hypothesize that through a targeted multidisciplinary intervention, the risk of rehospitalization within 30 days in high-risk patients can be reduced by at least 25%. Such a reduction would result in substantial individual and societal benefits.

Previous study hypothesis:

It is hypothesized that through a targeted multidisciplinary intervention over 5 days after hospital discharge, the risk of rehospitalization within 30 days in high-risk patients can be reduced by at least 25%.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 30/09/2022, Ethikkommission Nordwest- und Zentralschweiz EKNZ (Hebelstrasse, 53 4056, Basel, Switzerland; +41 (0)61 268 13 50; eknz@bs.ch); ref: 2022-01040

Study design

Pragmatic single-center randomized open-label superiority trial

Primary study design

Interventional

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Risk for unplanned rehospitalizations in multimorbid patients discharged from the hospital

Interventions

Current interventions as of 29/10/2024:

The multimodal intervention includes several transitional care components per discharge phase: pre-discharge, bridging, and post-discharge interventions.

Pre-discharge interventions

Patients will receive individualized discharge management by an APN of the Hospital@Home team in addition to standard-of-care counseling. For each patient, the APN will instruct self-management, conduct medication reconciliation and review, assess the needs for post-discharge care coordination, improve the discharge summary and care plan, and involve the family as needed.

Bridging interventions

Bridging interventions such as coordination and planning of outpatient follow-up appointments with primary care physicians or community nurses and availability of material and medication upon discharge (coordination with pharmacies) are used as needed. In addition to patient education, communication with the outpatient healthcare team (primary care physicians, specialists, and community nurses) and family members ensures that patients are aware of appointments and changes to their care plan. A scheduled home visit can also serve as a bridging intervention to directly hand over patients to the community nurses (e.g., for patients with intravenous therapy or supply and/or drainage systems).

Post-discharge intervention

Patients will receive structured telephone follow-up daily for 5 days (weekdays only) following

the discharge. The telephone calls will focus on the following aspects:

- Assessment of symptoms and vital signs, general well-being, and organizational issues;
- Potentially adjustments to medications based on the treatment plan;
- Medication adherence, structured needs assessment to organize missing medications or material (e.g. wound dressings), identification of the need for home visits and follow-up visits;
- Counseling on health-related issues, patient and family education;
- Organizing and coordinating additional follow-up visits with primary care physicians and specialists;
- Organizing and coordinating care with community nurses (if necessary)

During the first 5-days post-discharge, the Hospital@Home team may conduct home visits, if needed. Home visits are performed by APN and/or a physician and are aimed at avoiding the need for rehospitalization by early treatment adjustments. Patients in the intervention group will be able to call a hotline to contact the Hospital@Home team during workdays to ask questions or to receive help.

Control procedures

Patients assigned to the control arm will receive the current standard-of-care counselling before hospital discharge. The nurse care manager team will provide a summary of discharge recommendations and organize outpatient care. In each patient the responsible resident physician will review the discharge medication, conduct a medication reconciliation, and explain the updated medication

Previous interventions:

Patients with a BARRS-Score ≥ 5 will be randomised after providing written informed consent. The researchers will use blinded group allocation by randomizing patients into an intervention and a control group using the study software RedCap. A randomization list will be generated by a statistician otherwise not involved in the study. The randomization will be stratified based on the presence/absence of active cancer and/or heart insufficiency.

Patients will be randomly assigned into two parallel groups:

1. Intervention: The Hospital@Home care team will follow up patients for 5 days
2. Control group: usual discharge without follow-up

Patients in the intervention arm will – in addition to the standard-of-care counseling prior to hospital discharge – receive a multidisciplinary, coordinated 5-day care at their home, starting on the day after discharge. Members of the Hospital@Home study team will coordinate the discharge management prior to discharge, such as ensuring that all the necessary paperwork is present, organizing and coordinating post-discharge outpatient care, organizing any necessary material and medication and follow-up appointments with primary care physicians.

Intervention Type

Other

Primary outcome(s)

Current primary outcome measure as of 29/10/2024:

The primary outcome is the rate of first unplanned rehospitalizations within 30 days after discharge from the index admission. An unplanned rehospitalization is defined as an unscheduled admission to any hospital and any division within 30 days after discharge.

Previous primary outcome measure:

The rate of unplanned rehospitalizations in the high-risk group (BARRS score ≥ 5) at 30 days after discharge. Planned (elective) rehospitalizations will not be counted as events. This information is collected by a phone call at 30 days after discharge.

Key secondary outcome(s)

Current secondary outcome measures as of 29/10/2024:

1. The rate of unplanned rehospitalizations in the high-risk group (BARRS score ≥ 5) 18 days after discharge.
2. Change in quality of life between discharge and 30 days after discharge, using the EQ5D-5L
3. Quality of life after 30 days, using the EQ5D-5L
4. Death within 30 days
5. Health care use (e.g., physicians' visits, emergency department visits) within 30 days
6. Patient satisfaction with the discharge management at 30 days, using a clinic-specific questionnaire (TEA)

Previous secondary outcome measures:

1. The rate of unplanned rehospitalizations in the high-risk group (BARRS score ≥ 5) 18 days after discharge. This information is collected by a phone call 30 days after discharge
2. Quality of life measured by EQ-5D questionnaire at discharge and 30 days after discharge
3. Patient satisfaction with discharge management, as measured by the Transition Evaluation Assessment Tool (TEA) at 5 days after discharge

Completion date

31/01/2027

Eligibility

Key inclusion criteria

Current participant inclusion criteria as of 25/11/2024:

Patients ≥ 18 years old with a high risk (estimated risk $\geq 20\%$) for unplanned rehospitalization who consent to participate are included in this RCT, if they meet the following inclusion criteria:

1. Hospital inpatients with a high risk (estimated risk $\geq 20\%$) for unplanned rehospitalization (BARRS-Score of ≥ 5 points) who are scheduled for discharge to their home
2. Patient and/or proxy must be able to give written informed consent
3. Patient and/or proxy must be able to communicate in German

Previous participant inclusion criteria as of 29/10/2024 to 25/11/2024:

1. Hospital inpatients with a high risk (estimated risk $\geq 20\%$) for unplanned rehospitalization (BARRS-Score of ≥ 5 points) scheduled for discharge to their home
2. Patient must be able to give written informed consent

Previous participant inclusion criteria:

1. Hospital inpatients with a BARRS-Score of ≥ 5 points scheduled for discharge to their home
2. Patient must be able to communicate in the German language
3. Patient must be able to give written informed consent

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Mixed

Lower age limit

18 years

Upper age limit

100 years

Sex

All

Total final enrolment

0

Key exclusion criteria

Current participant exclusion criteria as of 25/11/2024:

Patients are excluded if they meet the following criteria:

1. Discharge to other institutions (e.g., rehabilitation facilities, nursing homes);
2. Patients or proxy that are not able to understand the trial (e.g., cognitive impairment, language barrier);
3. Anticipated death within 30 days of the trial period;
4. Planned hospitalization within the next 30 days;
5. Unacceptable distance for home visits (>20 km away from the hospital);
6. Prior participation in the current trial (electronic health record will be labeled).

Previous participant exclusion criteria as of 29/10/2024 to 25/11/2024:

1. Discharge to other institutions (e.g., rehabilitation facilities, nursing homes);
 2. Patients that are not able to understand the trial (e.g., cognitive impairment, language barrier);
 3. Anticipated death within 30 days of the trial period;
 4. Planned hospitalization within the next 30 days;
 5. Unacceptable distance for home visits (>20 km away from the hospital);
 6. Prior participation in the current trial (electronic health record will be labelled).
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Previous exclusion criteria:

1. Patients discharged to other institutions, including rehabilitation facilities, nursing homes
2. Inability/unwillingness to give informed consent, e.g. due to cognitive impairment or language barrier
3. Patients who are scheduled for a planned hospitalization within the next 30 days
4. Patients who live more than 20 km away from the hospital
5. Previous participation in the trial

Date of first enrolment

04/05/2023

Date of final enrolment

31/12/2026

Locations

Countries of recruitment

Switzerland

Study participating centre

CTU Cantonal Hospital of Baden, Canton Aargau

Kantonsspital Baden AG

Im Ergel 1

Baden AG

Switzerland

5404

Sponsor information

Organisation

Kantonsspital Baden

ROR

<https://ror.org/034e48p94>

Funder(s)

Funder type

Charity

Funder Name

Stiftung Kardio

Funder Name

Department of Health, Canton of Aargau

Results and Publications

Individual participant data (IPD) sharing plan

Upon completion of the study, the datasets generated analysed will be made available upon reasonable request from the clinical trial unit KSB (ctu@ksb.ch). Data that will be shared, will need to comply with legal restrictions and be fully anonymized.

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol article		20/03/2026	23/03/2026	Yes	No