

Medium-term outcomes in patients with ascites treated with the alfapump® system: the TOPMOST study

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
15/07/2019	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
27/08/2019	Completed	<input type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
04/12/2025	Signs and Symptoms	<input checked="" type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

Ascites is a build-up of fluid between the two layers of the peritoneum. This is a membrane that lines the tummy (abdomen). Ascites is caused most commonly by liver cirrhosis and by cancer. The Sequana Medical alfapump® system is designed to remove ascites from the peritoneal cavity as it forms and move the ascites to the bladder where it is eliminated by the patient through normal urination.

The alfapump® system is a battery-powered pump that is implanted subcutaneously and is connected to two catheters; one to collect ascites from the peritoneal cavity and one to deliver the ascites to the bladder. The pump battery is charged transcutaneously (through the skin) by the patient. The alfapump® system is implanted using minimally invasive surgery and the physician can monitor the amount of ascites removed and adjust the removal rate using a dedicated computer (the alfapump® programmer) provided with the alfapump® system. According to current clinical guides, the treatment of choice for refractory ascites is repeated paracentesis to evacuate fluid, associated with the administration of albumin.

In a series of Clinical Studies, the alfapump® system was shown to be an alternative to paracentesis in patients with refractory ascites due to liver cirrhosis.

This study will create an international registry of patients fitted with the alfapump® system. The main aim of the registry is to monitor the performance of the alfapump® system.

Who can participate?

All patients implanted with an alfapump® system

What does the study involve?

The TOPMOST is an international registry involving patients who are implanted with an alfapump® system according to its intended use and not participating in another clinical study in which the alfapump® is studied. Patients of both genders are eligible for this Registry. The Registry will include commercial implants of the alfapump® system implanted until 31 December 2023. It is expected this would be up to 400 patients.

Charts of all consecutive patients enrolled will be reviewed to obtain baseline data, safety and performance data on the alfapump® up to 24 month follow-up and standard of care lab results.

At select sites, additional questionnaire and health status information including nutrition, diet, quality of life, physical activity and body measurements will be taken. Also data on concomitant medication, ECOG status, Frailty Score and further standard of care lab results will be collected.

What are the possible benefits and risks of participating?

There is no direct benefit for patients from participating in the Registry. Participation in this registry gives doctors and regulatory authorities the possibility to follow how the alfapump® system is working and whether there are any problems with the system.

This is not a treatment study and there are no additional interventions or visits planned at the hospital so no additional risks are expected for patients included in the registry.

Where is the study run from?

Sequana Medical NV, Zurich, Switzerland

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

October 2018 to December 2025.

Who is funding the study?

Sequana Medical NV, Ghent, Belgium

Who is the main contact?

Mr Jeroen Capel,

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Contact information

Type(s)

Public

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Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

ClinicalTrials.gov (NCT)

NCT04326946

Protocol serial number

2018-AAR-012

Study information

Scientific Title

inTernatiOnal alfaPuMp® cOhort STudy (TOPMOST)

Acronym

TOPMOST

Study objectives

The purpose of the study is to collect information on how the implanted ALFApump® system is working and how patients are doing before and after having received the ALFApump®. The information collected will tell how much fluid is being transported with the help of the pump, and if there is any need for extra paracentesis. The study will also collect information on what patients, as well as the physician, thinks about using the system.

Ethics approval required

Old ethics approval format

Ethics approval(s)

1. Approved 06/09/2018, Swiss Ethics Committees on research involving humans (Haus der Akademien, Laupenstrasse 7, CH-3008, Bern; info@swissethics.ch; +41 31 306 93 95), ref: 2018-01173
2. Approved 16/11/2018, Ethics Committee University of Leipzig (Karl-Sudhoff-Institut für Geschichte der Medizin und der Naturwissenschaften, Käthe-Kollwitz-Straße 82, 04109 Leipzig; +49-(0)341 / 97 154 90; ethik@medizin.uni-leipzig.de), ref: 291/18-ek
3. Approved 18/06/2019, Ethics Committee Friedrich-Alexander-University of Erlangen (Erlangen-Nuremberg, Krankenhausstraße 12, 91054 Erlangen+49 9131 85-22270; ethikkommission@fau.de), ref: 201_19Bc
4. The alfaPump® system is commercially available in the European Economic Area (EEA), receiving CE marking for commercial activity under CE 616535 with Notified Body number 0086 (BSI).

Study design

Multi-centre open label observational cohort study.

Primary study design

Observational

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Refractory ascites due to liver cirrhosis or malignant ascites

Interventions

All consecutive patients treated with the alfapump® during the enrollment period are eligible for participation.

Patients are followed for a period of up to 24 months post-implant or until 30 days post explant.

The alfapump® system is implanted using minimally invasive surgery lasting 45 minutes. The procedure is typically performed under general anaesthesia but can also be performed under local anaesthesia with sedation. The pump is implanted under the skin in the right half of the abdomen via an incision of 3-4 cm. The catheters are implanted via a small open incision. The System is fully implanted, allowing the patient normal mobility and activity.

The study is a registry and no experimental interventional procedures are planned.

Patients will be attending visits according to routine care.

Intervention Type

Device

Phase

Phase IV

Drug/device/biological/vaccine name(s)

Not applicable.

Primary outcome(s)

Patient survival with a functional alfapump® system at 6 months.

(An alfapump® system is considered functional when: capable of communication, can be charged, there is an intact fluid pathway from peritoneal space to the bladder through the system, pump transported 80% of the target daily volume in the past 30 days)

Key secondary outcome(s)

1. Occurrence of (major) reportable events, adverse events, incidents and device deficiencies requiring a surgical intervention
2. Safety at 1-month post-implant for procedure-related incidents
3. Requirement for paracentesis
4. Clinical performance of the alfapump® system by monitoring ascites transport
5. Changes in occurrence of reportable events, all incidents as well as device deficiencies from up to 6 months pre-implant and post-implant.
6. Clinical impact of the alfapump® system by monitoring patient's clinical status, clinical

chemistry, haematology laboratory data, MELD-Na, and ECOG

7. Quality of life changes

8. Overall survival

Completion date

31/12/2025

Eligibility

Key inclusion criteria

1. All patients implanted with an alfapump® system as per intended use

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Total final enrolment

37

Key exclusion criteria

1. Younger than 18 years
2. Pregnancy
3. Inability to operate the Smart Charger to recharge the alfapump®
4. Participating in another study in which the alfapump® is studied

Date of first enrolment

17/10/2018

Date of final enrolment

31/12/2023

Locations

Countries of recruitment

Austria

Germany

Switzerland

Study participating centre

Inselspital
Freiburgstrasse
Bern
Switzerland
3010

Study participating centre

Universitätsklinikum Leipzig - Department für Innere Medizin Sektion Hepatologie
Liebigstraße 20
Leipzig
Germany
04103

Study participating centre

Chirurgische Universitätsklinik Erlangen
Krankenhausstr. 12
Erlangen
Germany
91054

Study participating centre

University of Vienna
Medical University of Vienna
Spitalgasse 23
Vienna
Austria
1090

Study participating centre

Universitätsklinikum Jena
Klinik für Allgemein-, Viszeral- und Gefäßchirurgie
Am Klinikum 1
Jena
Germany
07747

Sponsor information

Organisation
Sequana Medical NV

Funder(s)

Funder type

Industry

Funder Name

Sequana Medical NV

Results and Publications

Individual participant data (IPD) sharing plan

All data generated or analysed during this study will be included in the subsequent results publication

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes