Using oxygen-deprived red blood cells to treat burns and blood cancers at Haukeland University Hospital

Submission date 24/01/2025	Recruitment status No longer recruiting	[_] Pr [_] Pr
Registration date 18/03/2025	Overall study status Completed	[_] Sta [_] Re
Last Edited 26/03/2025	Condition category Circulatory System	[_] Ind [X] Re

-] Prospectively registered
-] Protocol
- Statistical analysis plan
-] Results
-] Individual participant data
- X] Record updated in last year

Plain English summary of protocol

Background and study aims

Anaemia results from either the acquired loss of red blood cells due to disease, injury, or surgery or the inability of the patient to produce their own red blood cells due inherited disorders. Substantial evidence suggests decreased quality of red blood cells from substances that accumulate during red blood cell storage. Hypoxic storage, where the oxygen content is reduced to low levels, has been indicated by data from in vitro and pre-clinical studies to improve red blood cell quality. The use of hypoxic red blood cells for blood transfusion may result in improved outcomes, both in acute and chronic transfusion settings. It could lead to improved oxygen delivery and contribute to a decrease in the required number of red blood cell transfusions and decreased risk of iron overload. Furthermore, hypoxic blood may improve recovery in burn patients.

The Hemanext ONE system can be used to manufacture and store hypoxic red blood cells in special blood bags that do not allow any oxygen into the blood. Thus, the cells are not damaged during storage. This investigation aims to study safety of hypoxic red blood cells in 2 patient groups: acute burn and haematologic malignancies.

Who can participate?

Male and female adult patients (at least 18 years) with acute burn or haematologic malignancies, who require transfusion of red blood cells.

What does the study involve?

The study involves transfusion of two units of hypoxic red blood cells manufactured with the Hemanext ONE System. Patients will be enrolled, transfused and followed-up until their subsequent transfusion or 28 days (+/- 1 day) after transfusion, whichever comes first. There are 5 scheduled visits during the study including physical examination, vital signs and blood samples.

What are the possible benefits and risks of participating?

There are no clear immediate benefits to participate in this study, but preclinical data support the hypothesis that hypoxic blood has the potential to be of benefit to patients suffering from hematological malignancies. The results of the study may eventually lead to improved transfusion treatment of patients with hematological malignancies. There are no clear disadvantages to participating in the study. No further risk is expected with the transfusion of Hemanext ONE System beyond what is already known with blood transfusion in general, including:

- Transfusion-associated graft versus host disease
- Transfusion-related acute lung injury (TRALI)
- Post transfusion purpura
- Transfusion-associated circulatory overload (TACO)
- Transfusion-transmitted infection (TTI)
- Allergic reaction
- Transfusion-associated dyspnea (TAD)
- Febrile non-hemolytic transfusion reaction (FNHTR)
- Hypotensive transfusion reaction
- Delayed hemolytic transfusion reaction (DHTR)
- Delayed serologic transfusion reaction (DSTR)
- Acute hemolytic transfusion reaction (AHTR)
- Major cardiac event (MCE)
- Haematoma (bruise)
- Arterial puncture
- Delayed bleeding (re-bleeding)
- Nerve injury/irritation
- Localized infection/inflammation
- Deep venous thrombosis (DVT)
- Arteriovenous fistula
- Compartment syndrome
- Brachial artery pseudoaneurysm
- Vasovagal reactions

Where is the study run from?

The study is run from Haukeland University Hospital, Bergen, Norway.

When is the study starting and how long is it expected to run for? February 2022 to May 2024

Who is funding the study? The study is funded by Hemanext Inc. (USA), the company that produces the blood bags.

Who is the main contact? Laurel Omert, MD, laurel.omert@hemanext.com

Contact information

Type(s) Principal Investigator

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Additional identifiers

EudraCT/CTIS number Nil known

IRAS number

ClinicalTrials.gov number NCT05549232

Secondary identifying numbers PRO-CLIN-0012

Study information

Scientific Title

A single center, pilot clinical investigation of surgical bleeding in burn patients, and chronically transfused patients with haematological malignancies, who are transfused with hypoxic red blood cells manufactured with Hemanext ONE system

Study objectives

Anaemia results from either the acquired loss of red blood cells (RBCs) due to disease, injury, or surgery or the inability of the patient to produce their own RBCs due inherited disorders. The primary therapeutic goal of RBC transfusion is to increase the oxygen-carrying capacity of the blood and prevent tissue hypoxia. Substantial evidence suggests decreased quality of RBCs from substances that accumulate during RBC storage.

Hypoxic storage, where the oxygen content of RBC units is reduced to low levels (e.g., less than 20% oxy-haemoglobin prior to refrigeration and maintained throughout storage), has been indicated by data from in vitro and pre-clinical studies to reduce oxidative stress, counteract RBC metabolic impairments, and improve RBC deformability and oxygen delivery. The use of hypoxic

RBCs for blood transfusion may result in improved outcomes, both in acute and chronic transfusions settings.

The Hemanext ONE system can be used to manufacture and store hypoxic RBCs for any patient requiring a blood transfusion. It received its CE Mark in April 2021. The Post Market Clinical Evaluation Plan for Hemanext ONE requires collection of safety and efficacy data for the device. This pilot investigation was conducted to begin to inform on the safety of the blood product resulting from processing leukocyte reduced RBCs with the device.

The investigation aimed to study safety of hypoxic RBCs in 2 patient groups: acute burn and haematologic malignancies. Burn patients undergoing surgical excision routinely require large quantities of blood to be transfused. Patients with haematological malignancies often require regular transfusions.

Ethics approval required

Ethics approval required

Ethics approval(s)

Approved 27/06/2022, REK sør-øst B (Regional Ethics Committee South East B) (Gullhaugveien 1-3, Oslo, 0484, Norway; +4722845511; rek-sorost@medisin.uio.no), ref: 381996

Study design

Single-center prospective open-label single treatment clinical investigation

Primary study design

Interventional

Secondary study design Non randomised study

Study setting(s)

Hospital

Study type(s)

Safety

Participant information sheet

Not available in web format, please use contact details to request a participant information sheet.

Health condition(s) or problem(s) studied

Safety of hypoxic RBCs in 2 patient groups: acute burn and haematologic malignancies.

Interventions

Transfusion of two units of Red Blood Cells, Leukocytes reduced, O2/CO2 reduced (single transfusion course) manufactured with the Hemanext ONE System, combined with PRN standard Red Blood Cells, as applicable. Patients will be enrolled, transfused and followed-up until their subsequent transfusion or 28 days (+/- 1 day) post-transfusion, whichever comes first.

Intervention Type

Device

Pharmaceutical study type(s)

Not Applicable

Phase Not Applicable

Drug/device/biological/vaccine name(s)

Hemanext ONE

Primary outcome measure

Adverse events in patients are monitored by investigators during transfusion visit and by phone 24 hours and 7 days (+/- 1) after the transfusion visit.

Secondary outcome measures

1. Blood pH, pO2, pCO2, HCO3 and base (excess) is measured by following the standard practice at the site before, during and after the transfusion (Burn patients group only).

2. Haemoglobin level is measured by following the standard practice at the site before the following transfusion.

3. Frequency of all AEs, including transfusion related AEs are monitored by investigators up to 7 days (+/- 1) post-transfusion.

4. AEs are monitored by investigators up to 28 days after the initial transfusion, or up to subsequent transfusion episode, whichever comes first.

5. Frequency of all AEs are monitored, through the assessment of patient's diary, when applicable, from enrolment up to 7 days (+/-1 day) post-transfusion.

6. Vital signs are monitored by investigators over the course of the transfusion and up to 15 minutes post-transfusion.

Overall study start date

11/02/2022

Completion date

16/05/2024

Eligibility

Key inclusion criteria

A. Hematological malignancies patients group:

- 1. Male or female patients at least 18 years of age
- 2. Patients expected to require > 2 units of red blood cells in a single transfusion event

3. Patients who have the capacity to consent to particpate and are willing to comply with the study procedures.

4. Patients identified by a Transfusion hemoglobin trigger of less than 9 g/dL

5. Patients with a documented diagnosis of leukemia, myelomatosis or MDS requiring chronic transfusions

B. Burn patients group:

1. Male or female patients at least 18 years of age

2. Patients who have the capacity to consent by themselves to participate to the clinical investigation

3. Smaller burn patients, hospitalized with a Total Body Surface Area (TBSA%) burn \geq 10% and \leq 50%

4. Patients expected to require > 2 unit of red blood cells in a single transfusion event

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex Both

Target number of participants 20

Total final enrolment

22

Key exclusion criteria

- A. Both patients groups:
- 1. Patients with any positive antibody screening test
- 2. Patients for whom consent has not been obtained
- 3. Patients with a known haemolytic anemia (congenital or acquired)
- 4. Patients < 18 years old
- 5. Patients with a known or suspected pregnancy
- 6. Patients with a history of major transfusion reactions
- 7. Patients whom the Investigator deems clinical trial participation is not in their best interest.

B. Burn patients specific exclusion criteria:

1. Patients who do not have the capacity to consent by themselves to participate to the clinical investigation

2. Patients hospitalized with a Total body surface area (TBSA%) burn more than 50%

3. Patients with combined trauma in need of blood transfusions for treatment other than the burn excision

Date of first enrolment

30/08/2022

Date of final enrolment

16/04/2024

Locations

Countries of recruitment Norway

Study participating centre

Haukeland University Hospital Jonas Lies vei 65 Bergen Norway 5021

Sponsor information

Organisation Hemanext Inc.

Sponsor details 99 Hayden Avenue, Suite 620 Lexington, MA United States of America 02421 +1 847 722 57 10 hemanext@linkmedical.eu

Sponsor type Industry

Funder(s)

Funder type Other

Funder Name Investigator initiated and funded

Results and Publications

Publication and dissemination plan

The results of this study may be published or presented at scientific meetings by the investigator after publication of the overall study results or 1 year after the end of the study (or study termination), whichever comes first.

For all publications relating to the study, the investigator will comply with recognized ethical standards concerning publications and authorship, including those established by the International Committee of Medical Journal Editors. Authorship of publications for the overall study results will be determined by mutual agreement and in line with International Committee of Medical Journal Editors.

Intention to publish date

16/05/2025

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are available from the corresponding author on reasonable request laurel.omert@hemanext.com

IPD sharing plan summary

Available on request