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A study comparing the effects of using an automated electronic system and a paper-based system of recording patients observations on the length of stay in a hospital for trauma patients

Submission date
10/05/2012

Recruitment status
No longer recruiting

Registration date
16/05/2012

Overall study status
Completed

Last Edited
02/11/2018

Condition category
Other

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[X] Results
[_] Individual participant data

Plain English summary of protocol

Background and study aims
In current practice, each time a set of observations (blood pressure, heart rate etc) are performed they are recorded on a paper chart. Each result is given a score. The scores for each observation are manually summed, and the total score is recorded on the paper chart. If the total score is too high, the nurse at the bedside will respond by asking for help from a doctor or senior nurse. This method for helping doctors and nurses to recognise when patients are becoming unwell is called an Early Warning System. In 2007 the National Institute of Clinical Excellence advised that wards use an Early Warning System and recommended that the use of electronic systems to facilitate the process should be investigated. The study will test whether recording medical observations electronically can help doctors and nurses to improve outcomes for patients.

Who can participate?
Any patient over 16 years of age admitted to the John Radcliffe Hospital (wards 2A and 3A) in Oxford over an eight month period.

What does the study involve?
Whilst a patient is in hospital their medical observations will be recorded either using the paper-based system or using the electronic system. This will depend on which part of the ward a patient is initially admitted to. The nursing staff are guided to perform exactly the same observations on the patients regardless of whether the results of those observations are being recorded on paper or electronically.
To investigate whether there are any differences between the paper-based and electronic methods of recording the observations we ask trained nursing staff to look at the patients' medical notes.

What are the possible benefits and risks of participating?
The implementation of an electronic based track and trigger system may improve relevant patient outcomes in comparison to an optimised paper based version. There are no known risks for this study. Only data collected from the patients stay will be used.

Where is the study run from?
The study is being carried out on the Trauma unit at the John Radcliffe Hospital (wards 2A and 3A)

When is study starting and how long is it expected to run for?
August 2011 to July 2012

Who is funding the study?
Biomedical Research Council (UK)

Who is the main contact?
The main contact for any questions regarding this study is the OTEST study research team, who can be contacted on 01865 572282.
Dr Peter Watkinson, Principal Investigator for the study and Intensive Care Consultant, can be contacted via The John Radcliffe Hospital, Headley Way, Oxford OX3 9DU. Tel: 01865 741166.

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
N/A

Study information

Scientific Title

Oxford Track-&-Trigger Electronic System for Trauma (OTEST) - a randomised stepped wedge trial comparing the effects of an integrated electronic 'Track-and-Trigger' system and a paper-based 'Track-and-Trigger' system on the length of trauma patients stay from admission date to 'fit for discharge'

Acronym
OTEST

Study objectives
Will the implementation of an electronic based track and trigger system improve relevant patient outcomes in comparison to an optimised paper based version?

The null hypothesis in this trial is that the electronic 'Track-and-Trigger' system will not affect the time between admission to the ward and 'fit-to-discharge' in comparison to a paper-based system; the alternative hypothesis that the electronic 'Track-and-Trigger' system will affect the time between admission to the ward and 'fit-to-discharge' in comparison to a paper-based system.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Cambridgeshire 2 Research Ethics Committee, 17/02/2011, ref: 11/HO308/11

Study design
Single-centre randomised stepped wedge interventional study

Primary study design
Interventional

Study type(s)
Screening

Health condition(s) or problem(s) studied
Patients admitted to a trauma ward

Interventions
Control 'paper system' arm:
A paper evidence-based 'track-and-trigger' system on which physiological observations of blood pressure, pulse, oxygen saturation, temperature and conscious level are recorded is used. A score is manually assigned to each observed physiological value dependent on that value's position within the range of values seen in hospitalised patients. The scores for each physiological value are manually summed to provide a single (summary) measure of the patient's physiological status. The frequency of subsequent observations and the clinical response to a patient's physiological status are then set by following an algorithm based on the summary measure. The control arm is essential to allow comparison with the current standard system, a comparison recommended by the National Institute for Clinical Excellence. Observation time is no longer than the patient's hospital stay. No follow-up other than survival at 30 days after admission.

Intervention 'electronic system' arm:
Physiological observations will be entered onto a tablet personal computer or personal digital assistant both integrated with a central display station. Scores will be automatically assigned and summed. The time of the next observation set and the appropriate clinical response will be automatically displayed according to the escalation algorithm. Visual warnings will be displayed if an observation set or clinical response is delayed. This study will use the VitalPAC v 1.2 system (The Learning Clinic, UK) to provide this functionality. The product and its software will be fully localised to both trust standards and those specific to the trauma unit (and has already been deployed in several NHS Trusts). Observation time is no longer than the patient's hospital stay. No follow-up other than survival at 30 days after admission.

Intervention Type
Other

Phase
Not Applicable

Primary outcome(s)
Time from ward admission to 'fit to discharge'. 'Fit to discharge' is defined as either discharged from the ward to home or alternative care or accepted by social services as a 'delayed discharge'. This primary outcome measure has been chosen instead of using length of stay as a significant number of patients have an extended length of stay whilst waiting for suitable support mechanisms to be put in place outside hospital.

Key secondary outcome(s))
Patient outcome measures:
1. Hospital length of stay
2. In hospital mortality
3. 30 day mortality following ward admission
4. Unplanned admissions to intensive care
5. Number of cardiac arrest calls

Compliance measures:
1. Appropriate escalation in observation frequency
2. Appropriate escalation in clinical intervention

Completion date
01/07/2012

Eligibility

Key inclusion criteria
All patients over 16 years of age admitted to two trauma wards over an eight month period

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Adult

Sex
All

Key exclusion criteria
1. Patients whose treatment plan is palliative at admission to the trauma ward
2. Patients with a learning disability
3. Patients unable to speak English and without a suitable translator
4. Patients in the custody of HM Prison Service

Date of first enrolment
31/08/2011

Date of final enrolment
01/07/2012

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Oxford University Hospitals NHS Trust
Oxford
United Kingdom
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Sponsor information

Organisation
Oxford University Hospitals NHS Trust (UK)

ROR
https://ror.org/03h2bh287

Funder(s)

Funder type
Research council

Funder Name
Biomedical Research Council (UK)

Results and Publications

Individual participant data (IPD) sharing plan
The data sharing plans for the current study are unknown and will be made available at a later date.

IPD sharing plan summary
Data sharing statement to be made available at a later date

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	31/10/2018		Yes	No