Trainee-led evaluation of the need for intershift recovery among emergency department doctors in the United Kingdom: the TIRED-UK study

Submission date	Recruitment status No longer recruiting	Prospectively registered		
09/09/2019		[X] Protocol		
Registration date 07/11/2019	Overall study status Completed	Statistical analysis plan		
		[X] Results		
Last Edited	Condition category	Individual participant data		
05/11/2020	Other			

Plain English summary of protocol

Background and study aims

Emergency departments (ED) provide patient care 24 hours a day all year round. This requires staff to work long consecutive shifts that can result in fatigue. It is recognised that fatigue negatively impacts productivity, exacerbates the risk of human error, and could be associated with occupational burnout. From a patient perspective staff with excess fatigue could led to reduced experience of care and increased patient safety concerns. Fatigue may be measured by a variety of approaches including psychometric testing, assessment of reaction speeds, and personal diaries. However, these methods are impractical for providing rapid assessment within a working population and have limited validity. With rates of burnout as high as 60% in ED doctors there is a need for effective measurements of fatigue in order to produce evidence-based solutions.

The 'need for recovery' (NFR) scale is a validated questionnaire originally developed in the Netherlands, to assess how work demands affect inter-shift recovery. It features eleven items requiring a 'yes' or 'no' response, takes only a few minutes to complete, and shows high acceptability amongst surveyed populations. An online survey using the NFR has been trialled by this research group in a single centre ED and was shown to be an acceptable means of measuring staff fatigue.

This study is a survey aiming to answer the question 'What is the baseline need for recovery (NFR) score among ED doctors in the UK and which factors influence NFR?'. It will achieve this by conducting a national study to characterise the baseline NFR score in ED doctors across the UK and determine whether there are any associations and differences between NFR scores and demographic, occupational, personal wellbeing, rota characteristics, or geographical region variables. The survey will be open for the period of one month and will take participants approximately 15 minutes to complete.

Who can participate?

Doctors with full or provisional registration with the General Medical Council who have been employed in their main role as an ED doctor for the preceding month at the time of completion of the survey.

What does the study involve? Participants will complete an online questionnaire

What are the possible benefits and risks of participating?

This study has the potential to aid patient safety by providing individuals and employers with an indication of who is most at risk of increased NFR score, whether disparities exist between different staff groups, departments, and localities, and whether increased need for recovery is a predictor of future occupational burnout.

Where is the study run from?

When is the study starting and how long is it expected to run for? June 2018 to October 2019

Who is funding the study? Royal College of Emergency Medicine

Who is the main contact? Dr Laura Cottey, laura.cottey@nhs.net

Contact information

Type(s)

Scientific

Contact name

Dr Laura Cottey

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Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

41938

Study information

Scientific Title

Trainee-led evaluation of the need for Intershift Recovery among Emergency Department doctors in the United Kingdom

Acronym

TIRED-UK

Study objectives

What is the baseline need for recovery (NFR) score among Emergency Department (ED) doctors in the UK and which factors influence NFR?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not required. Research limited to involvement of staff as participants (no involvement of patients/service users as participants)

Study design

Observational; Design type: Cross-sectional

Primary study design

Observational

Study type(s)

Other

Health condition(s) or problem(s) studied

Fatigue and recovery in doctors

Interventions

This study is being conducted as part of the Trainee Emergency Research Network collaborative (TERN), an initiative by the Royal College of Emergency Medicine to widen access to research by ED doctors and specifically trainees.

The methodology for the survey has been designed with regard to the Checklist for Reporting Results of Internet Esurveys (CHERRIES).

The design is a cross-sectional online survey with additional ED demographic data collected by a local TERN representative. Each participant will complete the survey once. This design has been

chosen for ease of access by participants and has been amended following PPI consultation where proposed questions were reviewed.

The 57-item cross sectional survey has been previously trialled during a feasibility study and was found to be acceptable to participants and to have good face validity. The survey will seek to gather information on demographic characteristics (6 items), operational/rota characteristics (25 items), NFR questionnaire (11 items), perceptions relating to burnout (2 items) and one free text item to seek participant's views on suggestions on recovery.

The survey has been designed to minimise respondent fatigue and can be completed in 10-15 minutes using a smart phone app, computer or laptop.

In addition to the individual doctor surveys, a nominated TERN representative will collect anonymous site-specific data from EDs which is collected locally and nationally. This methodology has been chosen as previous surveys have found when participants are asked questions which requires them to calculate shift lengths and maximum number of shifts based over a certain period this reduces responses. This data is the same for each group of doctors on a specific rota and therefore can be collected from the roster without burdening participants. The information collected will also include broader information on the characteristics of the ED, staffing, rota pattern, leave allocation, teaching &

training, consultant/ senior supervision, case-mix, specialist designation (e.g. Major Trauma Centre, Hyper- acute Stroke centre, etc.) which can then be associated with individuals' responses.

Intervention Type

Other

Primary outcome(s)

Baseline Need for Recovery (NFR) score amongst ED doctors in the UK assessed by NFR questionnaire.

Key secondary outcome(s))

Assessed by 57-item survey:

- 1. Demographic characteristics
- 2. Occupational characteristics
- 3. Well-being characterisitics

Completion date

10/10/2019

Eligibility

Key inclusion criteria

1. Doctors with full or provisional registration with the General Medical Council who have been employed in their main role as an ED doctor for the preceding month at the time of completion of the survey

This includes; ED consultants, ED specialist training doctors, ED associate specialist and staff grades doctors, acute common care stem trainees, GP trainees, Foundation Year one and two doctors, clinical fellows and trust grade doctors.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Total final enrolment

4247

Key exclusion criteria

1. Doctors whose main place of employment is outside of the ED. This includes speciality doctors employed in specialties other than Emergency Medicine. This will be identified in the demographic questions asked in the survey

2. EDs designated as Type 2, 3, or 4 by NHS England

Date of first enrolment

03/06/2019

Date of final enrolment

15/07/2019

Locations

Countries of recruitment

United Kingdom

England

Northern Ireland

Scotland

Wales

Study participating centre NHS Greater Glasgow and Clyde

J B Russell House Gartnavel Royal Hospital 1055 Great Western Road Glasgow United Kingdom G12 0XH

Study participating centre University Hospitals Coventry and Warwickshire NHS Trust

Clifford Bridge Road Coventry United Kingdom CV2 2DX

Study participating centre Frimley Health NHS Foundation Trust

Portsmouth Road Camberley United Kingdom GU16 7UJ

Study participating centre Cambridge University Hospitals NHS Foundation Trust

Addenbrookes Hospital Hills Road Cambridge United Kingdom CB2 0QQ

Study participating centre Croydon Health Services NHS Trust

London Road Thornton Heath United Kingdom CR7 7YE

Study participating centre University Hospitals Of Derby And Burton NHS Foundation Trust

Royal Derby Hospital Uttoxeter Road Derby United Kingdom DE22 3NE

Study participating centre East Kent Hospitals University NHS Foundation Trust

Kent & Canterbury Hospital

Ethelbert Road Canterbury United Kingdom CT1 3NG

Study participating centre Medway NHS Foundation Trust

Windmill Road Gillingham United Kingdom ME7 5NY

Study participating centre Northumbria Healthcare NHS Foundation Trust

Rake lane North Shields United Kingdom NE29 8NH

Study participating centre Royal Liverpool Hospital

Prescot Street Liverpool United Kingdom L7 8XP

Study participating centre Aintree University Hospital NHS Foundation Trust

Fazakerley Hospital Lower Lane Liverpool United Kingdom L7 9AL

Study participating centre Great Western Hospital

Marlborough Road Swindon United Kingdom SN3 6BB

Study participating centre Royal Devon and Exeter NHS Foundation Trust

Barrack Road Exeter United Kingdom EX2 5DW

Study participating centre St Helens And Knowsley Hospital Services NHS Trust

Whiston Hospital Warrington Road Precot United Kingdom L35 5DR

Study participating centre Northwick Park Hospital

Watford Road Harrow United Kingdom HA1 3UJ

Study participating centre Gloucester Royal Hospital

Gloucestershire Hospitals NHS Foundation Trust Great Western Road Gloucester United Kingdom GL1 3NN

Study participating centre University Hospital Southampton NHS Foundation Trust Southampton United Kingdom SO16 6YD

Study participating centre Southport And Ormskirk Hospital NHS TrustSouthport

United Kingdom PR8 6PN

Study participating centre Imperial College Healthcare NHS Trust London United Kingdom W2 1NY

Study participating centre
NHS Blackburn With Darwen CCG
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Study participating centre
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KY15 5UP

Study participating centre Lewisham And Greenwich NHS Trust London United Kingdom SE13 6LH

Study participating centre
University Hospitals Plymouth NHS Trust
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Torbay And South Devon NHS Foundation Trust
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United Kingdom
TQ2 7AA

Study participating centre University Hospitals of Leicester NHS Trust

Leicester United Kingdom LE1 5WW

Study participating centre Manchester University NHS Foundation Trust Manchester United Kingdom M13 9WL

Study participating centre
Mid Cheshire Hospitals NHS Foundation Trust
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CW1 4QJ

Study participating centre
Southern Health & Social Care Trust
Craigavon Area Hospital
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BT63 5QQ

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University Hospitals Birmingham NHS Foundation Trust
Birmingham
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Study participating centre
Cardiff & Vale University LHB
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CF14 4XW

Study participating centre South Eastern Health & Social Care

Dundonald United Kingdom BT16 1RH

Study participating centre Taunton And Somerset NHS Foundation Trust Taunton United Kingdom TA1 5DA

Study participating centre
South Tees Hospitals NHS Foundation Trust
Middlesborough
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Study participating centre
University Hospitals Bristol Nhs Foundation Trust
Bristol
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BS1 3NU

Study participating centre NHS Lothian Edinburgh United Kingdom EH1 3EG

Study participating centre
The Newcastle Upon Tyne Hospitals Nhs Foundation Trust
Newcastle
United Kingdom
NE7 7DN

Belfast Health & Social Care Trust

Belfast United Kingdom BT9 7AB

Study participating centre Pennine Acute Hospitals NHS Trust

Manchester United Kingdom M8 5RB

Study participating centre Guy's And St Thomas' Nhs Foundation Trust London United Kingdom SE1 9RT

Study participating centre
Portsmouth Hospitals Nhs Trust
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Chelsea And Westminster Hospital Nhs Foundation Trust
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SW10 9NH

Study participating centre
St George's University Hospitals Nhs Foundation Trust
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SW17 0QT

James Paget University Hospitals Nhs Foundation Trust

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E9 6SR

Airedale Nhs Foundation Trust

Keighley United Kingdom BD20 6TD

Study participating centre
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London
United Kingdom
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Study participating centre
North Middlesex University Hospital Nhs Trust
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Study participating centre Bedford Hospital Nhs Trust Bedford United Kingdom MK42 9DJ

Maidstone And Tunbridge Wells Nhs Trust

Maidstone United Kingdom ME16 9QQ

Study participating centre East And North Hertfordshire Nhs Trust

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Study participating centre Lewisham And Greenwich Nhs Trust London United Kingdom SE13 6LH

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Study participating centre
Harrogate And District Nhs Foundation Trust
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HG2 7SX

Study participating centre
Salford Royal Nhs Foundation Trust
Salford
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M6 8HD

Study participating centre
Sheffield Teaching Hospitals Nhs Foundation Trust
Sheffield
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S5 9AU

Surrey And Sussex Healthcare Nhs Trust

Redhill United Kingdom RH1 5RH

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Western Sussex Hospitals Nhs Foundation Trust
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University Hospitals Of North Midlands Nhs Trust
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Study participating centre

Doncaster And Bassetlaw Teaching Hospitals Nhs Foundation Trust

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NHS Ayrshire and Arran

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Study participating centre
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Bolton Nhs Foundation Trust
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Study participating centre
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Study participating centre
Brighton And Sussex University Hospitals Nhs Trust
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Sheffield Children's Nhs Foundation Trust

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Study participating centre Mid Cheshire Hospitals Nhs Foundation Trust

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Study participating centre Yeovil District Hospital Nhs Foundation Trust Yeovil United Kingdom BA21 4AT

Study participating centre NHS Forth Valley Stirling United Kingdom FK8 1DX

Sponsor information

Organisation

University Hospitals Plymouth NHS Trust

ROR

https://ror.org/05x3jck08

Funder(s)

Funder type

Research organisation

Funder Name

Royal College of Emergency Medicine; Grant Codes: GR222017

Results and Publications

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 from Dr Tom Roberts, TERN@rcem.ac.uk. Anonymised data to both individual and site will be available for one year after the study. This will be available to principal investigators. Others may request the data and such requests will be dealt with on a case by case basis.

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	02/11/2020	05/11/2020	Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes
<u>Protocol file</u>	Study website	20/02/2019	07/11/2019	No	No
Study website		11/11/2025	11/11/2025	No	Yes