

The SCIN (Skin Care Intervention in Nurses) Trial

Submission date 13/06/2013	Recruitment status No longer recruiting	☒ Prospectively registered ☒ Protocol
Registration date 21/06/2013	Overall study status Completed	☐ Statistical analysis plan ☒ Results
Last Edited 23/10/2019	Condition category Skin and Connective Tissue Diseases	☐ Individual participant data

Plain English summary of protocol

Background and study aims

Hand dermatitis (eczema) is a common cause of discomfort and loss of productivity in the NHS workforce with treatment costs amounting to £125 million/year. Different preventive measures are used to reduce hand dermatitis in healthcare workers but it is not clear how effective these are. To prevent infections spreading between patients, nurses have to wear gloves and wash their hands a lot of the time. This constant contact with detergents and water often leads nurses to develop hand dermatitis, which in turn may mean that they are more likely to carry harmful bacteria on their hands. The risk of getting hand dermatitis is particularly high in nurses with a past history of allergic disease such as asthma and childhood eczema and among nurses who work in intensive care units (ICUs) where patient contacts and hand washing are more frequent. Hand dermatitis can be reduced by avoiding unnecessary overuse of gloves and washing of hands, drying hands properly after washing, and regular use of hand creams (moisturisers). However, hand care by nurses is often poor. We will study nurses at high risk of hand dermatitis (student nurses with a past history of allergic disease and ICU nurses) and test the effectiveness and value for money of an online training programme (Behavioural Change Package or BCP) which aims to change nurses beliefs and behaviours in looking after their hands.

Who can participate?

This study aims to recruit, through the participating trust's Occupational Health (OH) department, at least 40 ICU nurses and 40 first year student nurses at each of 26 NHS trusts in England.

What does the study involve?

NHS trusts will be randomly allocated to one of two groups. 13 trusts will receive BCP and 13 trusts to act as controls. Nurses in the control trusts will be provided with the same written information as the nurses in trusts receiving BCP, and will be encouraged to report hand dermatitis early to their OH department. They will otherwise receive advice in accordance with their local trust's existing policies. Nurses receiving BCP will be encouraged to develop detailed plans to help them change their behaviours, supported by online reminders. The student nurses will be provided with personal tubes of moisturising cream. On ICU wards, facilities for washing and drying hands will be reduced and dispensers of moisturising creams installed. The participants will be asked to complete a questionnaire at the start of the study about past and current hand dermatitis, allergies, beliefs about hand care, use of gloves, soaps, alcohol hand rubs, moisturisers, activities outside work which may lead to dermatitis and current quality of

life. BCP will then be implemented in the allocated trusts. We will ask the nurses to complete a shorter questionnaire soon after they have received the BCP and again at the end of 12 months. In addition, any nurses who report hand dermatitis at any stage of the study will be asked to have their hands photographed and checked for harmful germs. Nurses with hand dermatitis will be treated according to normal practice in their trust. Two dermatologists from the study team will review the photographs to grade the severity of dermatitis. We will also randomly check the hands of nurses who do not have dermatitis.

What are the possible benefits and risks of participating?

All nurses participating in the study will benefit from intensive hand care advice. If the programme is effective, those in the BCP-receiving group will benefit from a lower risk of developing hand dermatitis. The results of the study will be circulated to the participants so that all nurses in the study will benefit from increased awareness of the importance of hand care at work. The study has been designed to have few, if any, adverse effects on the study participants. The programme is not painful or intrusive and should have little inconvenience. Once an individual has developed irritant hand dermatitis the outlook is poor. Individuals may develop sleep disturbance and hampered leisure activities.

Where is the study run from?

This study is being led by Guy's and St Thomas' NHS Foundation Trust but involves senior researchers and experts from King's College, London; the University of Southampton; Imperial College, London; University College London; the University of Manchester and Nottingham University Hospitals NHS Trust.

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

After a feasibility study in a hospital in Wales, the main study will start in September 2014 and will last until May 2017.

Who is funding the study?

The project is funded by the National Institute for Health Research (NIHR), UK

Who is the main contact?

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Contact information

Type(s)

Scientific

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Additional identifiers

Protocol serial number
HTA 11/94/01

Study information

Scientific Title
A cluster randomised controlled trial of a behavioural change package to prevent hand dermatitis in nurses working in the National Health Service

Acronym
SCIN

Study objectives
It is hypothesised that a behavioural change intervention to improve hand care, based on the theory of planned behaviour and implementation intentions, coupled with provision of hand moisturisers, can produce a clinically useful reduction in the occurrence of hand dermatitis when compared to standard care in at-risk nurses working in the National Health Service (NHS).

More details can be found at <http://www.nets.nihr.ac.uk/projects/hta/119401>
Protocol can be found at http://www.nets.nihr.ac.uk/__data/assets/pdf_file/0006/81186/PRO-11-94-01.pdf

Ethics approval required
Old ethics approval format

Ethics approval(s)
NRES Committee London - City Road & Hampstead, original application approved on 31/10/2013 (ref: 13/LO/09810) and substantial amendment application approved on 05/09/2014. Protocol v14 approved 05/11/2014.

Primary study design

Interventional

Study design

Four-year cluster randomised controlled trial

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Hand dermatitis

Interventions

1. Nurses in intervention trusts will receive the bespoke online behavioural change package (BCP). This will be developed by members of the study team with expertise in dermatology, occupational medicine, nursing, and health psychology and care will be taken to ensure compatibility with current guidance on infection control. It will include advice: on when and when not to use gloves; on when to use antibacterial hand rubs; on when and how to wash and dry hands; on when to use moisturising cream; and to contact OH early if hand dermatitis occurs. As part of the package, nurses will be asked to form implementation intentions for performing behaviours in their workplace. These will be recorded, and participants will subsequently be reminded of them and offered the opportunity to revise them. The package will be supported by provisions to encourage adherence, such as moisturising creams. It will be actively reinforced over the course of the study by consistent messages on skin care from local OH and control of infection teams, and from line management.
2. Nurses in control trusts will be provided with an advice leaflet about optimal hand care and encouragement to contact their occupational health department early if hand dermatitis occurs.

Intervention Type

Behavioural

Primary outcome(s)

The primary outcome is the change in point prevalence of visible hand dermatitis from baseline to the end of follow up.

Methodology: All study participants will be asked to contact their occupational health department early if they develop hand dermatitis. Their hands, wrists and forearms will be photographed using a standardised method. The images obtained will subsequently be assessed independently by two dermatologists blinded to other information about the participant, to ascertain whether any hand dermatitis is present. A participant will be classed as a case if both observers note visible hand dermatitis. In the event of disagreement, between the two dermatologists a third arbitrator will be consulted.

Key secondary outcome(s)

Secondary outcomes will be the difference between intervention and control trusts in:

1. The incidence of new episodes of hand dermatitis presenting to the OH department over the 12-months of follow-up
2. Severity of visible hand dermatitis from baseline to the end of follow-up
3. Days lost from sickness absence and days of modified duties because of hand dermatitis per 100 days of nurse time during the 12-months of follow-up
4. The change in estimated prevalence of hand colonisation by pathogenic bacteria (with or

without associated dermatitis) from baseline to the end of follow-up

5. The change from baseline to after completion of the BCP, and to the end of the 12-month follow-up in beliefs about dermatitis prevention behaviours.

6. The change from baseline to the end of follow-up in the reported frequency of use of alcohol hand rubs for hand cleansing, hand-washing with water, use of recommended techniques for drying hands after washing, use of moisturising creams and use of gloves for different durations (for student nurses, who will not have started clinical attachments at the beginning of the study, this will reduce to differences between the intervention and control trusts at the end of the follow-up)

7. The change from baseline to the end of follow-up in quality of life score

We will also document the reported participation in the behavioural change package, reasons given for not participating, and comments on its content.

Completion date

31/05/2017

Eligibility

Key inclusion criteria

1. All NHS trusts in England which train nurses, have an in-house occupational health (OH) service and have at least one intensive care unit (ICU)
2. First year student nurses and ICU nurses

Participant type(s)

Health professional

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Total final enrolment

1956

Key exclusion criteria

In order to avoid the risk of student nurses moving placements from an intervention to a control trust (or vice versa), only one trust in each city or town will be invited to participate. London is an exception to this where three trusts were identified in which student nurses did not move to neighbouring trusts during their training.

Date of first enrolment

01/09/2014

Date of final enrolment

01/12/2015

Locations

Countries of recruitment

United Kingdom

England

Scotland

Wales

Study participating centre

Guy's and St Thomas NHS Foundation Trust

The Education Centre

75-79 York Road

Waterloo

London

United Kingdom

SE1 7EH

Study participating centre

Cambridge University Hospitals NHS Foundation Trust

Cambridge Biomedical Centre

Robinson Way and Long Road

Cambridge

United Kingdom

CB2 0QQ

Study participating centre

East Kent University Hospitals NHS Foundation Trust

Kent & Canterbury Hospital

Ethelbert Road

Canterbury

United Kingdom

CT1 3NG

Study participating centre

Frimley Health NHS Foundation Trust

Frimley Park Hospital

Portsmouth Road

Frimley

Camberley
United Kingdom
GU16 7UJ

Study participating centre
Homerton University Hospital NHS Foundation Trust
Homerton Row
London
United Kingdom
E9 6SR

Study participating centre
Hull and East Yorkshire Hospitals NHS Trust
OH Department
Castle Hill Hospital
Castle Road
Cottingham
United Kingdom
HU16 5JQ

Study participating centre
Industrial Diagnostics
Atherstone House
The Sidings
Merry Lees Industrial Est.
Merry Lees
United Kingdom
LE9 9FE

Study participating centre
Kettering General Hospital NHS Foundation Trust
Rothwell Road
Kettering
United Kingdom
NN16 8UZ

Study participating centre
Lancashire Teaching Hospitals Foundation NHS Trust
LTH Occupational Health
Royal Preston Hospital
Sharoe Green Lane

Preston
United Kingdom
PR2 9HT

Study participating centre

NHS Lothian

NHS Lothian Occupational Health Service
Astley Ainslie Hospital
Grange Loan
Edinburgh
United Kingdom
EH9 2HL

Study participating centre

Luton and Dunstable University Hospital NHS Foundation Trust

Luton & Dunstable University Hospital
Lewsey Road
Luton
United Kingdom
LU4 0DZ

Study participating centre

Milton Keynes NHS Foundation Trust

Milton Keynes Hospital
Standing Way
Milton Keynes
United Kingdom
MK6 5LD

Study participating centre

Norfolk and Norwich University Hospitals NHS Foundation Trust

20 Rouen Road
Norwich
United Kingdom
NR1 1QQ

Study participating centre

Northampton General Hospital NHS Trust

Occupational Health Department
Billing House
Cliftonville

Northampton
United Kingdom
NN1 5BD

Study participating centre

North Tees NHS

University Hospital of North Tees
Hardwick
Stockton on Tees
United Kingdom
TS19 8PE

Study participating centre

Nottingham University Hospitals NHS Trust

Queens Medical Centre Campus
Nottingham Occupational Health
Derby Road
Nottingham
United Kingdom
NG7 2UH

Study participating centre

Oxford University Hospital NHS Trust

John Radcliffe Hospital
Headington
Oxford
United Kingdom
OX3 9DU

Study participating centre

Papworth Hospital NHS Foundation Trust

Papworth Everard
Cambridge
United Kingdom
CB23 3RE

Study participating centre

Pennine Acute Hospital NHS Foundation Trust

North Manchester General Hospital
Delaunays Road
Crumpsall

Manchester
United Kingdom
M8 5RB

Study participating centre
Plymouth Hospitals NHS Trust
Kingstor House
Derriford Hospital
Plymouth
United Kingdom
PL6 8DH

Study participating centre
Royal Berkshire NHS Foundation Trust
21 Craven Road
Reading
United Kingdom
RG1 5LE

Study participating centre
St George's Healthcare NHS Trust
St George's Healthcare NHS Trust
Blackshaw Road
Tooting
London
United Kingdom
SW17 0QT

Study participating centre
The Ipswich Hospital NHS Trust
Heath Road
Ipswich
United Kingdom
IP4 5PD

Study participating centre
The Leeds Teaching Hospitals NHS Trust
St. James's University Hospital
Beckett Street

Leeds
United Kingdom
LS9 7TF

Study participating centre

The Newcastle Upon Tyne Hospitals NHS Foundation Trust
Campus for Aging and Vitality (NGH)
Westgate Road
Newcastle Upon Tyne
United Kingdom
NE4 6BE

Study participating centre

University Hospitals Birmingham NHS Foundation Trust
Woodlands Nurses Home
Selly Oak Hospital
Raddlebarn
Birmingham
United Kingdom
B29 6JF

Study participating centre

University Hospitals of Leicester NHS Trust
Leicester Royal Infirmary
Infirmary Square
Leicester
United Kingdom
LE1 5WW

Study participating centre

University Hospital Southampton NHS Trust
Southampton General Hospital
Tremona Rd
Southampton
United Kingdom
SO16 3YD

Study participating centre

Sheffield Hospital NHS Trust
Northern General Hospital
Herries Road

Sheffield
United Kingdom
S5 7AU

Study participating centre
York Teaching Hospital NHS Foundation Trust
Centurion House
Centurion Park
Tribune Way
Clifton Moor
York
United Kingdom
YO30 4RY

Study participating centre
University of Manchester
The Mill
Sackville Street
Manchester
United Kingdom
M13 9PL

Study participating centre
University of Salford
Allerton Building
Frederick Road Campus
Salford
United Kingdom
M6 6PU"

Study participating centre
Wye Valley NHS Trust
County Hospital
Stonebow Rd
Hereford
United Kingdom
HR1 3AW

Study participating centre
West Suffolk NHS Foundation Trust
West Suffolk Hospital

Hardwick Lane
Bury St Edmunds
United Kingdom
IP33 2QZ

Study participating centre

Old Nurses Home

Glangwilli General Hospital
Carmarthen
United Kingdom
SA31 2AF

Study participating centre

Ty Hafren Bronglais

General Hospital
Aberystwyth
United Kingdom
SY23 1ER

Study participating centre

Prince Philip Hospital

Dafen Road
Llanelli
United Kingdom
SA14 8QF

Study participating centre

Withybush General Hospital

Fishguard Road
Haverfordwest
United Kingdom
SA61 2PZ

Sponsor information

Organisation

Guy's and St Thomas' NHS Foundation Trust (UK)

ROR

Funder(s)

Funder type

Government

Funder Name

Health Technology Assessment Programme

Alternative Name(s)

NIHR Health Technology Assessment Programme, Health Technology Assessment (HTA), HTA

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/10/2019	23/10/2019	Yes	No
Protocol article	protocol	17/03/2016		Yes	No
HRA research summary			28/06/2023	No	No