Merger of CCGs – what difference will it make for Primary Care research?

The NHS Long Term Plan set out the direction of travel for the NHS over the next decade with a range of implications for Primary Care research. Among them was the move towards more collaborative working between CCGs, and for Integrated Care Systems (ICSs) to cover England by April 2021. Importantly, the plan anticipated that there would be typically one CCG per ICS. This accelerated impetus towards CCGs working more strategically, at a larger scale. Across the West Midlands, the last year has seen CCGs merging and others sharing working arrangements (such as joint Accountable Officers) in preparation for merger by 2021.

What are the implications for Primary Care research?

Commissioning at a larger scale should allow advantages for the NHS and for patients. Working at a larger geography, covering populations of over one million, means CCGs are more able to drive collaboration between partner organisations, including health providers and local government to join up care and transform health, so creating opportunities for research across traditional interfaces. It should allow CCGs to more strategically plan for, and intervene in, long-term health conditions, to target groups with common characteristics and to address inequalities. Primary Care research should play an important part in generating evidence that supports more effective care and outcomes, responding to gaps in knowledge through designing and delivering research focused around the commissioning priorities and needs of local populations.

Inevitably, there is a risk of disruption to Primary Care research as new CCG arrangements come into place. The CRN Primary Care team will continue to work to ensure that such risks are minimised, that the merged CCGs are fully engaged with the importance of such research and are on board with getting systems running to ensure that the day-to-day delivery of Primary Care research continues to run smoothly.

We are looking forward to opportunities to work with the merged CCGs to innovate and develop new approaches to driving the development and delivery of research across more integrated healthcare system.

In this edition we feature articles on:

- Increasing physical activity in older people with joint pain (iPOPP), see page two
- Colour COPD, sputum colour charts to guide antibiotic self-treatment of acute exacerbation of COPD, see page four
- The impact of Giant Cell Arteritis on patients’ lives, see page 13

If you would like to contribute to Participate or for further information, please contact Jenny Oskiera, email jenny.oskiera@nihr.ac.uk
Cancer: Life Affirming Survivorship Support in Primary Care

**RENEWED Online Study**

An estimated 2.5 million people in the UK are cancer survivors (people who have finished primary treatment for cancer, whether or not they are cured), with this number on the increase. Anxiety and depression are common within this population as are fatigue and lack of physical activity. Studies show that healthy lifestyle changes and support for improving psychological wellbeing could improve quality of life for cancer survivors. The study aim is to evaluate an internet-based intervention (Renewed) to help support lifestyle changes and improve psychological wellbeing for breast, colon and prostate cancer survivors to improve quality of life. The RENEWED online intervention provides patients who have finished primary treatment for breast, colorectal or prostate cancer with self-management support for a healthy lifestyle and improved mood. This may lead to an improvement in their quality of life and prevention of cancer recurrence, with the primary outcome looking at whether using the web-based Renewed programme results in a difference in quality of life at six-month follow-up compared to treatment as usual. The long-term aim is for the intervention to potentially help cancer survivors with other forms of cancer.

We have now completed recruitment, exceeding the original target of 2,500 participants. This has taken nearly 500 practices across England, Wales and Scotland with over 50,000 Docmail letters sent out. Thank you for your support and contribution to reaching this target.

**Increasing Physical activity in Older People with joint Pain (iPOPP)**

**Study background**

Physical activity levels in older people with chronic musculoskeletal pain are low. Lower activity levels are associated with increased pain and disability. Walking is a straightforward way of increasing physical activity, which is accessible, inexpensive and low impact.

iPOPP is a three-arm randomised controlled trial which aims to test whether a brief behavioural intervention increases average step count compared to usual primary care or receiving a pedometer and activity diary in the post in adults aged 65 years and over with chronic musculoskeletal pain.

We are looking for approximately 57 practices to take part in the study. We need a practice population size of 400,000 within the West Midlands in order to recruit a total of 1,085 patients. Each practice will provide approximately 20 participants.

**What will be asked of practices?**

- Allow access to CRN staff to conduct a practice list search for potentially eligible patients
- GP to screen patient list for ineligible patients, CRN to complete Docmail invites for suitable patients
- Provide clinic time and space for a Health Care Assistant to deliver the walking intervention to patients (n=6, based on a list size of 7,000), which includes 2 x 30 minute appointments, the latter of which may be a telephone consultation

If you are signed up to the CRN Research Sites Initiative Scheme, this study will be paid at Grade One, £300.

**What are the benefits of participating?**

For patients this trial offers the opportunity to:

- get involved in research and potentially receive a programme of support to increase physical activity levels

For practices this trial offers the opportunity to:

- give their patients the chance to be involved in research
- participate in research which can be reported as part of appraisal and revalidation
The IMPPP study - Improving Medicines use in People with Polypharmacy in Primary Care

IMPPP is a large randomised clinical trial looking at how practice pharmacists and GPs, with the help of a new computer tool, can work together to improve the use of medicines in patients who are prescribed multiple medications in Primary Care.

What does the study involve?
The trial will operate in 54 GP sites across Bristol and West Midlands. Each practice will recruit 50 patients over a six-month period, and the participants will be followed up for a further six months.

Each participating practice will identify 260 eligible patients to be invited at the start of the study. Once patients have been invited to participate in the study, the practices will be randomised to either the intervention or control group. Practices in the control group will be asked to continue their usual care.

For practices in the intervention arm, the trial will involve GPs and practice pharmacists working together to deliver a structured polypharmacy medication review. Reviews will be conducted in batches over a six-month period. The study will fund additional time required for a practice pharmacist to undertake reviews. Where a practice does not have a pharmacist, one will be provided to undertake the reviews.

Practices will be provided with an IT tool which will support the case-finding, study administration and monitoring, and delivery of the polypharmacy medication review.

Practices will receive training for GPs and pharmacists, regular feedback and financial incentives for each full review completed. Funding will also be provided to cover the cost of reviews, initial trial set-up and clinical training.

For further information or if you are interested please contact Jenny Simm - Research Facilitator, contact details on back page. Prof. Carolyn Chew-Graham – Co-Investigator (Keele University), email: c.a.chew-graham@keele.ac.uk phone: 01782 734717. Dr Deborah McCahon – Trial Manager, email: deborah.mccahon@bristol.ac.uk

Attack

Aspirin To Target Arterial Events in Chronic Kidney Disease

is a pragmatic multicentre open-label randomised controlled trial to determine whether the addition of low-dose aspirin to usual care reduces the risk of major vascular events in people with chronic kidney disease who do not have pre-existing cardiovascular disease. This is a very simple study, with a very low workload for participating practices.

This study, run by the same researchers as those managing the HEAT, FAST GOUT and ALL HEART Primary Care studies, is running across the West Midlands, with over 1,000 patients recruited to date. Participating practices would receive service support costs to cover their time to help with this important study, and support would be provided.

Would your practice be interested in helping us with this national study?

The Trial Manager is Jen Dumbleton on 0115 823 1053 or jennifer.dumbleton@nottingham.ac.uk

What is the clinical and cost effectiveness of using a goal-directed allopurinol-based treat-to-target protocol in people with recurrent gout flares?

Patients will be randomised to either:
A: Treat to Target ULT or B: Usual GP care

Core Practice Activities

- Database search to identify patients with gout (CRN Support)
- GP to screen patient list for any inappropriate/ineligible patients
- Mail-out (DocMail or paper copies) and reminder after four weeks if no-response (CRN Support)
- Provision of suitable clinic room for patient visits with CRN Research Nurse
- Screening (30 minutes)
- Baseline, one year and two years post randomisation (one hour each)
- Practice Nurse required to initiate ULT as per the T2T protocol and will receive face-to-face training by the study team
- Data extraction of consented patients records
- Display study poster [Provided by study team]

Practice Target: six to seven patients

If your practice is interested in taking part, or for more information, please contact Gerri Mulcahy, CRN Research Facilitator, contact details on back page.
**Colour COPD**

- Sputum colour charts to guide antibiotic self-treatment of acute exacerbation of COPD

Colour COPD is a pragmatic multicentre, randomised controlled trial to determine whether the addition of a sputum colour chart to the existing self-management plan provided to patients with COPD improves their use of antibiotics and steroids and reduces the number of exacerbations they experience in a 12-month period. This is a very simple study, with a very low workload for participating practices.

This study will be running across the West Midlands and Greater Manchester. We are looking for 80 GP practices in total across the two locations. Participating practices would receive service support costs to cover their time to help with this important study, and support from the trial team will be provided.

**Would your practice be interested in helping us with this study?**

The trial team can be contacted on colourcopd@trials.bham.ac.uk or 0121 414 8137

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**Electronic Risk Assessment for Cancer - ERICA**

The ERICA study is a large randomised controlled trial assessing the clinical and cost effectiveness of six electronic risk assessment tools (eRATs) for bladder, kidney, lung, colorectal, ovarian and oesophago-gastric cancers in general practice. We will recruit 530 English practices to compare the effect of eRATs (vs usual care) on: cancer staging at time of diagnosis, cost to the NHS, patient experience of care, and service delivery.

We hope to see a 4-5% increase in the proportion of early stage cancers diagnosed if the intervention is successful.

The trial runs for two years, with the software being available on EMIS, Vision and SystmOne. A pop-up appears when a patient aged 40+ has recorded symptoms/test results with a 2+% risk of one of the six cancers. A symptom checker is present for recording additional clinical events, leading to the recalculation of a new risk score. GPs decide the next appropriate course of action themselves. We estimate one to two pop-ups per GP per week.

For the main trial outcome, we are not asking practices to collect data; this is provided by National Cancer Registration and Analysis Service. Practices may choose to take part in nested studies involving giving feedback on the eRATs. Participating practices receive £470.55 if randomised to the intervention arm and £204.40 in the control arm. We provide full support for practices and will liaise with CCGs to arrange software installation.

**Principal Investigator:** Professor Willie Hamilton, CBE.

For more information, please contact us:

- **Tel:** 01392 726555
- **Email:** erica@exeter.ac.uk
- **Web:** www.theericatrial.co.uk
- **Twitter:** @EricaTrial

**For the ERICA study, we estimate one to two pop-ups per GP per week.**

Absenteism and presenteeism costs the NHS approximately £2.4 billion per year and is associated with worse patient outcomes. The main causes of NHS staff absenteeism are musculoskeletal complaints and mental ill-health. Lifestyle factors such as smoking, obesity and low levels of exercise leading to poor cardiovascular health are also important factors.

**Enhancing The Health Of NHS Staff** is a multicentre, randomised controlled pilot trial of an employee health screening clinic for NHS staff. The aim of this NIHR funded trial is to evaluate the effectiveness and cost-effectiveness of a complex intervention in reducing absenteeism and presenteeism in NHS staff, comparing a hospital-based staff health screening and referral clinic with usual care. This pilot trial is due to start later this year. We aim to recruit 480 participants across three NHS Hospital Trusts in the West Midlands and Herefordshire.

**What will it involve for participants?**

- Participants will provide written consent and complete on-line questionnaires at baseline, 26 and 52 weeks
- Participants randomised to either attend the staff health clinic and receive assessment for their musculoskeletal, mental and cardiovascular health (or lifestyle advice for those <40 years) or usual care – they would not attend the staff health clinic but would see their GP if they had any health concerns

**What will it involve for GP practices?**

We will inform you if any of your patients consent to participate, notify you of any test results and potential actions that you may wish to consider, and may invite you to tell us about your experience of receiving information from the trial and the acceptability of the process.

If you would like to find out more please contact the trial team on 0121 414 8137 or ethos@trials.bham.ac.uk

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**West Midland Wide Studies**
Join Dementia Research - Innovation and Improvement Project

By Claire Brown, Research Nurse

Join Dementia Research (JDR) is an NHS and dementia charity initiative aimed at making it easier for people with and without dementia to participate in research studies. JDR provides people with the opportunity to register their interest and be matched with suitable research studies. Anyone over 18 can sign up online, by telephone, by completing a registration form, or through expressing interest via a JDR kiosk.

Recruitment to JDR has been slow, both in the West Midlands, and nationally. As part of the CRN West Midlands Primary Care wider Patient and Public Involvement and Engagement (PPIE) Strategy 2017/18, a group of Research Nurses, Research Facilitators and a Patient Research Ambassador planned an Innovation and Improvement (I&I) project to look at this issue. This included a smaller I&I project, looking specifically at the use of the JDR kiosks and how to maximise their impact.

The working group met every quarter to plan and monitor progress. The project looked at raising awareness of JDR in different settings with a variety of approaches. Impact was measured through qualitative descriptions (feedback) and quantitative measures (JDR registrations, expressions of interest via kiosks).

The kiosks were found to promote best engagement when staffed, they were rarely used when left unattended. In future, JDR kiosks should be focussed on larger events and venues to maximise expressions of interest while ensuring efficient use of staff time, and used in conjunction with leaflets and paper registration forms to offer alternative avenues for engagement. They need to be in a location where there is good Wi-Fi, or with CRN staff who can access mobile phone connectivity.

Promoting JDR with a stand at events such as Memory Walks and the 5k Chocolate Run proved particularly successful, as well as attending events like Memory Cafes and group sessions such as Singing for the Brain. Group members also attended practice nurse forums, patient participant group meetings and linking in with care homes; all were found to be an effective way to promote JDR and the JDR Learn Tool. Collaboration with CCGs has also been very successful; it was piloted in Herefordshire where they have included JDR as one of their five-year Dementia Strategy priorities, and set local targets for registrations.

Pharmacies were shown to be a more challenging area to promote JDR, with data suggesting only one registration resulted from promotion at four pharmacies during the project. Holding events in local libraries also showed varying degrees of success, the overall feedback being that it was quite time-consuming for CRN staff, with few resulting registrations as footfall on the day was unpredictable. Feedback from libraries has been that they are keen to support by displaying promotional materials.

Although the I&I project ended in December 2019, the group decided to continue working together to promote JDR and ensure the work to raise awareness continues.

For more information about JDR: https://nhs.joindementiaresearch.nihr.ac.uk/

Primary Care Clinical Research Lead

CRN West Midlands Primary Care are pleased to announce that Dr David Shukla has been appointed as the new CRN West Midlands Primary Care Clinical Research Lead (CRL); he will be starting in the role on 1 March 2020. David is a GP working in a busy teaching practice in Dudley and has been working with us as Primary Care Clinical Research Specialty Lead in the Central region since 2016. Prior to that, he was a GP champion for Primary Care.

Congratulations to Dr Shukla, we very much look forward to working with him in this new capacity. David will continue to undertake the CRSL role for the Central region.

Practice Pack

You may have seen a previous article about the Practice Pack that contains all the information you need to understand how the Clinical Research Network works with you and your staff and also to promote research generally to patients. The pack contains general research information and patient facing materials. This is being currently being circulated. If you have not received these materials please contact your local Research Facilitator.
Improving Practice Nurses’ Awareness of Join Dementia Research

Research Nurse Eleanor Hoverd and Public Research Champion Will Ryder spoke to Practice Nurses from Coventry & Rugby CCG about Join Dementia Research (JDR) at their Practice Nursing Forum. With only 0.9% of dementia sufferers registered with JDR locally (see Figure 1) it is vital that health professionals are aware of JDR and how to signpost patients and carers, so they have the chance to register and take part.

Very few Practice Nurses at the forum had heard of JDR, five years after the launch of the Challenge on Dementia 2020 (DoH 2016) which had key aspirations that by 2020 there would be:
- Equal access to diagnosis for all
- GPs providing a lead role in coordination and continuity of care for people with dementia
- Every person diagnosed with dementia having meaningful care following their diagnosis
- All NHS staff having received training on dementia appropriate to their role

The opportunity to speak to Practice Nurses with a Research Champion, or member of the public that volunteers to raise health research awareness, highlighted the need for improved communication on JDR.

A Research Champion’s perspective on what patients and carers want to know, and how to approach the subject of JDR during consultation was valuable and well received. Interestingly, recent JDR figures for Coventry & Rugby show that the most popular method for recruiting people with dementia is through the newspaper (see Figure 2). Practice Nurses do not feature as a recruitment source. Health professionals are in an ideal position to share information about dementia research with their patients and it poses the question as to whether many are aware of JDR and how to signpost patients and carers.

If you are interested in hearing more about JDR at a health professional event, please contact Eleanor Hoverd, Research Nurse on eleanor.hoverd@nihr.ac.uk

West Midlands update as at February 2020
- 22.42% of volunteers on the JDR system have enrolled into a study (national average 22.42%)
- 44.29% of volunteers with a confirmed diagnosis have enrolled onto a study (national average 36.13%)

These current results show what a great tool JDR can be to share research opportunities with members of the public, and for researchers to identify volunteers to contact.

If you would like any further information about JDR, promoting JDR or to receive JDR promotional literature, please do feel free to contact your local Primary Care team member or Jacqueline Smart on jacqueline.smart@nihr.ac.uk

Join Dementia Research (JDR)

Firstly, many thanks again to those who support raising the awareness of JDR amongst the contacts that you have developed.
The Clinical Research Network is working in collaboration with the University of Southampton and the Queen Elizabeth Hospital Birmingham on a trial which aims to determine the effectiveness of adding spironolactone to standard topical treatment, compared with placebo and standard topical treatment, for moderate or severe persistent facial acne in adult women.

Spironolactone has been used off-licence in acne for over 30 years due to its antiandrogenic properties and American guidelines suggest it has a role in treating acne, although there is little evidence of its benefit. If shown to be effective, spironolactone could reduce the use of antibiotics for acne, be cost-effective (cheaper than antibiotics) and would be more suitable for long-term use than other second line oral treatments.

**What is involved for practices?**

Practices are only asked to identify patients as a Participant Identification Centre. Patients will be asked to contact the research team, led by Dr Agustin Martin-Clavijo, Consultant Dermatologist, based at the Queen Elizabeth Hospital.

Practices will be asked to:

- Undertake a database search
- GP to check for exclusions
- Mail out using Docmail to eligible patients – women aged 18 years or older with a diagnosis of acne vulgaris
- Identify patients opportunistically during GP consultations
- Display recruitment materials

Participants and their GPs will be sent a letter at 24 weeks informing them whether the participant received spironolactone or a placebo. It is possible participants may consult after this point if they wish to take spironolactone. The consultant dermatologist’s letter will suggest potential action for the GP, if their patient asks to be prescribed spironolactone, as is usual care in some secondary care clinics. Participants will have their baseline urea and electrolytes checked by study team and ongoing monitoring is not necessary in this group.

**Practice remuneration**

**Service Support Costs:** to cover search and list checking £54.86 (pro rata based on the number of patients identified at the practice).

**Opportunistic recruitment:** £27.00 to identify one patient.

**Research Costs:** £72.00 to cover practice set up and mailing to cover practice set-up, search, screening and mailing.

Please note that this study also qualifies as part of the Research Site Initiative (RSI) Scheme and attracts an additional £100 payment to participating RSI practices.

**Funding**

SAFA is funded by the National Institute for Health Research HTA Programme. The study is currently recruiting from GP practices within CRN West Midlands Central area.

For more information or to express an interest in this study, please contact your local facilitator, Sheila Bailey, details on back page.

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**Retirement of Dr Soong Loy Yap, Maypole Health Centre, Birmingham**

The CRN WM (Central) region are very sad to see Dr Loy Yap, Lead GP for research at Maypole Health Centre, Birmingham retiring.

Maypole Health Centre joined the Research Sites Initiative scheme in 2014 and prior to that was research active working on a number of studies. Over the years he has recruited 538 patients. He was a very enthusiastic ambassador for primary care research and saw the benefits not only for his patients, but his own personal development as a GP.

We would like to thank Dr Yap for all his hard work and enthusiasm and it was always a pleasure working with him. We wish him a long and happy retirement.
**SNACKTIVITY to Promote Physical Activity and Reduce Future Risk of Disease in the Population (Work Package 1)**

There is strong evidence that being active and sitting less is important for health. Guidance states that adults should, over a week, complete at least 150-minutes of moderate to vigorous intensity physical activity (e.g. 30 minutes per day on five days of the week or through blocks of ten minutes or more). However, few people manage to achieve the current physical activity guidance because they have to make big changes to their lifestyle to achieve it, which can be too daunting and difficult.

The Clinical Research Network (CRN) is working in collaboration with Loughborough University and Birmingham Community Healthcare NHS Foundation Trust on a study called Snacktivity. Snacktivity aims to investigate whether encouraging frequent physical activity snacks of between two and five minutes throughout the whole day is a better way to get people more active and meet the recommended 150 minutes of moderate to vigorous intensity physical activity than either 30 minutes of activity on five days or in periods of ten minutes.

This project has received a five-year programme grant funded by the National Institute for Health Research (NIHR) and involves five related work packages. Work Package 1 (WP1) is primarily aimed at developing Snacktivity Intervention for testing in later feasibility and effectiveness trials. The person-based approach will be followed for developing complex interventions. The study team plans to conduct a series of studies to build and develop the Snacktivity intervention specification, to ensure it is underpinned by public opinion and is fit for purpose.

**Work Package 1 consists of three sub studies:**
- WP1a: public survey
- WP1b: focus groups to inform Snacktivity intervention design
- WP1c: development of the digital intervention component (Snackapp)

**Work Package 1a**
The first step in this programme grant is to conduct a survey of people’s views of Snacktivity. WP1a has recruited six practices across Birmingham with each to conduct a mail out to a random selection of 1,000 patients.

**Practices are requested to:**
- Conduct a pre-written search supported by the CRN inviting patients aged 18-80 to take part in the survey
- Mail a random selection of an estimated 1,000 patients at the practice, supported by the CRN

*Please note that practices are not being asked to screen the list as this is a low risk study. However, if a practice does wish to screen the list the cost for the time taken will be reimbursed.*

**Practice remuneration**
Practices will be paid £46.42 Service Support Costs for the set-up time for the study.
Practices that are part of the RSI scheme will receive a Study Level One payment of £300.00.

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**PUBLIC INVOLVEMENT AT THE HEART OF THE SNACKTIVITY TEAM**

Working with members of the public is central to the Snacktivity project. Magdalena Skrybant, Snacktivity’s Public Involvement Lead, led a successful process in July/August 2019 to recruit public contributors to the team. Snacktivity definitely captured the public imagination: the advert attracted over 70 Expressions of Interest, which were converted to 36 completed applications.

Ten public contributors from a range of backgrounds, and with different experiences of physical activity, were recruited to join Snacktivity’s Public Advisory Group (PAG). To welcome the PAG members to the Snacktivity team, an induction event was held, which included an overview of the Snacktivity project and an opportunity to meet the Chief Investigator, Amanda Daley, Snacktivity programme team and a number of investigators on the Snacktivity project.

Since the PAG was launched, its members have certainly been kept busy. The PAG has met three times and been involved in over 15 tasks, which include: providing input to the design of the Snacktivity logo; helping design patient-facing materials, including the Snacktivity survey and helping researchers understand what instructions to provide to participants when unboxing the wearable that will be used in the trial.

There is great enthusiasm for the project amongst Snacktivity PAG members and no shortage of requests for input from the PAG amongst Snacktivity researchers. Magdalena Skrybant will ensure that public involvement aligns to the NIHR Standards for Involvement, which were launched in November last year, and reported using GRIPP2, the international guidance for reporting public involvement in research.
Are Referrals to Hospital from out-of-hours Primary Care Associated with National Early Warning Scores?

By Samuel Finnikin, Gail Hayward, Fay Wilson, Daniel Lasserson

Background

The National Early Warning Scores (NEWS) is used in various healthcare settings to augment clinical decision making, and there is growing interest in its application in primary care. This research aimed to determine the distribution of NEWS among patients in UK out-of-hours (OOH) general practice and explore the relationship between NEWS and referral of patients to hospital.

Methods

A historical cohort study using routinely collected data from the Birmingham out-of-hours general practice research database. This includes patients who attended a large out-of-hours general practice provider in the West Midlands, between July 2013 and July 2018. All adults who were seen face to face, who had a full set of physiological observations recorded were included. NEWS was calculated post hoc, and subsequent hospital referral was the outcome of interest.

Results

A NEWS was calculated for 74,914 consultations. 46.9% of patients had a NEWS of 0, while 30.6% had a NEWS of 1. Patients were referred to hospital in 8.5% of all encounters. Only 6.9% (95% CI 6.3% to 7.5%) of the 6,878 patients referred to hospital had a NEWS of ≥5. Of the 1,509 patients with a NEWS ≥5, 68.6% (95%CI 66.2% to 70.9%) were not referred to hospital. When considering how NEWS was related to hospital referral, the Area Under the Receiver Operating Characteristic (AUROC) for patients seen in their own home was 0.731 (95%CI 0.681 to 0.787). For patient seen in treatment centres, the AUROC was 0.589 (95% CI 0.582 to 0.596).

Conclusions

Patients seen in out-of-hours general practice have low physiological acuity. Those referred to hospital have a slightly higher NEWS overall. NEWS is poorly associated with hospital referral, although the association is stronger for patients seen at home compared with patients seen in treatment centres. Implementing NEWS-based referral from OOH general practice is likely to increase hospital admissions.

Funding Available for GPs to Help with a Study to Understand the Nature and Scale of Avoidable Harm in Prisons

The Universities of Manchester, Cardiff and Nottingham and Greater Manchester Mental Health NHS Foundation Trust have been successful in obtaining funding for an NIHR Policy Research Programme looking at understanding the nature and scale of avoidable harm in prisons. The study will involve nurses and GPs (with five years’ post-qualification clinical experience) reviewing medical records and collecting data from prisons.

We are going to conduct a pilot in the first instance, with two prisons. We will then roll out further to approximately 14 other prisons. Funding will be provided for the data collection time in the prisons (at £225 per session/£56.25 per hour). Travel expenses (mileage) to the prisons will also be reimbursed. The training and data collection will commence late 2019 subject to approvals, and will continue through 2020 until the data collection is complete.

We would be grateful if you could provide the following information:

- Your potential availability (2020, e.g. number of sessions of data collection per week you might be able to commit to)
- Whether you have any experience as a GP in prisons/prison healthcare or interest in this area
- Whether you have experience of using SystmOne and number of years’ experience
- Number of years since qualifying as a GP
- How far you might be willing to travel to conduct data collection (to enable the research team to look at the logistics of the sampled prisons and GP availability)

If you are interested in finding out more information and in being involved in this project, please email the Research Officer, Christina Sheehan at christina.sheehan@nottingham.ac.uk

Dr Sam Finnikin is a CRN research active GP based at Ley Hill surgery, Sutton Coldfield
**Stakeholder Perceptions of Preventive Approaches to Rheumatoid Arthritis:**

**Qualitative study of healthcare professionals’ perspectives on predictive and preventive strategies**

We would like to invite you to take part in an interview study about strategies to predict the development of rheumatoid arthritis (RA) and treatments to reduce the risk of developing RA.

In the future it is possible that increasing numbers of people will be offered tests to predict their likelihood of developing RA. It may also be possible to treat people to reduce their risk of RA. The purpose of the study is to understand your thoughts regarding these strategies, what factors you think may influence their use and the potential for integrating them into clinical practice.

If you agree to take part in the study, you will be invited to take part in a one to one interview which will last around 30-60 minutes. Interviews can be done either face to face or on the telephone. During the interview, we will ask you to complete a short background questionnaire. We will then present you with two short scenarios describing individuals who present with some concerns about their health. After this, you will be asked some questions based on these scenarios as well as questions relating to predicting and preventing the development of RA.

If you would like further information, please contact Imogen Wells on lXW703@student.bham.ac.uk

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**Tasmin5s: Towards an Integrated Self-Monitoring Solution for Stroke/TIA: The BP Together Study**

People who have survived previous stroke or TIA are at particularly high risk of a subsequent stroke, with sub-optimal control of blood pressure a key reason. Previous research into GP supervised self-monitoring and self-management solutions have shown the approach to be less effective in post stroke patients. Tasmin5s hopes to build on this previous research and determine whether an integrated intervention combining self-monitoring/management of hypertension with a simple tele-monitoring system results in lower blood pressure than usual care for people with uncontrolled hypertension.

The study is a randomised controlled trial with participants randomised (1:1) to either:

- **Intervention:** Patients will self-monitor their blood pressure and send readings to an integrated monitoring system which GPs will be able to review monthly and direct patients to titrate their medication, request adherence testing or refer to a specialist

- **Usual care:** use of equivalent BP targets to the self-monitoring arm

All participants will receive a Stroke Association guide including recommendations for appropriate lifestyle intervention, lipid lowering medication and target blood pressures: ‘How to reduce your risk of stroke guide’.

This study is open to GP practices in the Central region if you are interested please contact your facilitator, Alastair Mobley, details on back page.

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**Are you a Primary Care GP or Nurse Working in an Area of High Ethnic Diversity or High Socio-Economic Deprivation?**

**Do you come into contact with parents or carers of children who have excess weight?**

We are looking for volunteers to take part in a study about having healthier weight conversations with parents or carers of primary school-age children.

We would like to talk to you about your experiences and concerns.

**Interested?**

Your participation will take no longer than 60 minutes and will be arranged at a time and location to suit you. You will receive £75 as a thank you for your time.

Please follow this link to access the Participant Information Leaflet http://bit.ly/38noSts and contact the study team if you would like to take part.

Phone: 0121 414 7869

supportstudyiahr@contacts.bham.ac.uk

www.birmingham.ac.uk/support-study
CRN West Midlands 2019 VIP Awards

Commercial Studies - Delivering to Time & Target Award

Citation: The practices accepting this award on behalf of Primary Care have contributed significantly to the overall metric of recruitment to time and target. They have truly embedded research as a core business activity within their practices and the dedication from their research teams and engagement from their Principal Investigators are what sets them apart from their counterparts. As a result of their consistent performance they have been able to establish a good reputation with commercial sponsors and Clinical Research Organisations. They act as advocates for clinical research within Primary Care and maintain strong working relationships with the CRN.

WINNER: SHERBOURNE MEDICAL CENTRE

This award went to the organisation(s) which showed greatest contribution to delivering recruitment to time and target for commercial studies.

Consideration was given to the number of commercial studies opened, recruitment of first patient, recruitment to time and target, approval times, using commercial income to build capacity to take on additional studies and developing relationships with commercial partners.

The practice table shows those practices that contributed to HLO2a: the proportion of commercial contract studies, achieving or surpassing their recruitment target during their planned recruitment period, at confirmed CRN sites for Primary Care.

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Our congratulations and thanks go to our well-motivated and high-achieving practices.

Patient Research Ambassadors change to Research Champions

The Clinical Research Network Coordinating Centre has previously described and supported the roles of both Patient Research Ambassadors and Join Dementia Research Champions. These roles were recently reviewed and the decision has been made to refer to both as Research Champions in future.

Research Champions include patients, carers and members of the public, who may or may not have taken part in a research study.

What they all have in common is that they are passionate about getting more people involved in research so that better care and treatments can be developed for everyone.

Research Champions volunteer their time to help spread the word about health and care research to patients and the public. They also help research and clinical staff understand more about the experiences of those who take part in research.

Please make sure to use the new name from now on.

If you would like more information about the role of the Research Champions in Primary Care or have any questions please contact Eleanor Hoverd, Research Nurse, eleanor.hoverd@nihr.ac.uk

Narratives Experiences Online (NEON)

Understanding the impact of mental health recovery stories

Background to the study

The NEON study has collected mental health recovery narratives from around the world. NEON is now running three online trials looking at whether accessing stories in this collection can help people with experience of psychosis or other mental health problems, and carers. NEON is currently recruiting across England. Visit www.recoverystories.uk for more information.

What will it involve for GP practices?

• Displaying a poster and some postcards about the study in the surgery (to April 2021)
• Circulating the website details as widely as possible www.recoverystories.uk

How do I get involved?

Posters can be downloaded from www.researchintorecovery.com/neontrials/promotion

If you would like copies of the posters and cards to be sent to your practice please email neon@nottingham.ac.uk

Geoff Robson
Research Champion Case Study

Our Research Champion (Primary Care) Geoff Robson shares his story:

1. A brief overview

I spent 40 years working for the rail industry, the majority in Project Management with the last 15 years developing and delivering major infrastructure projects. Having fully retired in mid-2017 I wanted to pursue some new challenges and decided to volunteer for some form of work within the NHS, although at the time I had no real knowledge about the opportunities.

Having become involved as a Research Champion, previously Patient Research Ambassador (PRA), my experience of health research has been very positive, although it does bring its challenges, as many GPs, and the majority of the public, lack full awareness.

2. How did you first hear about Research Champions?

Through my local GP Patient Participation Group I found out about the role of Lay Member on the Clinical Research Network West Midlands Partnership Group and put my name forward. Although my application was unsuccessful, I was told about the role of Research Champion within the Network and asked if I would be interested.

3. What made you decide to become a Research Champion?

I discussed this at length with Mary-Anne Darby (the CRN’s Head of Patient & Public Involvement and Engagement) who explained the role to me very well. To be actively involved in promoting research and helping others understand what research is about was a major motivation. As such, I agreed to take it on. The support I have had has been really helpful, with one to one meetings with Mary-Anne for the first six months, and meetings and communication with the various research teams.

4. Why do you think NHS research is important?

Research is key to finding treatments and cures for diseases and conditions that affect the lives of many people, thus reducing the burden on e.g. doctors’ surgeries, hospitals and care homes. The role of Research Champion provides an opportunity to inform and support those in the health industry to understand the type of research being undertaken and to ensure that the right level of public involvement is achieved.

5. What activities have you been involved with?

I have been involved in promoting involvement in dementia research including a radio interview, and organising an event at my local library. I helped at an event in a retirement village in Birmingham, contributed at meetings and presented at the PRA Away Day, among other activities. I am currently developing a plan to raise research awareness among student doctors, and the impact it will have in the future; I have become a co-applicant in a submission for the Parkrun Research Project and reviewed a paper aimed at research into the control of antibiotic use. This has been very rewarding and has enabled me to use my experience from previous employment to review the documents.

The work I have done to date has helped inform GPs and the public about what research involves and the tools available to help people get involved. It has also helped me to use my knowledge and experience of communication, presentations and document review to support various research initiatives. This year I am looking forward to helping engage with care homes regarding Join Dementia Research (JDR) and providing support as needed. I will also continue to support the JDR and Equality, Diversity and Inclusivity working groups.

6. What would you say to others who are considering getting involved in research?

Do it! Participation is the best way to add value to the research process, using experiences as a patient and carer. If you are considering becoming a Research Champion, there may be opportunities depending on the area you live in. You don’t need any specific experience of research or the health service, just a willingness to commit some time to promoting research through various means.

Remember, the role is voluntary so this does not necessarily mean that it will take up a great deal of time.
What is the Impact of Giant Cell Arteritis on Patients’ Lives?

Objectives
Clinical management of giant cell arteritis (GCA) involves balancing the risks and burdens arising from the disease with those arising from treatment, but there is little research on the nature of those burdens. We aimed to explore the impact of giant cell arteritis (GCA) and its treatment on patients’ lives.

Methods
UK patients with GCA participated in semi-structured telephone interviews. Inductive thematic analysis was employed.

Results
The overarching themes from analysis were:
• ongoing symptoms of the disease and its treatment and
• life-changing impacts

The overall impact of GCA on patients’ lives arose from a changing combination of symptoms, side effects, adaptations to everyday life and impacts on sense of normality. Important factors contributing to loss of normality were glucocorticoid-related treatment burdens and fear about possible future loss of vision.

Conclusions
The impact of GCA in patients’ everyday lives can be substantial, multifaceted and ongoing despite apparent control of disease activity. The findings of this study will help doctors better understand patient priorities, legitimise patients’ experiences of GCA and work with patients to set realistic treatment goals and plan adaptations to their everyday lives.
Approximately one in five people in the UK suffer from acute cough over the winter season. This makes it one of the most common reasons to visit a GP in the colder months, and subsequently costs the NHS around two billion pounds per year.

Many patients opt to seek relief from acute cough from over-the-counter (OTC) treatments, but very little research has been conducted into the effectiveness of OTC cough medicines; many of which have been described as ineffective by patients. Some trials into acute cough have been attempted in hospitals in the past, however, their success has been limited. This is likely to be related to the choice of setting, as cough patients rarely present at hospital, resulting in poor participant recruitment to the trials.

In 2015, the Rococo study took a different approach and, at the time of delivery, it was one of the largest multicentre, randomised controlled clinical trials in participants with coughs. The trial compared a branded OTC cough medicine ‘Unicough’ with a simple cough linctus to see if was capable of demonstrating significant reductions in acute cough symptoms. It successfully recruited 163 patients within just four months.

Rococo’s recruitment success was primarily down to its real-world approach. Instead of focussing on secondary care it looked at where patients first seek relief for an acute cough and, as a result, it was conducted exclusively through community-based sites. This included 14 pharmacies and four GP surgeries in England and resulted in Rococo being the first UK study to recruit participants seeking cough medicines from pharmacies.

Professor Surinder Birring, Consultant Respiratory Physician at King’s College Hospital London and Chief Investigator for the study, explains its significance for the future of community-based health research:

“This was an important study for community-based research as it was the first cough study to be done in a pharmacy setting; the most appropriate setting for the type of illness being studied. At the outset it was completely unknown how the trial was going to run. There were some initial concerns around gaining ethical approvals and some uncertainties around whether patients would want to be involved. However, the ethics board was very supportive of our requirements, and we concluded with results that clearly demonstrate the appropriateness of pharmacies as a research setting."

As outlined by Professor Birring, not only did the community-based approach maximise access to the target patient population, it also meant that study participants more closely resembled the broader population seeking cough medicines. This led to the generation of real-world data - more accurate and applicable results in terms of how effective the medicine is in the ‘real-world’ as opposed to a controlled clinical trial setting.

The recruitment success of Rococo was also in part down to support provided by the NIHR Clinical Research Network. Sinead Collinge, Industry Operations Manager for the West Midlands Clinical Research Network, explains how her team supported the study:

“The NIHR contributed significantly to efficient site identification and selection. We have a number of pharmacies in the area that are ‘research-ready’. By this we mean that they either have Royal Pharmaceutical Society ‘Research Ready’ status or they are already engaged in clinical research and have worked with the NIHR Clinical Research Network previously.”

The Royal Pharmaceutical Society runs a ‘Research Ready’ accreditation scheme for pharmacies in the UK. It requires a pharmacy to have a dedicated research lead and for all staff to have undertaken Good Clinical Practice (GCP) training which is mandatory for employees who will help to deliver a study. There are currently over 100 pharmacies registered. Sinead continues:

“Having access to research ready pharmacies really helps when it comes to getting a study up and running in a pharmacy setting. The staff already understand the principles that make the study run smoothly, such as how to collate, store and submit the data. But
Aim: Depression is usually managed in primary care, but most antidepressant trials are of patients from secondary care mental health services, with eligibility criteria based on diagnosis and severity of depressive symptoms. Antidepressants are now used in a much wider group of people than in previous regulatory trials. The clinical effectiveness was investigated of sertraline in patients in primary care with depressive symptoms ranging from mild to severe and tested the role of severity and duration in treatment response.

Method: The PANDA study was a pragmatic, multicentre, double-blind, placebo-controlled randomised trial of patients from 179 primary care surgeries in four UK cities (Bristol, Liverpool, London, and York). It included patients aged 18 to 74 years who had depressive symptoms of any severity or duration in the past two years, where there was clinical uncertainty about the benefit of an antidepressant. This strategy was designed to improve the generalisability of our sample to current use of antidepressants within primary care. Patients were randomly assigned (1:1) with a remote computer-generated code to sertraline or placebo, and were stratified by severity, duration, and site with random block length. Patients received one capsule (sertraline 50 mg or placebo orally) daily for one week then two capsules daily for up to 11 weeks, consistent with evidence on optimal dosages for efficacy and acceptability. The primary outcome was depressive symptoms six weeks after randomisation, measured by Patient Health Questionnaire, nine-item version (PHQ-9) scores. Secondary outcomes at two, six and 12 weeks were depressive symptoms and remission (PHQ-9 and Beck Depression Inventory-II), generalised anxiety symptoms (Generalised Anxiety Disorder Assessment seven-item version), mental and physical health-related quality of life (12-item Short-Form Health Survey), and self-reported improvement. All analyses compared groups as randomised (intention-to-treat).

Results/conclusion: 655 patients were recruited and randomly assigned - 326 (50%) to sertraline and 329 (50%) to placebo. Two patients in the sertraline group did not complete a substantial proportion of the baseline assessment and were excluded, leaving 653 patients in total. Due to attrition, primary outcome analyses were of 550 patients (266 in the sertraline group and 284 in the placebo group; 85% follow-up that did not differ by treatment allocation). No evidence was found that sertraline led to a clinically meaningful reduction in depressive symptoms at six weeks. The mean six-week PHQ-9 score was 7.98 (SD 5.63) in the sertraline group and 8.76 (5.86) in the placebo group (adjusted proportional difference 0.95, 95% CI 0.85–1.07; p=0.41). However, for secondary outcomes, evidence was found that sertraline led to reduced anxiety symptoms, better mental (but not physical) health-related quality of life, and self-reported improvements in mental health. Weak evidence was observed that depressive symptoms were reduced by sertraline at 12 weeks.

Importance: Sertraline is unlikely to reduce depressive symptoms within six weeks in primary care but we observed improvements in anxiety, quality of life, and self-rated mental health, which are likely to be clinically important. Our findings support the prescription of SSRI antidepressants in a wider group of participants than previously thought, including those with mild to moderate symptoms who do not meet diagnostic criteria for depression or generalised anxiety disorder.

Publication: https://www.thelancet.com/journals/lanpsy/article/PIIS2215-0366(19)30366-9/fulltext
Research Design Service

In an increasingly competitive research environment, securing funding to conduct health and social care research can be difficult and time consuming. The National Institute for Health Research (NIHR) Research Design Service provides expert advice and support to researchers developing research funding applications.

How we can help
We could help your team with:
• designing a research study
• research methods (qualitative and quantitative)
• identifying suitable sources of funding
• involving patients and public in research design
• identifying potential academic, clinical and lay collaborators
• identifying and refining the research question
• medical statistics
• health economics
• advice on common pitfalls
• interpreting feedback from funding panels

Who we can help
We can help you if you are:
• developing grant applications for applied health or social care research
• applying for personal fellowships
• writing applications to national, open, peer-reviewed funding streams

We support a broad range of people, including doctors, nurses and allied health professionals; patients and service users; academics and NHS and social care managers. Our priority is to support applications to NIHR research funding streams. We also support applications to Research Councils and other open, national, peer-reviewed funding programmes.

Why choose us
Expertise: Our research advisers offer a unique breadth of expertise and a proven track record in improving research applications.

Collaboration: We can help you to identify possible gaps in your research team and collaborators who can add value to your research, including health and social care clinicians, policy makers, academic researchers and patients and the public.

Tailored support: We can offer support tailored to the needs of your research team. Advice is available at face-to-face meetings, by telephone or email, at research clinics or as feedback from panel review meetings.

If you require further information please visit our website: https://www.rds-wm.nihr.ac.uk/ or contact us direct:

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Research Design Service

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