

# Arolwg o Brofiad Cyfranogion mewn Ymchwil (PRES)



Canolfan Cydlynu Rhwydwaith  
Ymchwil Glinigol NIHR  
(NIHR CRN CC)

Andi Skilton, Uwch Reolwr ar gyfer Cynnwys ac  
Ymgysylltu â'r Cyhoedd





Pa rwystrau neu gyfleoedd  
ydych chi'n eu gweld ar gyfer  
cyflenwi PRES?



# Cwestiynau PRES

FOR STAFF USE:

Study name/acronym: \_\_\_\_\_

Research site (Hospital/GP Practice): \_\_\_\_\_

Study IRAS/CPMS Number: \_\_\_\_\_

## Dewisol:

- oedrannau - 0-6;  
7-11; 12-15
- demograffeg
- gwybodaeth  
gyswllt

Please rate how strongly you disagree or agree with the following statements about your research experience by ticking the face or circle that matches your answer best.

**The information that I received before taking part prepared me for my experience on the study**

Strongly disagree      Strongly agree  I don't remember

**I feel I have been kept updated about the research**

Strongly disagree      Strongly agree  It is too early to tell

**I know how I will receive the results of the research**

Yes  Yes, to some extent  No

**I know how to contact someone from the research team if I have any questions or concerns**

Strongly disagree      Strongly agree

**The researchers have valued my taking part in the research**

Strongly disagree      Strongly agree

**Research staff have always treated me with courtesy and respect**

Strongly disagree      Strongly agree

**I would consider taking part in research again**

Strongly disagree      Strongly agree

2

Please use the boxes below to explain your answers to the above questions or provide any other feedback on your experience in research.

The comments that you write will be read in full by your healthcare provider and NIHR. We may use your comments in reports about research and for promotional activities, but we will remove any information that could identify you before publishing any of your feedback.

The NIHR Privacy Statement can be found here:  
[www.nihr.ac.uk/documents/nihr-privacy-policy/12242](http://www.nihr.ac.uk/documents/nihr-privacy-policy/12242)

**What was positive about your research experience?**

**What would have made your research experience better?**

**How long have you been taking part in this research study?**

Less than three months   
At least three months but less than one year   
At least one year but less than three years   
Three years or longer   
Not sure

**Is this the first research study you have taken part in?**

Yes  No

**Who completed the survey?**

The person taking part in the research   
The person taking part in the research with someone else   
Someone else on behalf of the person taking part in the research

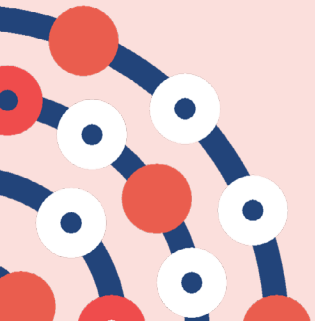
3

# Methodoleg PRES

- Wedi'i addasu a'i gydlynu'n rhanbarthol
- Timau cyflenwi'n ei ddosbarthu
- Ar bapur neu ar-lein
- 5 munud i'w gwblhau
- Ei roi yn ystod y cyfnod olaf neu ar yr un cyfnod ag ymgysylltu
- Ei gwblhau ar eu pennau eu hunain
- Unwaith fesul person fesul astudiaeth
- Nid oes angen cymeradwyaeth Pwyllgor Moeseg Ymchwil yr HRA



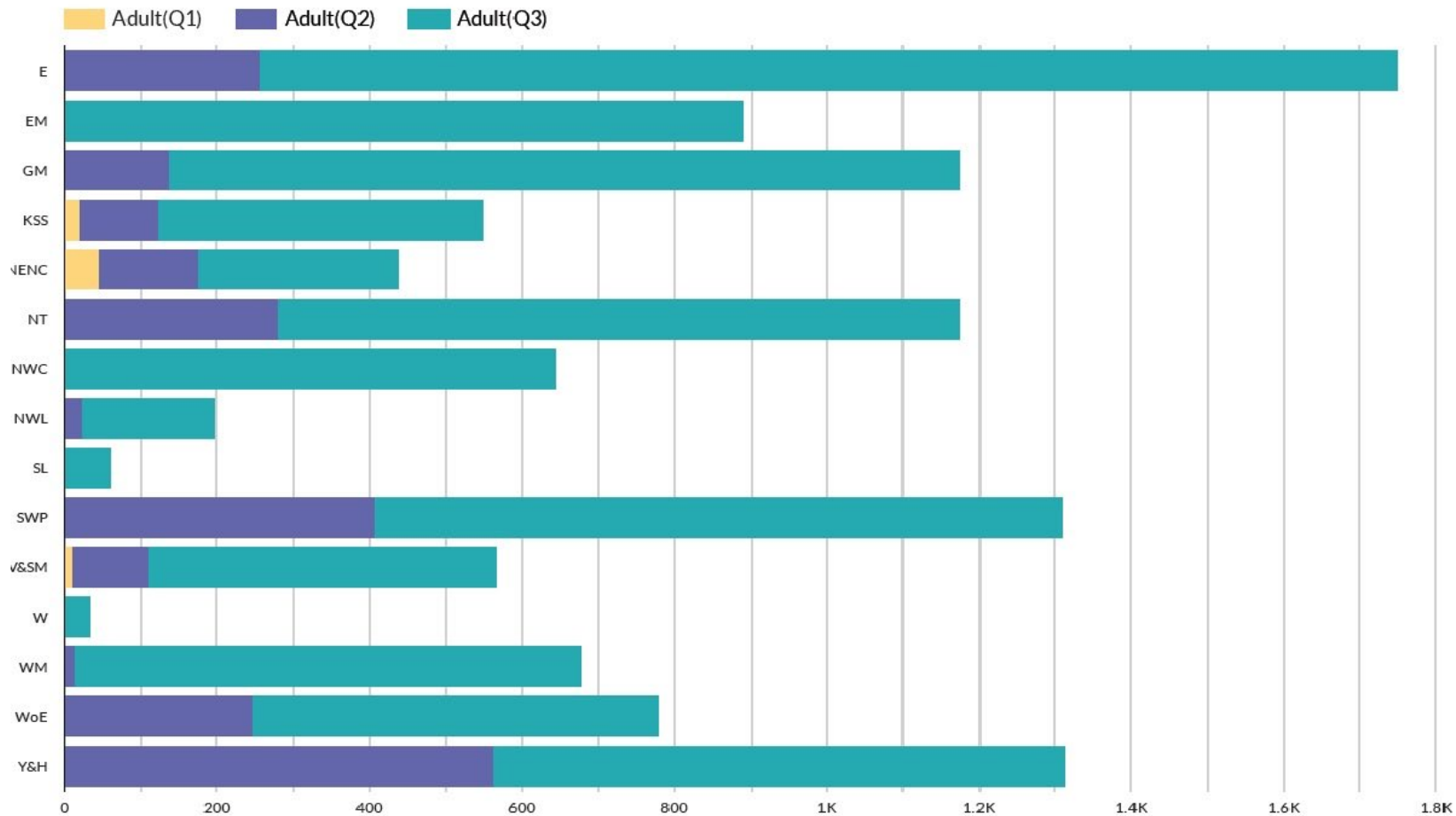
# Sut allech chi ddefnyddio canlyniadau PRES?



# C3 2020/21

- 11,562 o ymatebion oedolion
- 456 o astudiaethau
- 178/357 o Ymddiriedolaethau a Grwpiau Comisiynu Clinigol
- 75% o ymatebion ar gyfer Iechyd Cyhoeddus Brys
  - roedd 43% ar gyfer brechlynnau
- 49% yn ymatebion digidol

Astudiaethau y cafwyd y mwyaf o ymatebion iddyn nhw ( <i>brechlyn</i> )	<i>n</i>
SIREN	3109
Novavax ( <i>v</i> )	2839
Rhydychen/AstraZeneca ( <i>v</i> )	1905
ENSEMBLE 2 ( <i>v</i> )	414
Protocol Nodweddu Clinigol ar gyfer Haint Datblygol Difrifol	293



# Themâu enghreifftiol o PRES 20/21

## Brechlynnau COVID

Mae ofnau ynglŷn â diogelwch safleoedd yn codi fel mân broblem ond problem i'w hystyried ar hyn o bryd. Yn ôl un cyfranogwr, nid oedd wedi derbyn unrhyw wybodaeth am sut roedd yr ysbyty'n cael ei ddiogelu rhag COVID

Nid yw cyfranogwyr yn cyrraedd gyda disgwyliadau realistig ynglŷn â hyd ymweliadau ac amseroedd aros – mae'r amseroedd aros yna'n teimlo'n ddi-drefn pan nad ydy cyfranogion yn gwybod beth sy'n digwydd

Mae defnyddio apiau a dyddiaduron yn drysu rhai cyfranogwyr ac nid ydyn nhw'n deall yn iawn lle i gael cefnogaeth yn hyn o beth.

Mae llawer o gyfranogwyr yn derbyn llai o gyfathrebu nag roedden nhw'n ei ddisgwyl ynglŷn â chynnydd yr astudiaeth a'r hyn y mae disgwyl iddyn nhw ei wneud e.e. atgoffyn am eu hymweliad nesaf

Mae cyfranogwyr eisiau mwy o eglurder ynglŷn â sut y bydd rhoi brechlynnau cymeradwy ar waith yn effeithio ar yr astudiaeth y maen nhw'n rhan ohoni a hefyd maen nhw eisiau'r wybodaeth ddiweddaraf pan fydd materion yn cael sylw yn y newyddion, fel treialon yn dod i ben oherwydd diogelwch



# Themâu enghreifftiol o PRES 19/20

Mewnwelediadau	Argymhellion
<p>Dygwyd sylw at y ffaith bod cyfathrebu rheolaidd gydol y broses ymchwil yn bwysig i lawer o bobl.</p>	<p>Gallai sicrhau bod pobl yn derbyn y wybodaeth ddiweddaraf am newidiadau a gweithdrefnau ymchwil ym mhob cam o'r prosiect helpu i gadw cymhelliant pobl i barhau.</p>
<p>Mae pobl yn cymryd rhan mewn astudiaethau am amrywiaeth o resymau unigol, yn rhai personol ac yn rhai anhunanol.</p>	<p>Mae deall beth sy'n cymell rhywun i gymryd rhan yn gallu'ch helpu i ddeall sut i wneud iddyn nhw deimlo'u bod wedi ymgysylltu'n llawnach.</p>
<p>Mae'r berthynas â'r tîm ymchwil yn cael effaith o ran p'un a yw cyfranogion yn cael profiad ymchwil positif ai peidio.</p>	<p>Gellid gwneud profiadau gyda'r tîm ymchwil yn well fyth trwy amserlennu a pharodrwydd gwell a sicrhau bod y tîm ymchwil ar gael i ateb cwestiynau a rhoi sylw i bryderon sy'n codi.</p>

Filter Results By:

Partner Organisation  Study Name  Study ID  Speciality

Paper or Electronic?  Commercial or Non-Commercial  Age Group

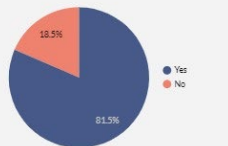
Year

Total surveys completed: **1,125**

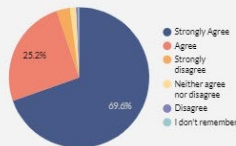
How long have you been taking part in this study?



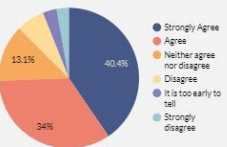
Is this the first study you have taken part in?



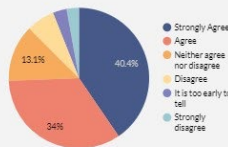
The information that I received before taking part prepared me for my experience on the study.



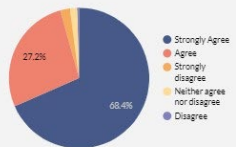
I feel I have been kept updated about the research.



I know how I will receive the results of the research.



I know how to contact someone from the research team if I have any questions or concerns



Search...



Research Participant Experience Survey Report 2018-19

Published: 20/05/2019 | Read Time: 32 minutes | Version: 4 | Print this document

Contents

What is this Report about?

This report shares results of the Research Participant Experience Survey 2018-19. The survey is about the experience of participants in health research. Individuals were surveyed between April 2018 and March 2019. The survey has been conducted annually by the National Institute of Health Research since 2015-16. It is carried out to help continually improve the experience of taking part in health research. It gives participants a chance to feedback on what went well and what could be improved and this report is produced and shared with a wide range of stakeholders in the health research community. This year the title of the report has changed from 'Patient Research Experience Survey' to 'Research Participant Experience Survey'. This is because individuals that take part in research are typically referred to as 'participants' and may not necessarily be patients.

A note on the terms used in the report

Where the expression 'n%' is used it means the number of people who answered that question. For example, 'n=807' means 807 answered the question.

Participants: an individual who took part in a health research study

Respondents: individuals who filled in this survey to feedback on their experience of participating in health research

1. What is this Report about?
  1. A note on the terms used in the report
  2. Acknowledgements
  3. Introduction
  3. Executive Summary
2. Delivery
  1. Design
  2. Research Staff:
  3. Study Information:
  4. Study organisation and environment:
3. Recommendations
  1. Recommendation
    1. Insight from Survey
  2. Recommendation
    1. Insight from Survey
  3. Recommendation
    1. Insight from Survey

Search...



Patient Research Experience Survey results 2019/20 - CRN Greater Manchester

Published: 07/09/2020 | Read Time: 3 minutes | Version: 1 | Print this document

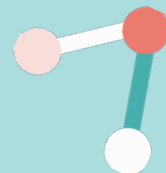
Contents

Greater Manchester patient responses - data overview

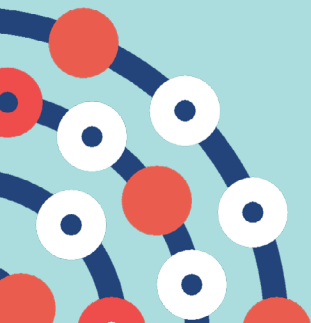
Which health area was the study in?

- Meningitis - 908**
- Cardiovascular - 119
  - Diabetes - 94
  - Lung - 64
  - Cancer - 103
  - Breast cancer - 45
  - Renal - 37
  - Urology - 40
  - Prostate cancer - 38
  - Neurology - 21

1. Greater Manchester patient responses - data overview
  1. Which health area was the study in?
  2. I was given all the information I needed in relation to the study.
  3. How old are you?
  4. How did you find out about this study?
  5. It is important to me to know the overall results of the research study. (rated on a scale of 1 to 5, where 1 is strongly disagree and 5 is strongly agree)
  6. I had / am having a good experience taking part in research. (rated on a scale of 1 to 5, where 1 is



# Enghreifftiau o PRES



Lift No.11

RESEARCH TODAY  
SAVES LIVES  
TOMORROW!

Did you know...

that Broomfield Hospital encourages patients to take part in research?

Research is for everyone and can be anything from letting us analyse your health records, to trying out a different or new treatment to improve your health.

Please ask your healthcare professional and GP about research today.

@MEHTresearch

“I WAS WELL  
INFORMED  
& LOOKED  
AFTER”

“THE STUDY  
PROVED  
BENEFICIAL  
& HELPFUL”

NHS  
Mid and  
South Essex  
NHS Foundation Trust



NIHR CRN East Midlands

@NIHRCRNEastMids

It's fantastic to see that so many people who have taken part in #COVID19 research are interested in being involved in more research studies in the future!

#ResearchVsCovid

NIHR | Clinical Research Network  
East Midlands



of people who took part in COVID-19 research in the East Midlands said that they would consider taking part in research again

Figure from the Participant in Research Experience (PRE) followed by NIHR CRN East Midlands and regional partners.

NIHR | Clinical Research  
Network Eastern

Recruitment Enhancement Tool for Research Delivery Teams:  
*Learning from the experience of people who have taken part in research*

This document is a recruitment enhancement tool, designed for use by clinical research teams delivering research studies. The information contained in this document is based on feedback from over 7,000 research participants between 2015 and 2020 via the Participant in Research Experience Survey. The survey is produced by the NIHR Clinical Research Network Eastern and delivered annually by research teams across the Eastern region. The information in this document comes directly from people who have taken part in a research study, their words, their experiences and their ideas for improvement.

The aim of the document is to give research staff a better understanding of what is important to the participant during the delivery of the research, so that they can make improvements in the service they provide and enhance participant experience where possible. Improving participant experience will help to ensure recruitment and retention to the study is successful.

Your voice is important too!

Research delivery staff can be influencers. You may feel that you cannot control what national study teams provide, however you are the patient's advocate. Feeding back comments and suggestions from patients on how to improve study delivery may not result in changes to the current study protocol but it could influence future study design.

NIHR | Clinical Research Network  
Thames Valley and  
South Midlands

CRN TVSM staff:  
could public  
volunteers help you  
explain your study  
better to patients?





crncc.ppie@leeds.ac.uk

Dolenni ychwanegol:

- [Gwefan NIHR](#)
- [NIHR CRN](#)
- [Portffolio NIHR CRN](#)
- [PRES](#)

