

Getting involved in health research - a quick guide for members of the public

Whether you have a health condition, have cared for someone who has, or are interested in learning more about research, your opinion on how we can create better medications and treatments for the NHS is valuable.

Patient and public involvement in research means that the public have a role in influencing, designing and carrying out research. Some examples of ways to be involved are reviewing lay summaries, participating in discussions with researchers and sharing your experiences.

We have put together some basic information for the public to find out more about what **'Involvement'** means in health research and how to go about getting started.



Getting involved in research is different to taking part in a study (known as participation) or learning more about research from researchers (known as public engagement) - both of these are important too.

Research needs you to get **involved!**



Most people or their families will have experience of using healthcare services in the NHS at some point in their lives.

We need as many voices from all backgrounds to make sure we are doing the right research to benefit everyone.



Those **involved** in research can help researchers

- design research studies to suit patients
- find and reach out to the best audiences
- communicate in the right way
- identify ways to measure outcomes
- provide the best solutions for patients
- share the results of research

What kind of research can I get involved in?

There are lots of different kinds of research that you can get involved in, and you don't need to have a specific health condition to contribute.

Examples might include hospital or community-based studies about specific health conditions, or studies of health in the course of normal life, such as childhood, pregnancy, ageing and bereavement.

What would I be expected to do?

Researchers need your thoughts about their research on how they can make their studies better for patients. Your questions and concerns about research can also be really valuable.

Most involvement activities ask you to provide your feedback in some way - either in person at a meeting or discussion group, or via email or phone.

What are the benefits of being involved?

- Opportunity to create positive change
- Learn more about a health condition
- Discover more about research and healthcare
- Develop new skills and increase confidence
- A chance to 'give something back' to the NHS

Don't just take our word for it, ask one of our public contributors:



"Being part of the PPI group has proved an extremely rewarding experience and provides a sense of contributing something of real value, no matter how small that might be. It is a very positive and socially stimulating experience"

Barbara, Cambridge University Hospitals
Panel member

Interested in joining the Cambridge University Hospitals Patient and Public Involvement Panel?

Our PPI panel provides feedback and ideas to researchers to make sure their research is more accessible and it meets the needs of patients and the public.



A lot of PPI activities can be done from home.

If you are interested in joining or like to find out more information send an email to CUH.PPI@nhs.net

Where can I find out more?

Useful websites

Opportunities for getting involved in research
www.peopleinresearch.org

Public Information Pack on getting involved
www.nihr.ac.uk/documents/public-information-pack-pip-how-to-get-involved-in-nhs-public-health-and-social-care-research/27388

Get involved NHS England
www.england.nhs.uk/get-involved/

Be part of research (taking part in research)
www.bepartofresearch.nihr.ac.uk/

Applied Research Collaboration
www.arc-eoe.nihr.ac.uk/patient-public-involvement

Healthtalk www.healthtalk.org/

UK Clinical Research Facility
www.ukcrfnetwork.co.uk/patients-public/

Association of Medical Research Charities
www.amrc.org.uk/

Patient Led Research Hub
www.plrh.org

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