RARE DISEASE CTN
RARE DISEASE CLINICAL TRIAL NETWORK

PUBLIC AND PATIENT INVOLVEMENT (PPI)

Public and Patient Involvement(PPI) is a way of doing research that supports collaboration between people with lived experience and researchers.

PPI means that research is carried out WITH or BY patients or members of the public rather than ABOUT, TO or FOR them.

LEARN MORE:

PPI IGNITE

The national organisation overseeing PPI, website has events and resources: https://ppinetwork.ie/

EUPATI FUNDAMENTALS

Learn the basics of patient engagement: eupati.eu/eupati-fundamentals/

HRCI

A national leader in PPI, they and their members sit at the interface of the health research community and the general public: hrci.ie/about-us/our-work/ppi/

DIGITAL BADGE

Certification inPPI, available through UCC, consists of modules and an assignment: ucc.ie/en/ppi-ignite/digitalbadge/

BITESIZE COURSE

Aimstohelpyou beginyour PPI activities: xerte.cahss.ed.ac.uk/play.php?template_id=2019 #page1

The Rare Disease Clinical Trial Network partners with contributors who have experience in any rare disease. They form our PPI Panel.

Contributors are equal partners in research projects and improve the quality and relevance of research. Learn more: rarediseaseresearch.ie

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Ireland's Global University

HRCI Health Research Charities Ireland