

Using an online survey to paint a picture of service user involvement in Healthcare Associated Infection Research

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Service User Research Forum (SURF)

SURF is a lay group committed to making a contribution to Healthcare Associated Infections (HCAI) research. Members are involved in a range of different research projects. Activities include setting research priorities, assisting in the development of applications, working as members of a research team, reviewing information and disseminating results.

The Survey

150 researchers in the field of Healthcare Associated Infections were asked to complete an anonymous online survey about their views and experiences of public involvement. This poster presents the data from the 50 responses received, along with a selection of the SURF group's reflections on the results, presented in the speech bubbles.

The percentage of researchers who say they have included public involvement: It's good that they are thinking

about involvement, no matter how big or small this may be. Yes No 52% 48%

Why public involvement is not being included:

We need to find out why they feel it isn't relevant, they may have legitimate answers, or it may reveal the need for greater education

are involving the public:

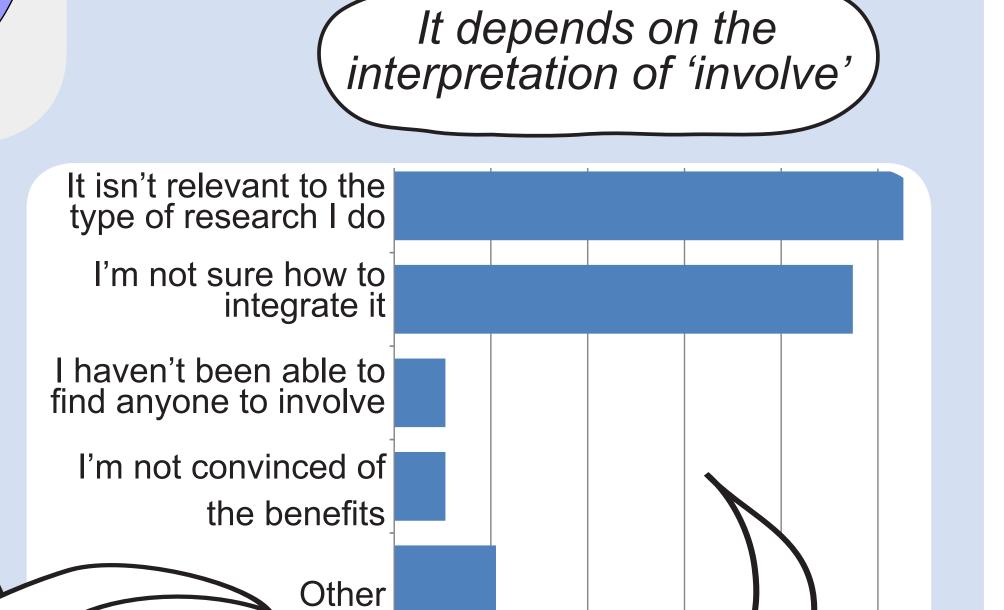
strategy

Input into recruitment

funding submission

Feedback on researcher

identified research priorities



All research is relevant as this will ultimately have an impact on the public, in health outcomes and the public purse,

20%

The benefits researchers reported:

Improved recruitment strategies

Rigour

Identify new perspectives

Improved team behaviour

Established new contacts and networks

Reminded me of end point of research and suffering associated with illness

...this makes me feel positive about my input...

particularly pleased to see the human element included, so often forgotten in the higher scheme of things

guidelines and recommendations Better written patient information

Enhanced development of

Improved acceptability to patients

...could be more emphasis on outcome - that it would be more trusted if lay members involved.

The challenges they experienced:

Identifying members of the public to involve

Time

Managing expectations about timescales involved in research

If the research takes a long time

it is difficult to feel part of the

team and keep up levels of

commitment

Having to adapt existing

misinterpreting results

ways of working

Service user

Found lay member agressive towards the team

Preoccupation with

personal experience

People often get involved because of adverse experiences, researchers need to understand this can impact on them. We don't mean to be aggressive - we are passionate!

Obtaining funding for involvement before grant secured

Researchers should think carefully about what they do and do not need help with from the public. Don't be afraid of not including us in everything.

Key stakeholders not supportive

The most popular Comment on roles responses involve for lay members 'easy' involvement Comment on ethical issues Include a lay co-applicant Input into study design Feedback on patient We're not surprised information by the most Help write patient information common responses, Input into recommendations/ they fit with our guidelines experience of being Advisory/ steering group involved member Input into lay summary for

0% 10% 20% 30% 40%

The most common ways researchers say they

..all a bit 'dry', more imaginative ways could be explored....

Do researchers think public involvement can improve research?

Interesting survey, would be good to follow up with interviews to get behind and inside the responses.

