Keynote presentation

**001 KP PATIENT AND PUBLIC INVOLVEMENT IN RESEARCH:
TENSIONS, CHALLENGES AND OPPORTUNITIES**

J Tritter. *School of Languages and Social Sciences, Aston University, Birmingham, UK*

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Increasingly research funders call for greater involvement of patients and the public in all aspects of research; an agenda that is even more prominent for public funding bodies. This agenda has been linked to a growing recognition of the relevance of patient experience in developing health services and patient-centred care and shared decision-making in clinical consultations. In this paper I explore the conceptual challenges in defining patient involvement and public involvement and the implications this has for public legitimacy in shaping research agendas and the research process. Patients have a different, and typically individual stake in research while members of the public as policy agents can enact a more collectivist orientation. Collaborative research is premised on different forms of expertise but is often framed as a contestation between different types of knowledge often reflecting the zero-sum model of decision making power that underpins Arnstein's Ladder of Citizen Participation. Such an approach fails to consider that members of a research team typically have different roles and responsibilities and that this does not necessarily diminish the collaboration. I conclude with reflections on how partnership in research should be based on the recognition of difference rather than equal roles and responsibilities. Such an approach, I argue may enhance rather than compromise research collaboration between academics, service users and members of the public.