The BMJ's own patient journey
Seeks to promote patient partnership by walking the talk

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A year ago The BMJ committed to setting up an international panel of patients, patient advocates, and clinicians to help us develop a strategy to advance the “patient revolution” in healthcare. This week we are launching it, with due thanks to the members of our advisory panel. Over the past six months they have stimulated, provoked, steered, and supported us to make landmark changes to our editorial processes. Changes that we hope will add to national and international efforts to improve the quality, safety, value, and sustainability of health systems through realising the transformative potential of working in partnership with patients, their families, communities, and advocacy groups. So what have we done?

Firstly, we have embedded patient peer review of research papers. We started in January with randomised controlled trials and have now extended it to all research papers where patient input would clearly be helpful. We are also calling for the submission of robust research papers that advance the science, art, implementation, and assessment of the impact of patient partnership, shared decision making, and patient centred care. Authors of research papers are being asked to document if and how they involved patients in defining the research question and outcome measures, the design and implementation of the study, and the dissemination of its results. Comment on the burden on patients of new interventions and treatments will also be required. This is not intended to be a box ticking exercise, but to encourage and properly report on collaborative research between patients and investigators.

Secondly, we are inviting more analysis and comment articles between patients and investigators. We have now extended this invitation to research papers. We may do this again, but our current initiative is not about one off gestures. We want partnership to become integral to how we work and think, as well as being something we advocate in healthcare. New expert patients on our editorial board and a newly appointed patient editor will help us in this quest and in time will enable us to meet a fourth pledge: to have patients participating in internal decision making committees. They are already flagging up innovatory initiatives and helping us to get the patient’s voice into the journal more. Past articles written by patients underline the self evident truth that people with experience of illness and navigating health services have much to teach us.

Our advisers have issued some clarion calls. One is to avoid tokenism. Initiatives to promote patient involvement and provide patient centred care are all too often poorly informed by patients and don’t promote partnership. A second is “do it well.” The failure to systematically collect and use the data on patient experience of care has been criticised. A third is to get the “authentic” voice of patients heard; not just that of the articulate minority. This is a challenge that we all grapple with, and one that is recognised by our patient advisers. We will continue to publish individual perspectives but are extending our links with patient networks and advocacy organisations to help tap into collective views. As we do this we recognise the importance of transparency. As in academic medicine, commercial influences are pervasive, and we are requesting and publishing conflict of interest statements from patient reviewers and authors exactly as we do with medical authors and reviewers.

The fifth tranche of our strategy is a campaigning one. We support patient control or co-ownership of personal health records and the “Patients included” initiative. Patients are thin on the ground in medical forums (the time and expense of attending deters many) and not routinely invited as speakers, and the dissemination of its results. We started in January with randomised controlled trials and have now extended it to all research papers where patient input would clearly be helpful. We are also calling for the submission of robust research papers that advance the science, art, implementation, and assessment of the impact of patient partnership, shared decision making, and patient centred care. Authors of research papers are being asked to document if and how they involved patients in defining the research question and outcome measures, the design and implementation of the study, and the dissemination of its results. Comment on the burden on patients of new interventions and treatments will also be required. This is not intended to be a box ticking exercise, but to encourage and properly report on collaborative research between patients and investigators.

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or on to steering committees. Here again our plan is to lead by example.

Our new strategy aligns with our “too much medicine” and “open data” campaigns and our support for “minimally disruptive medicine.” The BMJ remains a journal for doctors, but with Consumer Reports, which is partnered with the Choosing Wisely initiative, we are providing input into lay versions of the papers in our “overdiagnosis” series, to promote public awareness of the commercial and technological drivers shaping healthcare. Patients’ rights are another focus. Recent emphasis has been put on the right to be empowered to self-manage chronic conditions. Worldwide, the big issue remains the right to access good quality, affordable healthcare.

Patient partnership is a lot easier to talk about than to realise. It demands mutual respect and understanding. Our strategy is being implemented incrementally and will evolve as we evaluate its impact. But we are excited by its potential and hope readers will be too.

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