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Primary care research is important but there is general concern about the difficulty of linking research to patient care. Research development is even more problematic in primary care than in other disciplines. Policies to enhance general practice research include the creation of research networks, collaborations with research institutes, and early exposure to research during undergraduate teaching and specialty training. Early exposure prompts students to consider research as part of their future career, and better equips future practitioners to deliver evidence based patient care.

The importance of early exposure to research is generally accepted by both general practice registrars and directors of vocational training programs. Most programs include research curricula or related activities, including training in knowledge and skills, conducting research projects, or participation in research in daily practice. Although research curricula create more positive attitudes toward research, there are no indications that more registrars are participating in research, and we still know little about the long term effects of such curricula. Furthermore, most studies have assessed changes in registrars’ attitudes toward performing a research project or undertaking education in research skills during vocational training, rather than assessing registrars’ participation in research during daily practice.

This study analyses registrars’ participation in research tasks during their daily work with patients. The aim was to assess patient recruitment, factors influencing recruitment, and registrars’ views and suggestions with regard to participation.

Methods

Setting and design
Dutch general practice registrars undertake a 3 year specialty training program. They spend the first and third year in a training practice in the community, and in the second year they rotate between hospital posts.
In 2003 and 2004 our training program included participation of all third year registrars (70) in a controlled clinical trial as part of their core program.

The trial

A controlled clinical trial of the effectiveness of registrars using problem solving treatment (PST) – a brief psychological treatment to teach patients how to use their own skills to cope with problems – for patients with emotional symptoms. It is theoretically assumed that symptoms are reduced if problems can be resolved.\(^{13}\)

Registrars were randomly allocated to either the intervention group or the control group. Both groups recruited patients with emotional symptoms during their regular clinical work. We asked each registrar to recruit 4–6 patients who had presented for three or more consultations in the past 6 months, had a score of four or more on the 12 item general health questionnaire (GHQ-12), and who experienced emotional symptoms.

Exclusion criteria were severe medical illness, current contact with psychiatric services (or contact in the past year), current psychological treatment or past cognitive behavioral therapy, severe mental disorder, organic psychiatric disorder or substance misuse, active suicidal ideas, and lack of sufficient Dutch language to participate.

Registrars in the intervention group received a 2 day training course in PST\(^{14}\) and provided the psychological treatment to the patients they had recruited within 8 months. Registrars in the control group provided ‘care as usual’ and were asked to complete their patient recruitment within 4 months. The trial design was approved by the Medical Ethics Committee of the Radboud University Nijmegen Medical Centre, The Netherlands.

Outcomes and analysis

The authors administered a self developed questionnaire to explore registrars’ opinions about their participation in the trial, barriers they experienced in patient recruitment, and their opinions and suggestions with regard to enhancing research participation. Recruitment data were obtained from the trial records.

Recruitment data and scaled answers from the questionnaire were analysed with descriptive statistics and independent sample t-tests using SPSS statistical analysis software. The answers to open ended questions were independently ordered into categories.

Results

Sixty-seven of the 70 registrars participated in the trial (37 in intervention group [27 women] and 30 as controls [18 women]).

Registrars randomly allocated to the intervention or control group in 2003 expressed resistance to obligatory participation. The authors modified the process for the 2004 cohort, offering registrars an individual choice to participate in the training. Registrars who participated in PST training comprised the intervention group (17); the others, providing usual care, were regarded as the control group. The registrars were also offered more research assistance, and both groups were given 8 months to recruit patients.

Patient recruitment

The registrars in the 2003 intervention group recruited 83 patients; the registrars in the control group recruited 11 patients. The registrars in the 2004 intervention group recruited 66 patients; the control group 48 (Table 1). The authors explicitly asked registrars in 2004 to describe the difficulties they had experienced recruiting patients (Table 2).

Registrars’ opinions and suggestions

The questionnaire had a response rate of 84% (30 of 37 registrars in the intervention group, 26 of 30 registrars in the control group \(p=0.54\)).

Positive points reported by the registrars included the interesting and relevant nature of the topic, the opportunity to learn a new skill, becoming acquainted with and contributing to research and evidence based medicine, good research support, becoming more attentive

<table>
<thead>
<tr>
<th>Table 1. Recruitment of patients</th>
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<tbody>
<tr>
<td><strong>2003</strong></td>
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<tr>
<td>Mean number of patients per registrar</td>
</tr>
<tr>
<td>Intervention group</td>
</tr>
<tr>
<td>Control group*</td>
</tr>
<tr>
<td>Total</td>
</tr>
</tbody>
</table>

* Control group recruitment took 4 months in 2003 and 8 months in 2004 (intervention group recruitment took 8 months in both years)
General practice registrars and research – attitudes toward participation

RESEARCH

not differ from the moderate recruitment rates recorded by GPs: Peto et al.,15 for instance, found an average rate of 3.7 patients recruited per GP per annum. Registrars’ barriers and wishes are also comparable with those of GPs: moderate patient recruitment because of time pressures, the need for interesting, practice oriented and relevant projects, personal support, good information, and good feedback about the research results.3,15–17

Strengths and limitations

As far as the authors are aware, this is the first study exploring general practice registrars’ actual performance in research. The study explored the attitudes of registrars toward research rather than opinions of GPs or directors of vocational training programs. Furthermore, it studies registrars’ opinions about participation in a trial within their routine practice rather than requiring them to conduct a research project themselves.

The study is limited however, by the modest sample of registrars, all of whom belonged to the same training program and who participated in a single trial alone. However, the findings are similar to those of other studies concerning registrars’ appreciation of research experience.7,11

The study was compromised by the change made in 2004 to select registrars on the basis of their motivation. In 2003, however, the authors to diagnosing and treating emotional problems, and developing a critical view.

Negative points included the time investment required (this was especially mentioned by intervention group registrars). All registrars in 2003 criticised obligatory participation; only control group registrars were negative about obligatory participation in the 2004 cohort.

To improve participation, registrars suggested they be allowed to choose between several research projects to better match the research topic with their personal interests. They felt they needed to spend enough time in clinical practice training, early and good information about a research project, and involvement of their general practitioner tutors in patient recruitment.

In 2003, all respondents said that they wanted to be involved in research in the future; in 2004 this was the case for 14 of 15 intervention group respondents and nine of 13 control group respondents (Table 3).

Discussion

Sixty-seven registrars recruited 208 patients in total. Registrars expressed an interest in participation and appreciated contributing to research. They enjoyed learning a new skill and being more attentive to a particular disease and/or symptoms. Nevertheless, their patient recruitment rate was below the authors’ expectations.

Initially the obligatory nature of participation was considered to be an important barrier. Engaging registrars by offering the choice to voluntarily take part in the training did not however, result in major improvements in recruitment: the doubled patient recruitment rate in the control group can be attributed to the doubled recruitment period.

Registrars suggested that the option to choose an interesting and relevant research topic, the opportunity to learn new skills and a report of the research results would make participation in research projects more attractive.

The recruitment rate in this trial does not differ from the moderate recruitment rates recorded by GPs: Peto et al.,15 for instance, found an average rate of 3.7 patients recruited per GP per annum. Registrars’ barriers and wishes are also comparable with those of GPs: moderate patient recruitment because of time pressures, the need for interesting, practice oriented and relevant projects, personal support, good information, and good feedback about the research results.3,15–17

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Table 3. Requirements that make research projects attractive for participation during vocational training (2004, 31 respondents)

<table>
<thead>
<tr>
<th>Requirements of research project</th>
<th>Mean score on 5 point scale*</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Most important</strong></td>
<td></td>
</tr>
<tr>
<td>The topic should be interesting</td>
<td>4.7</td>
</tr>
<tr>
<td>The topic should be relevant to general practice</td>
<td>4.7</td>
</tr>
<tr>
<td>There should be a report of the research results</td>
<td>4.6</td>
</tr>
<tr>
<td><strong>Less important</strong></td>
<td></td>
</tr>
<tr>
<td>I should have enough time</td>
<td>4.4</td>
</tr>
<tr>
<td>I should learn something (eg. a skill)</td>
<td>4.2</td>
</tr>
<tr>
<td>The project should be well adapted to the practice</td>
<td>4.0</td>
</tr>
<tr>
<td>There should be feedback on my own performance</td>
<td>4.0</td>
</tr>
<tr>
<td><strong>Least important</strong></td>
<td></td>
</tr>
<tr>
<td>Participation should contribute to my own career</td>
<td>3.2</td>
</tr>
<tr>
<td>I should have a say in the project</td>
<td>2.8</td>
</tr>
<tr>
<td>There should be financial reward</td>
<td>2.4</td>
</tr>
</tbody>
</table>

* 1 = very unimportant, 5 = very important

Table 2. Important factors in the recruitment of patients (2004, 31 respondents)

<table>
<thead>
<tr>
<th>Response to question 'Did the following reason play a role in the inclusion of patients?' in questionnaire</th>
<th>Numbers of residents (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Yes</td>
</tr>
<tr>
<td>Lack of patients who met inclusion criteria</td>
<td>18 (58)</td>
</tr>
<tr>
<td>Lack of time</td>
<td>16 (52)</td>
</tr>
<tr>
<td>Patients refusing to start the intervention*</td>
<td>10 (67)</td>
</tr>
<tr>
<td>Patients refusing to participate in research</td>
<td>12 (39)</td>
</tr>
<tr>
<td>Too many administrative actions</td>
<td>12 (39)</td>
</tr>
<tr>
<td>Difficulties in explaining the research</td>
<td>8 (26)</td>
</tr>
<tr>
<td>Difficulties leaving the role as a GP and asking patients for research participation</td>
<td>8 (26)</td>
</tr>
<tr>
<td>Lack of patients with emotional symptoms</td>
<td>3 (10)</td>
</tr>
</tbody>
</table>

* Intervention group (15 respondents)
observed variation in the registrars’ selection and recruitment of patients, which was related to their individual motivation as expressed during supervision sessions. For this reason the authors believe the actual effects of the change were limited.

Implications for general practice

Research experience during medical school is associated with postgraduate research involvement. Assuming that this applies to registrars too – and assuming a desire for research to be part of the culture of family medicine – the authors suggest that researchers and training programs should offer research in such a way that registrars will find it an attractive activity in which to participate.

This requires attention to the wishes of registrars and availability of resources, and the development of a culture that motivates registrars to prioritise research rather than a culture that views research as ‘unnecessary’. The way in which the training environment values research is an important factor in how registrars respond to participation. This environment includes both the training program, training practice and the profession at large. Faculty play an important role in this. With their expertise and experience, enthusiastic faculty could successfully integrate research into vocational training. Finally, registrars might be motivated by colleague peers, namely registrars actively involved in research such as those with an academic registrar position.

Creating attractive research programs should motivate registrars to voluntarily participate in research. Research networks, departments of family medicine, and residency training programs must collaborate to develop programs that offer registrars the opportunity to participate in distinct research projects. Whether this increases registrars’ participation in research activities is a question for further study.

Conflict of interest: none declared.

Acknowledgments

We thank ZonMw, the Netherlands organisation for health research and development, which funded our study, and we kindly thank all the registrars who participated.

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