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Non-randomised controlled trial

Improving stroke patients' care: a patient held record is not enough

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Abstract

Background: Stroke patients' care in hospital tends to be poorly organised, with poor communication and a lack of information being frequent sources of complaint. The purpose of this study was to evaluate whether a patient-held record (PHR) would result in greater patient satisfaction and better care planning for stroke patients.

Methods: A time series control (6 months) - intervention (8 months) - control (6 months) was used among London teaching hospital general medical and geriatric medicine inpatient wards. All stroke patients admitted to the wards during the intervention phase received a PHR and were instructed in its use. Demographic, stroke severity, social factors and outcomes were collected from all stroke patients during all phases of the study.

Results: Of 252 stroke patients aged 46 to 98 years entered into the study, by six months after admission 118 (46.8%) had died. PHR and control group patients were well matched in terms of socio-demographic characteristics and pre-stroke ability. At six months after admission, 119 (97%) patients responded to the questionnaire. Just over half (56%, 13) of intervention group patients recalled receiving a PHR. Of those patients, 59% reported reading the PHR, 27% had lost their PHR, and two-thirds said they had difficulties encouraging staff to write in the PHR. Half felt that possession of the PHR was more trouble than it was worth. PHR group patients were more satisfied with the recovery they had made (79% vs. 59%, $p=0.04$), but felt less able to talk to staff about their problems (61% vs. 82%, $p=0.02$). PHR group patients reported receiving fewer explanations about their condition (18% vs. 33%, $p=0.12$) and treatment (26% vs. 45%, $p=0.07$), and were more afraid of asking doctors questions (21% vs. 4%, $p=0.01$) than controls. PHR group patients were no better prepared for hospital discharge than control group patients, and both groups were ill-informed about services and benefits that might have helped after discharge from hospital.

Conclusions: Stroke patients received poor information and explanations regardless of whether they received a PHR. A PHR did not appear to improve patient satisfaction or discharge planning, and may have reduced opportunities for communication and explanation.

Introduction

Poor communication and lack of information are among the most common complaints of people with stroke [1,2,3]. Stroke services are poorly organised in many countries with a lack of continuity of care between hospital and community [4]. Consequently it is not surprising that dissatisfaction with services, especially those received after discharge, is common [5]. Patient held records (PHRs) exist in a variety of formats [6,7] and have been used in many different settings [8,9,10]. Individualised information booklets have been used with stroke patients before [11], but do not provide a continued record of care, their sole role being to provide information. In other areas of health care, patients are enthusiastic about holding their own records since they allow access to information, increase autonomy and bring about a shared feeling of responsibility for health [12,13,14]. PHRs have not been evaluated for use with stroke patients before.

A PHR might benefit stroke patients by providing information, promoting a more active role in their care and the possibility of contributing to decision making. As many people are involved in the care of stroke patients, possession of a PHR might give patients the confidence to ask questions of staff and they might receive more explanation. A PHR used by all members of the rehabilitation team, both in and out of hospital, could improve communication between team members. Consequently, our main hypothesis was that stroke patients with a PHR would be more satisfied with their care and would receive better planned care than those without a PHR.

Methods

A controlled comparison study with a control-intervention-control (A1-B-A2) time series design was used. Phase A1 was an initial six month control period, phase B was an eight month intervention with use of the PHR for all stroke patients, and phase A2 was the six-month period immediately after withdrawal of distribution of the PHR. Stroke patients admitted during phase A1 and A2 were combined and used as control patients for comparison with those admitted during phase B. This time-series design was used as a randomised controlled comparison of individual patients would not have been feasible within a single hospital because of contamination of the control group. A contemporary comparison with another hospital would not have provided any control over the amount or type of therapy and care received. Cluster randomisation of several hospitals would have been possible but would have increased the costs of an exploratory study to an unreasonable level.

Our study was planned to have a target sample size of 75 patients in the PHR group and 150 in the control group which would have given 80% power to detect a difference of 20% between PHR and control groups in specific areas of patient satisfaction. Stroke admission rates were lower during the PHR phase, so this was extended.

The patient held record was an A5 (metric) pocket-sized booklet with a hard cover and the patient's name on the front. The telephone numbers of all relevant staff were included and space was provided for assessment and management decisions to be recorded. Patients could also record their own comments. The PHR did not replace the existing case notes kept by each professional group. Therapists were asked to write in the PHR every time they saw a patient but nurses and doctors who had multiple daily contacts with patients were not expected to do this. Therapists were asked to record details of each of the main needs or problems they dealt with and then to outline any action taken and any immediate or longer term outcome, as previously reported [15].

Preparatory meetings were held with groups of staff to discuss their views on the value, design and content of the PHR. Prior to introduction of the PHR, interviews were held with therapists to explore their views on the use of the PHR [15]. Letters were sent to all medical and other staff who had contact with stroke patients informing them of the aims of the study and how the PHR should be used. The PHR was explained verbally to groups of nurses on each of the wards and repeated whenever it was felt necessary by the researchers.

Patients were given the PHR by a researcher (MA) who obtained informed consent to take part in the study and were encouraged to ask their therapists to write in the PHR and were asked to take the PHR home and use it with any health or social services staff they were in contact with after discharge. In cases where patients were too cognitively impaired to use the PHR themselves, these tasks were entrusted to a carer. If patients were too ill in the initial stages of their admission or without carers, the use of the PHR was delayed until patients were better able to understand. The PHRs were collected at six months after stroke or at death.

Patients admitted with new or recurrent strokes defined using clinical criteria (i.e. acute focal or global neurological disturbance of presumed vascular origin) were recruited via twice weekly visits to each acute admitting general medical and geriatric medicine ward in the hospital. Demographic data and pre-stroke levels of ability (Barthel activities of daily living index [16] and Nottingham extended activities of daily living index [17]) were recorded at initial assessment. All patients were followed

up at six months following admission to hospital using postal questionnaires including a previously validated patient satisfaction questionnaire [3], the Barthel index and questions about information received on diagnosis, prognosis, treatment and rehabilitation, their involvement in treatment and plans for discharge from hospital, attitudes to recovery and their opinions of the PHR.

Analyses were carried out using SPSS. Categorical variables were compared between groups using chi-square tests or Fisher's exact test for expected counts less than 5. Logistic regression analysis was used to adjust comparisons between groups for other baseline variables. The study was approved by the Royal Free Hospital Local Ethics Committee.

Results

During the twenty months of the study, 252 stroke patients were entered into the study, with 87 in the intervention phase B and the remaining 165 in the two control phases A1 and A2. At six-months following admission, 118 (47%) had died with 40% in phase B and 51% in the control phases ($p=0.20$), and 11 were lost to follow up (nine could not be traced and two had left the country).

Therefore, 123 survivors received questionnaires at six months of whom 119 (97%) responded. Complete data were available on Barthel index but patient satisfaction data was less complete owing to difficulties in patients with specific cognitive impairments or confusion who could not answer these questions.

The PHR and control groups were comparable in socio-demographic characteristics as shown in Table 1. The majority of patients had suffered their first stroke but just over half had some problems with basic activities of daily living pre-stroke as shown by their Barthel scores. At six months, functional ability was similar in both groups.

Use of the PHR

Although all patients in phase B received the PHR, only 56% (22 out of 39) reported they had ever had one. Of these, nine reported that they had not read their PHR, six had lost it, 14 had had difficulties persuading staff to write in it, and only six felt that the PHR had kept them informed about their treatment. Eleven of the patients thought that the PHR was more trouble than it was worth.

Table 1: Baseline Characteristics in Phase A (control) and Phase B (patient held record)

	Phase A	Phase B	P-value ⁺
Total number of subjects N	165	87	-
Mean age (range)	78 (53-93)	79 (46-98)	0.62*
Sex n (%)			
Male	61 (37.0)	38 (43.7)	
Female	104 (63.0)	49 (56.3)	0.30
Race n (%)			
White	151 (91.5)	72 (82.8)	
Non-white	14 (8.5)	15 (17.2)	0.04
Living conditions n (%)			
Alone	79 (47.9)	33 (37.9)	
Not alone/institutionalised	86 (52.1)	54 (62.1)	0.13
Last employment n (%)			
Manual	73 (58.9)	41 (52.6)	
Non-manual	51 (41.1)	37 (47.4)	0.38
Pre-stroke Barthel n (%)			
0-14	31 (18.8)	16 (18.4)	
15-19	55 (33.3)	30 (34.5)	
20	79 (47.9)	41 (47.1)	0.98
Pre-stroke extended ADL n (%)			
< 6	58 (35.2)	32 (36.8)	
6-17	54 (32.7)	29 (33.3)	
18-22	53 (32.1)	26 (29.9)	0.93
Ward on admission n(%)			
General medical	56 (33.9)	43 (49.4)	
Geriatric	109 (66.1)	44 (50.6)	0.02

⁺ Chi-squared or Fisher's Exact Test * T-test

Patient satisfaction

There were few differences in satisfaction between PHR and control groups (see Table 2). PHR group patients were happier with the recovery they had made (79% vs. 59%, $p=0.04$) but they were significantly less satisfied that they could talk to staff about problems (61% vs. 82%, $p=0.02$). Both groups reported high levels of dissatisfaction with community services received. Adjustment for age, sex, race, admission ward and disability at six months in a logistic regression on patient satisfaction questions made little difference to the odds ratios of satisfaction associated with the PHR.

Information recalled

More control than PHR group patients reported that someone had explained about the chances of recovery,

the effects of stroke on their lives, the reasons of investigations and the findings (see Table 3). Patients in both groups reported wanting more information from doctors than from any other professional group.

Involvement

Most patients (68% PHR vs. 61% control, $p=0.39$) were not involved in discussions about their treatment with doctors or therapists. Slightly more (61%) PHR group patients than control patients (49%) reported that they had had little idea what was happening to them while they were in hospital. The vast majority of patients in both groups felt able to ask questions of therapists and social workers. However, PHR group patients were significantly more afraid to ask questions of doctors than control group patients (21% vs. 4%, $p=0.01$).

Table 2: Differences in patient satisfaction in Phase A (control) and Phase B (patient held record)

	Phase A		Phase B		P-value	Differences in % 2 B-A (95% CI)	Adjusted odds ratios (95% CI)§
	N	Satisfied n (%)	N	Satisfied n (%)			
Hospital Care							
Treated with kindness	57	53 (93.0)	38	34 (89.5)	0.71*	-3.5 (-15.3,8.3)	0.6 (0.1,3.2)
Staff attended to needs	57	47 (82.5)	37	31 (83.8)	0.87	1.3 (-14.1,16.8)	1.1 (0.3,3.4)
Able to talk about problems	56	46 (82.1)	38	23 (60.5)	0.02	-21.6 (-40.1,-3.1)	0.3 (0.1,0.9)
Given information re illness	58	39 (67.2)	38	20 (52.6)	0.15	-14.6 (-34.6,5.3)	0.5 (0.2,1.1)
Doctors done all they can	57	49 (86.0)	38	35 (92.1)	0.52*	6.1 (-6.3,18.6)	2.0 (0.5,8.5)
Happy with recovery	58	34 (58.6)	38	30 (78.9)	0.04	20.3 (2.2,38.5)	2.9 (1.1,8.5)
Satisfied with type of therapy	52	46 (88.5)	31	28 (90.3)	1.00*	1.9 (-11.7,15.4)	1.5 (0.3,7.1)
I have had enough therapy	56	31 (54.4)	35	24 (68.8)	0.21	13.2 (-6.9,33.4)	1.9 (0.7,5.0)
Home Care							
Satisfied with amount of contact with hospital	46	38 (82.6)	28	22 (78.6)	0.67	-4.0 (-22.8,14.7)	0.7 (0.2,3.0)
Information wanted about allowances/benefits	46	28 (60.9)	23	17 (73.9)	0.28	13.0 (-9.9,35.9)	2.0 (0.6,7.1)
Good preparation for return home	42	39 (92.9)	21	20 (95.2)	1.00*	2.4 (-9.6,14.4)	†
Satisfied with service	58	17 (29.3)	39	10 (25.6)	0.69	-3.7 (-21.7,14.4)	0.8 (0.3,2.2)

* Fisher's Exact Test (Two-Tail), §adjusted for age, sex, race, admission ward, disability at 6 months, † not estimatable

Discharge planning

Almost three-quarters of patients in both groups were given adequate notice of when they would be going home. More PHR than control group patients felt ready to go home at the point of discharge (82% vs. 60%, $p=0.14$), but control group patients were more likely to have been instructed in how to cope at home (49% vs. 33%, $p=0.11$). Just over a third of patients had been told about services at home that might be helpful, and only a quarter had had welfare benefits explained to them.

Recovery

Significantly more PHR than control group patients felt that they had made a complete recovery from the stroke (38% vs. 14%, $p=0.01$), and a higher proportion felt that they had "got their lives back together again" (48% vs. 31%, $P=0.08$). Despite this, two-thirds of patients in both groups needed help with everyday activities. Both groups reported a positive outlook, attributing recovery to their own efforts and determination, and had not lost hope.

Table 3: Comparison of information received by patients in phase A (control) and phase B (patient held record)

	Phase A		Phase B		P-value	Differences in % B-A (95% CI)
	N	Yes n (%)	N	Yes n (%)		
Someone explained to me about:						
The causes of my stroke	58	21 (36.2)	38	12 (31.6)	0.64	-4.6 (-23.9, 14.6)
My chances of recovery	58	31 (53.4)	38	13 (34.2)	0.06	-19.2 (-39.0, 0.6)
The possible effects of the stroke on my life	58	19 (32.8)	38	7 (18.4)	0.12	-14.3 (-31.6, 2.9)
The reasons for the tests I had	58	26 (44.8)	38	10 (26.3)	0.07	-18.5 (-37.5, 0.5)
The results of the tests I had	58	23 (39.7)	38	9 (23.7)	0.10	-16.0 (-34.4, 2.5)
How being treated in hospital might help	58	20 (34.5)	38	6 (15.8)	0.04	-18.7 (-35.5, -1.8)
Someone explained to me about:						
The reasons for needing physiotherapy	48	36 (75.0)	28	19 (67.9)	0.51	-7.1 (28.3, 14.1)
Exercises to do on my own	50	36 (72.0)	27	20 (74.1)	0.84	2.1 (118.6, 22.8)
The use of aids (eg stick)	39	31 (79.5)	20	16 (80.0)	0.96	5.1 (-21.1, 22.1)
The reason for needing occupational therapy	47	30 (63.8)	25	13 (52.0)	0.33	-11.8 (-35.8, 12.1)
The use of aids (in eg your kitchen/bathroom)	45	23 (71.1)	23	18 (78.3)	0.53	7.1 (14.3, 28.6)
The reasons for need speech therapy	24	14 (58.3)	18	11 (61.1)	0.86	2.8 (-27.2, 23.7)
The use of communication aids	21	12 (57.1)	17	7 (41.2)	0.33	-16.0 (-47.5, 15.6)
Would have liked more information from:						
Doctors	57	23 (40.4)	39	15 (38.5)	0.85	-1.9 (-21.8, 18.0)
Nurses	57	13 (22.8)	39	11 (28.2)	0.55	5.4 (-12.4, 23.2)
Physiotherapists	48	12 (25.0)	28	10 (35.7)	0.32	10.7 (-10.9, 32.3)
Occupational therapists	43	8 (18.6)	26	10 (38.5)	0.07	19.9 (-2.2, 41.9)
Speech therapists	26	6 (23.1)	19	5 (26.3)	0.80	3.2 (22.3, 28.8)
Social workers	30	10 (33.3)	21	5 (23.8)	0.46	-9.5 (-34.4, 15.3)

Discussion

This is the first evaluation of a PHR, an intervention that has worked reasonably well in other clinical circumstances. Our hypothesis that the PHR would increase patient satisfaction is not supported by our findings. It is possible that the time series design was responsible for these disappointing findings. Non-randomised comparisons run the risk of introducing selection bias and fail to control for other potentially confounding factors that may bias assessment of outcomes. In this case, while our patient case mix was similar at baseline, the mortality rate was a little lower in the PHR intervention phase and more people were admitted to general medical wards. However, adjustment for age, sex, race, admission ward and Barthel index did not materially alter our findings. The inclusion of a second control phase following withdrawal of the intervention strengthens a time series design and permits any non-specific effects not associated with intervention to be evaluated. As with any non-randomised comparison, our findings must be viewed with some caution.

There are several possible explanations for our negative findings. As death rates were higher than anticipated, our effective sample size for most comparisons was re-

duced but was still sufficiently powerful to detect clinically important differences of about 30% in satisfaction. It is possible that PHRs were given to patients who were simply too sick to use them, but we attempted to enter all available patients into the study to assess the usefulness of the PHR for all stroke patients. Our findings tended to show that, if anything, patients who received a PHR did rather worse for most outcomes than those who did not. Why should this be? Disinterest, as has been found elsewhere [18], may be a reason why so many patients did not know they had a PHR or did not bother to read it. Some difficulties in its use may be attributed to the high turnover of staff and routine rotation of therapists, but most therapy staff had been involved in the design and inception of the PHR and certainly wanted it to work. Although all of the medical consultants managing stroke patients agreed for their patients to be studied, none of them acted as "champions" for the innovation. This lack of medical involvement may have contributed to the disappointing findings and limited use of the PHR.

That patients would lose their PHR was anticipated by therapists at the start of the study [15], and reflects the negative views of patients' ability among health professionals [8,12,13,18]. Women who hold their antenatal

records rarely lose them [7,12,19], but pregnant women are not ill. We wanted to challenge the expectation that older, disabled patients would be unable to take responsibility by evaluating the PHR among patients who generally lack power [20]. But perhaps these patients were simply too ill - as suggested by the high mortality rate - to make use of a PHR.

Our findings contrast with those which suggest that patients with a PHR experience less difficulty talking to staff and feel empowered [10,21]. It is possible that in the PHR group communication problems were *caused* by the PHR. It may have heightened their expectations and hopes which do not appear to have been fully met. Discussing a poor prognosis is an extremely difficult task for health professionals [22]. Making written entries in a PHR may have increased this difficulty as it is easier to be vague verbally than in writing [15] and this may have deterred staff from writing in the PHR.

Conclusion

We expected the PHR to bridge the gap between hospital and the community, contributing to the continuity of care that has been found elsewhere [10]. This did not occur and our findings highlight the generally high levels of dissatisfaction with services received after leaving hospital. Introducing a PHR into stroke care, an area of health care recognised to be haphazard and fragmented [5], was an ambitious undertaking. Changing the well-established culture and practice of stroke patient care will take more powerful interventions than a PHR. PHRs have been shown to be effective in some clinical settings but such findings are not applicable to the more difficult circumstances of the acute hospital care of frail and often confused patients. This study has drawn attention to the generally poor levels of information and explanation given to patients and their lack of involvement in their care, regardless of whether they had a PHR.

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