Should we provide outreach rehabilitation to very old people living in Nursing Care Facilities after a hip fracture? A randomised controlled trial

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Abstract

Objective: to determine whether a 4-week postoperative rehabilitation program delivered in Nursing Care Facilities (NCFs) would improve quality of life and mobility compared with receiving usual care.

Design: parallel randomised controlled trial with integrated health economic study.

Should we provide outreach hip fracture rehabilitation to care homes

Age and Ageing 2019; 48: 373–380
doi: 10.1093/ageing/afz005
Published electronically 22 February 2019

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Setting: NCFs, in Adelaide South Australia.

Subjects: people aged 70 years and older who were recovering from hip fracture surgery and were walking prior to hip fracture.

Measurements: primary outcomes: mobility (Nursing Home Life-Space Diameter (NHLSD)) and quality of life (DEMQOL) at 4 weeks and 12 months.

Results: participants were randomised to treatment (n = 121) or control (n = 119) groups. At 4 weeks, the treatment group had better mobility (NHLSD mean difference −1.9; 95% CI: −3.3, −0.57; P = 0.0055) and were more likely to be alive (log rank test P = 0.048) but there were no differences in quality of life. At 12 months, the treatment group had better quality of life (DEMQOL sum score mean difference = −7.4; 95% CI: −12.5 to −2.3; P = 0.0051), but there were no other differences between treatment and control groups. Quality adjusted life years (QALYs) gained over 12 months were 0.0063 higher per participant (95% CI: −0.0547 to 0.0686). The resulting incremental cost effectiveness ratios (ICERs) were $5,545 Australian dollars per unit increase in the NHLSD (95% CI: $244 to $15,159) and $328,685 per QALY gained (95% CI: $82,654 to $75,007,056).

Conclusions: the benefits did not persist once the rehabilitation program ended but quality of life at 12 months in survivors was slightly higher. The case for funding outreach home rehabilitation in NCFs is weak from a traditional health economic perspective.

Trial registration: ACTRN12612000112864 registered on the Australian and New Zealand Clinical Trials Registry. Trial protocol available at https://www.anzctr.org.au/Trial/Registration/TrialReview.aspx?id = 361980

Keywords
hip fracture, rehabilitation, aged care, mobility, quality of life, older people

Key points
• A 4-week multidisciplinary postoperative rehabilitation program after hip fracture surgery conducted in nursing care facilities was associated with better mobility and survival at 4 weeks compared with usual care.
• The benefits did not persist once the rehabilitation program ended but a small gain in quality of life at 12 months in survivors was seen.
• The overall mortality rate was 46% at 12 months.
• The outreach rehabilitation program could not be considered cost-effective against current public funding thresholds.
• Future trials should explore different approaches to postoperative hip fracture recovery in this group, such as nursing care facility based rehabilitation approaches.

Background
Hip fractures are a common cause of suffering for residents of nursing care facilities (NCFs) and outcomes are poor [1]. Most residents have dementia and are frail. In a retrospective cohort study of 60,111 US Medicare beneficiaries living in nursing homes, only one in five patients who had been fully independent or required limited supervision/assistance walking at baseline survived to regain their pre-fracture level of walking 180 days after fracture [2].

Guidelines for hip fracture management promote prompt surgery, early mobilisation, and a team-based rehabilitation approach to restoring function and mobility [3]. The high risk of death and adverse outcomes means there is uncertainty about the benefits of health service resources allocated to rehabilitation in people living in NCFs [3]. We investigated the feasibility of providing a Comprehensive Geriatric Assessment and interdisciplinary rehabilitation program which was developed according to clinical guidelines [4, 5]. The aim of the study was to examine if a rehabilitation program in NCFs for people who were recovering from hip fracture surgery improved quality of life and mobility at 4 weeks and 12 months.

Methods/design
See Supplementary material, available at Age and Ageing online for the CONSORT checklist and protocol. The study was approved by the Southern Adelaide Clinical Human Research Ethics Committee. A randomised controlled trial with masked outcome assessments was undertaken between June 2012 and December 2014. A computer generated random sequence with random block sizes was used by a pharmacist external to the project to allocate people with a hip fracture, who had been treated surgically into: (a) 4-week ambulatory geriatric rehabilitation program (delivered in the NCF) (b) usual care. Recruitment occurred
on acute orthopaedic wards at three South Australian Hospitals.

**Participant procedures**

Participants were randomised in hospital, the intervention commenced within 24 h of return to the NCF, and on return all residents received usual medical care from their general practitioner. All hospitals had an Orthogeriatrics service that reviewed patients prior to discharge. Inclusion criteria were: a recent hip fracture treated surgically, aged 70 years or older, able to follow a one-step command, living in an NCF prior to injury, ambulant prior to fracture, and ready for discharge, providing self or proxy informed consent (full criteria listed in Supplementary material, available at Age and Ageing online).

Those allocated to the intervention received visits from a hospital outreach team who provided a Comprehensive Geriatrics Assessment, physiotherapy and nutritional assessment and care plan. Physiotherapy included mobility and task specific training, graduated muscle strengthening exercises and training of care staff and family. The geriatrician met families within a fortnight to discuss progress. The intervention was low intensity and involved 13 h of input.

**Measurements/procedures**

**Primary outcomes**

The primary outcomes were mobility autonomy (measured using the Nursing Home Life-Space Diameter (NHLSD)) and Quality of Life. The NHLSD has high intra-rater (0.922) and inter-rater (0.951) reliability and moderate positive correlation with other functional characteristics (e.g. social activity participation, dressing) [6]. It consists of four diameters scored on a scale of 0–5 and weighted, with possible scores ranging from 0 (bed- or chair-bound) to 120 (signifying leaving the facility daily). Care staff were asked to describe the level of independence and hands-on support each participant was receiving at baseline, 4 weeks and 12 months.

Quality of life was assessed with the 28-item DEMQOL and 31-item DEMQOL-Proxy which are condition specific measures designed to measure health-related quality of life for older people with dementia and their carers [7]. At baseline 90 participants completed the DEMQOL and 237 were completed by proxies (in 83 cases both an individual and proxy questionnaires were completed). At 4 weeks, the DEMQOL-Proxy was completed for 199 participants. The overall correlation between scores of the self-completed questionnaires and proxy questionnaires was poor at baseline ($r = 0.27538$, $P = 0.0117$) with proxies tending to score quality of life lower than individuals suggesting that different constructs were being measured. Where two questionnaires were available the DEMQOL-Proxy was used. The EuroQol five dimension–five level questionnaire (EQ-5D-5L) was administered to compare participants’ quality of life with other patient groups internationally [8].

**Secondary outcomes**

Physical dependency was measured by the Modified Barthel Index [9] and the Functional Recovery Scale [10]. Other measures included cognition (Mini-Mental State Examination: MMSE) [11], confusion or delirium (Confusion Assessment Method) [12], depression (Cornell Scale for Depression in Dementia) [13], pain (the Pain Assessment In Advanced Dementia scale: PAINAD) [14] and nutrition (The Mini-Nutritional Assessment) [15].

**Statistical analysis**

To assess minimally important differences in the DEMQOL index score, we needed 98 per group (intervention and control). After allowing for deaths and drop-outs of 20% the estimated sample size was $196 \times 1.2 = 236$ (118 per group). The detectable effect size between groups was conservatively selected as small to medium (0.10–0.25) as suggested by Cohen [16]. Calculations were based on two-tailed tests with power of at least 80% and significance level of 0.05.

Outcomes were evaluated using linear mixed models with a time-by-group interaction term. The covariates were group, time, time*group and baseline scores for the outcome variables.

To investigate survival from the randomisation to 4 weeks and 12 months, we used Kaplan–Meier and log rank test to test the between group difference. All data were analysed according to the intention-to-treat principle and performed with SAS, v9.3 (SAS institute) and R 3.11.

To assess the incremental cost-effectiveness of the intervention compared with usual care we examined incremental cost-effectiveness per unit increase in the NHLSD total score over 1-year follow-up. Utility-based outcomes were incorporated into the analysis, to generate a secondary outcome: incremental cost per Quality adjusted life year (QALY; based on DEMQOL-Proxy values). QALYs were calculated using the area-under-the-curve [17]. Cost effectiveness acceptability curves (CEACs) were constructed, depicting the probability of the intervention being more cost-effective compared with the usual care arm at different willingness-to-pay thresholds (see Supplementary Figures S4 and S5, available at Age and Ageing online) [18]. Further details on the cost effectiveness analysis are provided in the supplementary information.

**Results**

At the three participating hospitals 2,120 hip fracture patients were screened, 354 were eligible and following consent 240 participated (see Supplementary Figure S1, available at Age and Ageing online). In the majority of cases (97%) consent was obtained from family members due to cognitive impairment. Demographic and clinical characteristics were well balanced between groups (Table 1). The mean age was 88.6 years (SD: 5.6) and 13% had a prior hip fracture. The majority (87.9%) received surgical treatment within the first 24 h of admission (range: 0–5 days). Baseline pain (PAINAD) scored at rest was low 1.4 (SD: 1.7), only 23
Table 1. Baseline characteristics of study population

| Characteristic* | Control | Intervention |
|-----------------|---------|--------------|
|                 | N1 = 121 | N2 = 119    |
| Female sex—n (%) | 91 (75.2) | 87 (73.1) |
| Age—n (%) (range: 70–101) |          |              |
| 70–79            | 8 (6.6)  | 8 (6.7)     |
| 80–89            | 56 (46.3)| 62 (52.1)  |
| 90–95            | 44 (36.4)| 38 (31.9)  |
| >95              | 13 (10.7)| 11 (9.2)   |
| Age-mean (SD)    | 88.6 (5.7) | 88.6 (5.4) |
| Mini-Mental State Examination—mean (SD) | 8.5 (7.6) | 7.5 (8.0) |
| Medication Appropriateness Index—mean (SD) | 2.5 (1.9) | 2.3 (1.8) |
| Delirium—n (%)   | 41 (33.9) | 42 (35.3)  |
| Previous any fractures—n (%) | 47 (38.8) | 47 (39.5) |
| Previous hip fractures—n (%) | 16 (13.1) | 16 (13.6) |
| Type of hip of fracture (at baseline) |          |              |
| Extracapsular   | 58 (47.9) | 52 (43.7)  |
| Intracapsular   | 63 (52.1) | 67 (56.3)  |
| Extracapsular hip fracture-surgery type at baseline (n = 110) |          |              |
| Sliding hip screw | 8 (13.8)  | 15 (28.8)  |
| Intramedullary nail | 50 (86.2) | 37 (71.2)  |
| Intracapsular hip fracture-surgery type at baseline (n = 130) |          |              |
| Internal fixation | 18 (28.6) | 15 (22.4)  |
| Cemented Hemiarthroplasty | 28 (44.4) | 36 (53.7)  |
| Uncemented Hemiarthroplasty | 15 (23.8) | 15 (22.4)  |
| Total hip replacement | 2 (3.2) | 1 (1.5)    |
| Pre-fracture Mobility Aid indoor |          |              |
| None            | 20 (16.5) | 26 (21.9)  |
| Walking stick   | 5 (4.1)   | 2 (1.7)    |
| Walking frame   | 96 (79.3) | 89 (74.8)  |
| Personal assistance | 0 (0.0) | 2 (1.7)    |
| Pre-fracture Mobility Aid Outdoor |          |              |
| None            | 15 (12.4) | 23 (19.3)  |
| Walking stick   | 4 (3.3)   | 1 (0.8)    |
| Walking frame   | 76 (62.8) | 67 (56.3)  |
|Personal assistance | 1 (0.8) | 3 (2.5)    |
| Unable          | 25 (20.6)| 25 (21.0)  |
| Pre-fracture Mobility Assistance—Indoor |          |              |
| Independent     | 82 (67.8) | 73 (61.3)  |
| 1 x LA          | 16 (13.2) | 19 (16.0)  |
| 1 x MA          | 4 (3.3)   | 7 (5.9)    |
| 1 x S/B         | 19 (15.7) | 20 (16.8)  |
| Pre-fracture Mobility Assistance—Outdoor |          |              |
| Independent     | 41 (33.9) | 33 (27.7)  |
| 1 x LA          | 17 (14.1) | 18 (15.1)  |
| 1 x MA          | 4 (3.3)   | 10 (8.0)   |
| 1 x S/B         | 37 (30.6) | 35 (29.4)  |
| 2 x LA          | 1 (0.8)   | 6 (5.0)    |
| Unable          | 21 (17.4)| 26 (21.9)  |

*There was no significant difference (P < 0.05) between control and intervention groups for all above variables at baseline. Data are mean (SD) or n (%).

At 4 weeks, the treatment group achieved a better mobility score (NHLSD mean difference −1.9; 95% CI: −3.3 to −0.57; P = 0.0055) (Table 2). The treatment group also had better nutritional status than the control group (−0.65; 95% CI: −1.3, −0.05; P = 0.0338).

At 12-month follow-up, the treatment group had better quality of life as measured by DEMQOL sum scores (mean difference = −7.4; 95% CI: −12.5, −2.3; P = 0.0051). There were no other statistically significant differences between treatment and control groups.

At 4 weeks, the death rate was 8% in the intervention group and 18% in the control group (log rank test by the end of 4 weeks P = 0.048), and in the control group the number of deaths increased each week from one death (Week 1) to eight deaths (Week 4) (Figure 1). However, after the rehabilitation program, there was an increase in deaths in the intervention group. After 35 days, there was no statistically significant difference between groups in the probability of survival (Figure 1).

Adverse events

In total, 95 nursing home residents sustained one or more falls during the 4-week intervention with 56 people from the intervention group incurring 62.7% (n = 162) of the falls. Twelve people had hospital admissions including three hip fractures. In the usual care group, 39 people fell with 15 (38.5%) requiring hospital admission and one person sustained a hip fracture (see Supplementary material, available at Age and Ageing online).

Economic evaluation

Mean per participant 12-month Australian Medicare costs were higher in the intervention group than in the control arm (by $2,076 per patient) but these differences were not statistically significant (95% CI: $−220 to $4,360). Drivers of higher costs in the intervention were the intervention cost itself and higher drug costs. When the adjusted 12-month primary and secondary outcomes in the base case were considered (Supplementary material), the intervention was more effective than the control with participants reporting NHLSD totals scores that were higher by 0.3745 per participant (95% CI: 0.0063 to 0.742) and QALYs gained that were higher by 0.0063 per patient (95% CI: 0.0049 to 0.0076). The resulting incremental cost effectiveness ratios (ICERs) were $5,545 per unit increase in the NHLSD total score (95% CI: $244−15,159) and $328,685 per QALY gained (95% CI: $82,654−750,075). The ICER based on QALYs is substantially greater than the implicit cost-effectiveness threshold of $50,000 per QALY gained currently applied by regulatory bodies in Australia [19], implying that the intervention would not be considered cost-effective [17].
A 4-week multidisciplinary home rehabilitation program reduced mortality and improved mobility and nutritional status in people living in NCFs who had previously been walking but then fractured their hips. However, improvements were not sustained at 12 months. At 12 months, there was a small quality of life improvement in survivors.

The higher health costs associated with improved mobility in the intervention group were modest. However, the 12-month cost effectiveness estimates are prohibitively high at $5,545 per unit improvement in the NHLSD total scores and $328,685 per QALY gained. Estimates of the incremental costs per QALY gained from this trial are much higher than the recommended threshold of $50,000/QALY used in Australia suggesting that providing outreach

### Table 2. Baseline and four week data for primary and secondary outcomes

| Outcomes | Control | Intervention | Difference (95% CI) | P value |
|----------|---------|--------------|---------------------|---------|
|          | n       | Mean (SE)    | n                   | Mean (SE) |
| Primary outcomes |       |              |                     |         |
| Baseline | NHLSD   | 121 0 (0)    | 119 0 (0)           | _       | _ |
| Quality of life | DEMQOL sum score | 50 86.5 (1.2) | 40 86.2 (4.4) | 0.30 (−3.4, 4.0) | 0.8711 |
|               | DEMQOL index (utility) | 50 0.80 (0.04) | 40 0.79 (0.04) | 0.01 (−0.11, 0.13) | 0.8587 |
|               | DEMQOL-proxy sum score | 119 90.9 (1.0) | 118 92.1 (1.0) | −1.1 (−3.8, 1.6) | 0.4141 |
|               | DEMQOL-proxy index (utility) | 119 0.62 (0.02) | 118 0.54 (0.02) | −0.01 (−0.08, 0.06) | 0.7111 |
|               | EQ5D index (utility) | 119 0.22 (0.02) | 119 0.23 (0.02) | −0.01 (−0.08, 0.06) | 0.7788 |
| 4 Weeks     | NHLSD   | 96 6.3 (0.50) | 107 8.2 (0.47) | −1.9 (−3.3, −0.57) | 0.0055 |
| Quality of life | DEMQOL sum score | 45 88.3 (1.6) | 49 91.0 (1.7) | −2.7 (−7.3, 1.8) | 0.2370 |
|               | DEMQOL index | 67 0.68 (0.05) | 59 0.74 (0.04) | −0.06 (−0.20, 0.08) | 0.3896 |
|               | DEMQOL-proxy sum score | 94 93.7 (1.1) | 105 94.2 (1.0) | −0.52 (−3.5, 2.4) | 0.7305 |
|               | DEMQOL-proxy index | 116 0.54 (0.02) | 115 0.60 (0.02) | −0.06 (−0.11, 0.01) | 0.0784 |
|               | EQ5D index | 118 0.38 (0.02) | 115 0.43 (0.02) | −0.05 (−0.12, 0.01) | 0.1058 |
| 12 Months   | NHLSD   | 66 10.1 (0.69) | 60 10.5 (0.63) | 0.37 (−2.1, 1.3) | 0.6777 |
| Quality of life | DEMQOL sum score | 41 88.5 (1.6) | 29 95.9 (2.0) | −7.4 (−12.5, −2.3) | 0.0051 |
|               | DEMQOL index | 93 0.54 (0.03) | 87 0.48 (0.04) | 0.06 (−0.07, 0.19) | 0.3521 |
|               | DEMQOL-proxy sum score | 66 101.9 (1.3) | 60 98.7 (1.4) | 3.1 (−0.62, 6.9) | 0.1023 |
|               | DEMQOL-proxy index | 118 0.30 (0.02) | 117 0.24 (0.02) | 0.06 (−0.006, 0.13) | 0.0739 |
| Secondary outcomes |               |              |                     |         |
| Baseline    | PAINAD   | 121 1.4 (0.11) | 119 1.4 (0.11) | 0.00 (−0.31, 0.31) | 0.9824 |
|             | Modified Barthel Index | 121 9.6 (1.8) | 119 9.5 (1.8) | 0.08 (−4.9, 5.1) | 0.9735 |
|             | Cornell Scale for Depression in Dementia | 119 10.1 (0.42) | 119 10.0 (0.42) | 0.01 (−1.2, 1.2) | 0.9857 |
|             | Mini-Nutritional Assessment | 121 5.4 (0.20) | 119 5.3 (0.20) | 0.12 (−0.43, 0.68) | 0.6670 |
|             | Functional recovery | 121 1.8 (0.36) | 119 1.8 (0.36) | 0.01 (−0.98, 1.01) | 0.9798 |
|             | Delirium | 121 0.34 (0.04)a | 119 0.35 (0.04)a | 0.94 (0.55, 1.6)b | 0.8184 |
| 4 Weeks     | PAINAD   | 95 0.49 (0.13) | 107 0.51 (0.12) | −0.02 (−0.34, 0.29) | 0.8998 |
|             | Modified Barthel Index | 95 23.5 (2.0) | 107 24.4 (1.9) | −1.0 (−6.4, 4.5) | 0.7267 |
|             | Cornell Scale for Depression in Dementia | 96 10.6 (0.47) | 107 10.5 (0.44) | 0.15 (−1.1, 1.4) | 0.8097 |
|             | Mini-Nutritional Assessment | 96 6.2 (0.22) | 107 6.9 (0.21) | −0.65 (−1.3, −0.05) | 0.0338 |
|             | Functional recovery | 94 5.8 (0.40) | 107 6.0 (0.38) | −0.25 (−1.3, 0.84) | 0.6542 |
|             | Delirium | 95 0.13 (0.03) | 107 0.17 (0.04) | 0.75 (0.35, 1.6) | 0.4589 |
| 12 Months   | PAINAD   | 66 0.06 (0.15) | 60 0.05 (0.16) | 0.01 (−0.42, 0.44) | 0.9645 |
|             | Modified Barthel Index | 66 32.3 (2.4) | 59 27.4 (2.5) | 5.0 (−1.9, 11.8) | 0.1533 |
|             | Cornell Scale for Depression in Dementia | 66 9.3 (0.56) | 60 10.1 (0.59) | −0.8 (−2.4, 0.8) | 0.3262 |
|             | Mini-Nutritional Assessment | 66 8.0 (0.27) | 60 8.8 (0.28) | −0.73 (−1.5, 0.03) | 0.0592 |
|             | Functional recovery | 66 7.1 (0.48) | 60 6.2 (0.50) | 0.84 (−0.52, 2.2) | 0.2257 |
|             | Delirium | 66 0.18 (0.05)a | 60 0.22 (0.05)a | 0.77 (0.33, 1.8)b | 0.5486 |

*aPercentage.

A 4-week multidisciplinary home rehabilitation program reduced mortality and improved mobility and nutritional status in people living in NCFs who had previously been walking but then fractured their hips. However, improvements were not sustained at 12 months. At 12 months, there was a small quality of life improvement in survivors.

**Discussion**

The higher health costs associated with improved mobility in the intervention group were modest. However, the 12-month cost effectiveness estimates are prohibitively high at $5,545 per unit improvement in the NHLSD total scores and $328,685 per QALY gained. Estimates of the incremental costs per QALY gained from this trial are much higher than the recommended threshold of $50,000/QALY used in Australia suggesting that providing outreach

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rehabilitation is not likely to be value for money for a health service. One option to improve the cost effectiveness estimate would be to decrease the rehabilitation costs and extend the period of additional therapy by exploring models of rehabilitation where NCF staff are trained to deliver therapy for longer periods of time. However, for frail older people living in nursing homes where death is a common event and quality of life gains are modest, results of cost effectiveness approaches are unlikely to be favourable and decisions on allocation of resources to this group may need to include consideration of a community’s values. After this trial, a citizens’ jury process was conducted with randomly selected citizens which suggested that the community regards access to recovery or rehabilitation services for nursing home patients as a human rights issue and despite access to recovery or rehabilitation services for selected citizens which suggested that the community measures in people without dementia is unclear. The current study not only demonstrates the challenges of working with very old people who have high mortality rates but also the difficulties in assessing effective treatments and improvements in quality of life.

Conclusions

A rehabilitation program for older people living in NCFs after hip fracture surgery who were mobile pre-surgery showed improved mobility, nutritional status and survival compared to usual care at 4 weeks. These improvements did not persist at one year but there were small quality of life gains at 12 months in the survivors. The outreach rehabilitation program was not cost-effective. Further studies could investigate whether a longer-term or NCF-based rehabilitation approach following hip fracture is cost-effective.
Supplementary data mentioned in the text are available to subscribers in Age and Ageing online.

Declaration of Conflict of Interest: M Crotty had completed two previous clinical drug trials on community dwelling hip fracture patients: (1) Novartis (2016–2017) trial to evaluate iv bimagrumab on total lean body mass and physical performance in patients after surgical treatment of hip fracture and (2) Eli Lilly STEADY trial to investigate subcutaneous injections of LY2495655 in older patients who have fallen recently and have muscle weakness. M Chehade has received institutional grants from Stryker to support hip fracture research, but it does not pose a relevant conflict to this study. All other authors declare no competing interests.

Declaration of Sources of Funding: This study was supported by funding provided by the National Health and Medical Research Council (NHMRC) Partnership Centre on Dealing with Cognitive and Related Functional Decline in Older People (grant no. GNT9100000). The contents of the published materials are solely the responsibility of the Administering Institution, Flinders University, and the individual authors identified, and do not reflect the views of the NHMRC or any other Funding Bodies or the Funding Partners.

References

1. Braithwaite RS, Col NF, Wong JB. Estimating hip fracture morbidity, mortality and costs. J Am Geriatr Soc 2003; 51: 364–70.
2. Neuman MD, Silber JK, Magaziner JS, Passarella MA, Mehta S, Werner RM. Survival and functional outcomes after hip fracture among nursing home residents. JAMA Intern Med 2014; 174: 1273–80.
3. National Institute of Clinical Excellence (NICE). The management of hip fracture in adults. London 2011; Available from: http://www.nice.org.uk/nicemedia/live/11968/51532/51532.pdf.
4. Crocker T, Forster A, Young J et al. Physical rehabilitation for older people in long-term care. Cochrane Database Syst Rev 2013; 28: CD004294.
5. Australian and New Zealand Hip Fracture Registry (ANZHFR) Steering Group. Australian and New Zealand Guideline for Hip Fracture Care: Improving Outcomes in Hip Fracture Management of Adults. Sydney: Australian and New Zealand Hip Fracture Registry Steering Group, 2014.
6. Tinetti ME, Ginter SF. The nursing home life-space diameter. A measure of extent and frequency of mobility among nursing home residents. J Am Geriatr Soc 1990; 38: 1311–5.
7. Mulhern B, Rowen D, Braxier J et al. Development of DEMQOL-U and DEMQOL-PROXY-U: generation of preference-based indices from DEMQOL AND DEMQOL-PROXY for use in economic evaluation. Health Technol Assess 2013; 17: 1–140.
8. Herdman M, Gudex C, Lloyd A et al. Development and preliminary testing of the new five-level version of EQ-5D (EQ-5D-5L). Qual Life Res 2011; 20: 1727–36.
9. Shah S, Vanclay F, Cooper B. Improving the sensitivity of the Barthel Index for stroke rehabilitation. J Clin Epidemiol 1989; 42: 703–9.
10. Koval KJ, Zuckerman JD. Functional recovery after fracture of the hip. J Bone Joint Surg Am 1994; 76: 751–8.
11. Folstein MF, Folstein SE, McHugh PR. 'Mini-mental state'. A practical method for grading the cognitive state of patients for the clinician. J Psychiatr Res 1975; 12: 189–98.
12. Inouye SK, van Dyck CH, Alessi CA, Balkin S, Siegal AP, Horwitz RI. Clarifying confusion: the confusion assessment method. A new method for detection of delirium. Ann Intern Med 1990; 113: 941–8.
13. Alexopoulos G, Abrams R, Young R, Shamolani C. Cornell scale for depression in dementia. Biol Psychiatry 1988; 23: 271–84.
14. Wärdén V, Harley AC, Volcér L. Development and psychometric evaluation of the Pain Assessment in Advanced Dementia (PAINAD) scale. J Am Med Dir Assoc 2003; 4: 9–15.
15. Nourhashemi F, Guyonnets S, Ousset PJ et al. Mini nutritional assessment and Alzheimer patients. Nestle Nutr Workshop Ser Clin Perform Programme 1999; 1: 87–91.
16. Cohen J. A power primer. Psychol Bull 1992; 112: 115–59.
17. Drummond MF, Sculpher M, O’Brien B, Stoddart GL, Torrance GW. Methods for the Economic Evaluation of Health Care Programmes. Oxford: Oxford University Press, 2005.
18. Krichbaum K. GAPN postacute care coordination improves hip fracture outcomes. West J Nurs Res 2007; 29: 523–44.
19. Harris AH, Hill SR, Chin G, Li JJ, Walkom E. The role of value for money in public insurance coverage decisions for drugs in Australia: a retrospective analysis 1994-2004, Med Decis Making 2008; 28: 713–22.
20. Laver K, Gnanamanickam E, Ratcliffe J et al. A citizens jury to inform policy on rehabilitation for people in residential care with hip fracture. Innov Aging 2017; 1: 226.
21. Chua KC, Brown A, Little R et al. Quality-of-life assessment in dementia: the use of DEMQOL and DEMQOL-Proxy total scores. Qual Life Res 2016; 25: 3107–18.
22. Hounsome N, Orrell M, Edwards RT. EQ-5D as a quality of life measure in people with dementia and their carers: evidence and key issues. Value Health 2011; 14: 390–9.
23. Beaupre LA, Binder EF, Cameron ID et al. Maximising functional recovery following hip fracture in frail seniors. Best Pract Res Clin Rheumatol 2013; 27: 771–88.
24. Devlin NJ, Shah KK, Feng Y, Mulhern B, van Hout B. Valuing health-related quality of life: an EQ-5D-5L value set for England. Health Econ 2018; 27: 7–22.
25. Rowen D, Mulhern B, Banerjee S et al. Estimating preference-based single index measures for dementia using DEMQOL and DEMQOL-Proxy. Value Health 2012; 15: 436–56.
26. Olsen JA, Lamu AN, Cairns J. In search of a common currency: a comparison of seven EQ-5D-5L value sets. Health Econ 2018; 27: 39–49.
27. Trigg R, Jones RW, Knapp M, King D, Lacey IA. The relationship between changes in quality of life outcomes and progression of Alzheimer’s disease: results from the dependence in AD in England 2 longitudinal study. Int J Geriatr Psychiatry 2015; 30: 400–8.
28. Coucill W, Bryan S, Bentham P, Buckley A, Laight A. EQ-5D in patients with dementia: an investigation of inter-rater agreement. Med Care 2001; 39: 760–71.
Association of blood pressure with clinical outcomes in older adults with chronic kidney disease

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Abstract

Background: in chronic kidney disease (CKD), hypertension is associated with poor outcomes at ages <70 years. At older ages, this association is unclear. We tested 10-year mortality and cardiovascular outcomes by clinical systolic blood pressure (SBP) in older CKD Stages 3 and 4 patients without diabetes or proteinuria.

Methods: retrospective cohort in population representative primary care electronic medical records linked to hospital data from the UK, CKD staged by CKD-EPI equation (≥2 creatinine measurements ≥90 days apart). SBPs were 3-year medians before baseline, with mean follow-up 5.7 years. Cox competing models accounted for mortality.

Results: about 158,713 subjects with CKD3 and 6,611 with CKD4 met inclusion criteria. Mortality increased with increasing CKD stage in all subjects aged >60. In the 70 plus group with SBPs 140–169 mmHg, there was no increase in mortality, versus SBP 130–139. Similarly, SBPs 140–169 mmHg were not associated with increased incident heart failure, stroke or myocardial infarctions. SBPs <120 mmHg were associated with increased mortality and cardiovascular risk. At ages 60–69, there was increased mortality at SBP <120 and SBP >150 mmHg. Results were little altered after excluding those with declining SBPs during 5 years before baseline, or for longer-term outcomes (5–10 years after baseline).

Conclusions: in older primary care patients, CKD3 or 4 was the dominant outcome predictor. SBP 140–169 mmHg having little additional predictive value, <120 mmHg was associated with increased mortality. Prospective studies of representative older adults with CKD are required to establish optimum BP targets.

Keywords

blood pressure, chronic kidney disease, older people, cardiovascular outcomes