

BMJ Open Continuity of care with doctors – a matter of life and death? A systematic review of continuity of care and mortality

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ABSTRACT

Objective Continuity of care is a long-standing feature of healthcare, especially of general practice. It is associated with increased patient satisfaction, increased take-up of health promotion, greater adherence to medical advice and decreased use of hospital services. This review aims to examine whether there is a relationship between the receipt of continuity of doctor care and mortality.

Design Systematic review without meta-analysis.

Data sources MEDLINE, Embase and the Web of Science, from 1996 to 2017.

Eligibility criteria for selecting studies Peer-reviewed primary research articles, published in English which reported measured continuity of care received by patients from any kind of doctor, in any setting, in any country, related to measured mortality of those patients.

Results Of the 726 articles identified in searches, 22 fulfilled the eligibility criteria. The studies were all cohort or cross-sectional and most adjusted for multiple potential confounding factors. These studies came from nine countries with very different cultures and health systems. We found such heterogeneity of continuity and mortality measurement methods and time frames that it was not possible to combine the results of studies. However, 18 (81.8%) high-quality studies reported statistically significant reductions in mortality, with increased continuity of care. 16 of these were with all-cause mortality. Three others showed no association and one demonstrated mixed results. These significant protective effects occurred with both generalist and specialist doctors.

Conclusions This first systematic review reveals that increased continuity of care by doctors is associated with lower mortality rates. Although all the evidence is observational, patients across cultural boundaries appear to benefit from continuity of care with both generalist and specialist doctors. Many of these articles called for continuity to be given a higher priority in healthcare planning. Despite substantial, successive, technical advances in medicine, interpersonal factors remain important.

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INTRODUCTION

Medical science has advanced rapidly since the early 19th century. Major advances

Strengths and limitations of this study

- The first systematic review of continuity of care and mortality.
- We included studies working with patients with all conditions, of all ages and of all stages of conditions.
- We included articles investigating continuity with all kinds of doctors in any health system.
- We included articles using any clearly defined measure of continuity of care.
- A meta-analysis was not possible due to heterogeneity of continuity and mortality measures.

from the germ theory to the sequencing of the human genome have together generated much deeper understanding of the pathophysiology of disease with improved prevention and treatment. However, all these advances are mostly related to physical factors. Research on human aspects of medical care has lagged.

Internationally, there has been a decrease in the perceived value of personal contact between patients and doctors. An editorial in the *New England Journal of Medicine*¹ suggested that non-personal care should become the 'default option' in medicine.

One way to study interpersonal care is by measuring continuity of care. The definition of continuity of care that we have used previously² is repeated contact between an individual patient and a doctor. Such repeated contact gives patients and doctors the opportunity for improved understanding of each other's views and priorities. Continuity of care can be considered to be a proxy measure for the strength of patient–doctor relationships.³

There have been a variety of approaches to measure continuity and so far only three randomised controlled trials have been completed.^{4–6} These all showed continuity to be beneficial for patients over relatively short

periods. However, RCTs are problematic with pre-existing long-term human relationships, like marriage and parent–child relationships, as prospective randomisation is almost impossible. Some doctor–patient relationships last for decades and become highly personal, and therefore RCTs are unethical or impractical. Observational studies have inherent limitations, and investigating continuity of care has certain problems, in particular that of reverse causality; poor health or death early in the study leading to a low measured level of continuity.⁷ However, study teams are increasingly aware of this and use study designs and analytical methods to reduce and account for it.

There is a clear rationale for the effectiveness of continuity of care as doctors collect ‘accumulated knowledge’⁸ about an individual patient which they then use in subsequent consultations to tailor advice.

Continuity of care in general practice is associated with greater patient satisfaction,⁹ improved health promotion,¹⁰ increased adherence to medication¹¹ and reduced hospital use.¹² Given all these separate benefits, the question arises whether these extend to mortality rates. Death is clearly the most important and serious of all outcomes.

Since 2010, individual studies have emerged investigating whether continuity of care is associated with reduced mortality, including some with specialists.^{13–35} These reports represent a new development, underlining the interpersonal component of medical care.

Research question

Are higher levels of continuity of doctor care, in any setting, with any patient group, associated with changed mortality?

METHODS

Search strategy and selection criteria

For inclusion in this systematic review (without meta-analysis), articles must have been published in the peer-reviewed literature, in the last 21 years, in English. We searched the databases of MEDLINE, Embase and the Web of Science from 1996 to 2017 by searching for ‘continuity’ OR ‘continuity of care’ together with terms for a medical doctor/physician and terms indicating death or mortality in the title or abstract (see online supplementary information—example search strategy). In addition, references of articles selected were hand-searched for additional relevant citations.

Experimental and observational study designs were considered including controlled trials, cohort studies (prospective and retrospective) and case–control studies. Systematic reviews and meta-analyses were excluded. Study participants could include any patient group, including entire populations or groups of patients with a specific disease or other feature.

Articles must have compared measured degrees of continuity of care with doctors (of any kind) to mortality rates. Any valid measure of continuity was considered,

including continuity being lost or absent and articles where the continuity measure was a single appointment or visit by a general practitioner/family physician during a hospital stay. Articles about organisational continuity and general staffing numbers were excluded.

As an outcome measure, any measure of mortality was accepted, that is, all-cause, time/age-limited or cause-specific. When complications or hospital admissions were combined with death rates, we sought a separate measure of mortality alone. If this was not available, studies were excluded.

Two pairs of reviewers checked the search results and decided independently whether papers met the eligibility criteria. Initially, the title and abstract of each citation was screened. The full texts of selected articles were then examined. Disagreements were resolved by discussion, and PHE independently had the deciding vote.

Data items

The variables and outcomes extracted included basic information: authors, date and country. We also extracted study design, study population (any particular condition, setting, age group, any other inclusion or exclusion criteria and selection method), numbers of patients, measure of continuity, length of continuity measurement and doctor type (generalist doctor including general practitioner, family physician and primary care physician or specialist). We extracted the period of time for the mortality measurement, and any overlap with or interval between mortality and continuity measurement periods. We also extracted whether mortality was all-cause or a disease-specific cause or limited to a particular group, how mortality was assessed and confounding factors tested or accounted for. We also extracted an estimation of any association found, with risk ratio or OR where possible and whether higher continuity was linked to an increased or decreased mortality risk. Data were extracted independently by two reviewers (of DJPG, EW, AT and KSL), using the data-extraction table designed for this review. Disagreements were resolved as described previously.

Risk of bias

The quality and risk of bias were assessed independently for individual studies by two reviewers using the Newcastle-Ottawa Scale.³⁶ We also assessed relevant areas of bias in terms of the timing of continuity and mortality measurement and confounding factors considered. For continuity of care and mortality, there is a particular potential for bias in that the worsening of health status before death may cause either decreased or increased continuity of care (reverse causality),⁷ so we noted whether this had been considered and adjusted for in study design. In terms of bias across studies, we considered publication bias and reporting bias in terms of whether mortality was the primary outcome.

Data analysis

Studies were analysed for a relationship between continuity of care and mortality rates, and whether this relationship was an inverse one (ie, greater continuity of care led to lower mortality rates) or not. For each study, we sought a risk metric (ie, relative risk ratio, HR or OR) from an adjusted model of data analysis in order to minimise the risk of selection bias and confounding. Where these statistical metrics were not reported, we provided any other available comparison measure.

Patient involvement statement

DJPG is a member of the St Leonard’s Practice Patient Participation Group as well as the Patron of the National Association for Patient Participation. As such, he is a patient representative as well as an author. The research question and outcomes were therefore conceived by a patient from the practice based on the priorities,

experience and preferences stated by patients at successive national patient conferences.

RESULTS

Study selection

After removal of duplicate results, 726 peer-reviewed publications were identified. No previous systematic reviews or trials on this subject were found. Of the 726 papers identified, 43 papers were selected for full-text review (figure 1). Articles were then excluded if continuity was not clearly measured or was the dependent variable,^{37–42} if the continuity of care measure was not clearly with a doctor or doctors only^{35 43–49} and if mortality was not analysed or not analysed separately at any point^{50–52} (eg, if it was expressed only as a composite outcome with hospitalisation). This left 22 studies for inclusion.

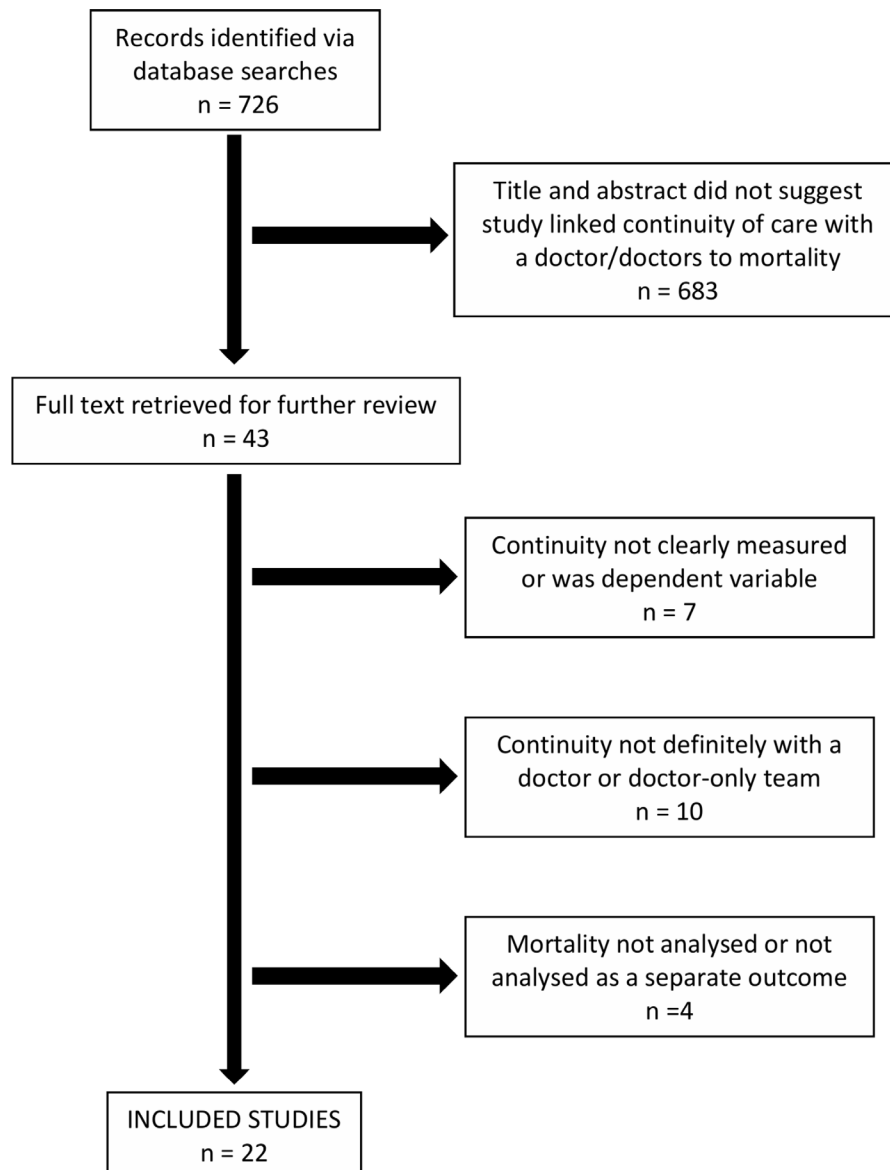


Figure 1 Study selection flow diagram.

Study characteristics

As shown in [table 1](#), the majority of included reports (15, 68.2%) were of retrospective cohort studies, often using insurance data. There were four prospective cohort and three cross-sectional studies. No randomised controlled trials were found. A number of cohort studies included large numbers of patients (median 16855). All of the reports were published since 2010. The studies were carried out in nine different countries; the majority were from North America (Canada 6, USA 5). Seven were from Europe (England 3, France 2, Croatia 1 and the Netherlands 1). There were two from Taiwan and one each from Israel and South Korea.

Nine (40.9%) of the studies investigated continuity with a general practitioner/family physician/primary care physician, 3 were with specialists only^{17 19 20} and 10 included continuity with doctors of any kind. Eight studies (34.8%) selected patients during or following an index hospitalisation.^{15–18 20 25 26 29} Five studies studied patients with diabetes^{22 23 27 30 31} and three studies focused on older patients.^{13 24 31}

The continuity measures used are reported in [table 1](#). The most common measure used was the Usual Provider of Care (UPC) index which was used in 10 studies (45.5%).^{13 16 17 21 23 25 26 29–31} Six studies used more than one measure, some only for sensitivity analysis.^{13 21 25 26 28 29} One study¹³ was designed to compare the association of different continuity measures with outcomes, including mortality. One article¹⁸ used the occurrence of a supportive visit by a family physician to a patient in hospital and another¹⁴ simply took loss of contact as meaning loss of continuity. Three studies^{32–34} used the results of a question or questions from the annual UK national General Practice Patient Survey.

The length of time over which continuity was measured (when not a survey response or hospital visit indicating a relationship) varied greatly between studies, from a single weekend in hospital¹⁷ up to 17 years.²⁴ The median length of continuity measurement was 2 years (IQR 3.75).

Most studies (20, 90.9%) reported all-cause mortality. One study³² investigated premature mortality; under the age of 75. Another³³ used premature coronary heart disease mortality as the primary outcome. The length of time for recording deaths also showed a large variation between studies, from 30 days to up to 21 years. The median follow-up time was 2.5 years (IQR 4.4).

Most of the studies investigated a large number of potential confounding factors ([table 1](#)). All studies working at the level of individual patients included some measure of health status including LACE index, comorbidities, previous healthcare usage and other measures. Most studies looked at age and sex and 14 (63.6%) used a measure of deprivation, social status or income.

Results of individual studies

Of the 22 studies, 18 (81.8%) showed that greater continuity of care was significantly associated with lower mortality. Of these, 16 (72.7% of the 22) were with lower

all-cause mortality ([table 2](#)). Two studies found no association of greater continuity of care with subsequent mortality during¹⁷ or following¹⁶ a hospital stay. One study found that continuity was not significantly associated with mortality except in general practices in the least deprived areas.³² One study¹³ investigated a range of continuity measures. They found that all insurance claims-based measures showed that higher levels of continuity were associated with higher mortality rates but greater continuity as reported by patients was associated with reduced mortality. This is the only study showing any association of increased continuity with increased mortality.

Due to the heterogeneity of study continuity and mortality measurements, it was not possible to combine them to produce an estimate of effect size; however, [table 2](#) shows the risk ratio, OR or HR from individual studies where available.

Risk of bias within studies

Using the Newcastle-Ottawa Scale,³⁶ all 22 studies were rated as high quality, with nine 10 studies (40.9%) gaining maximum scores from both reviewers independently ([table 1](#), supplementary table). No study was scored less than 7 out of 9 by any reviewer. As all these studies were cohort or cross-sectional studies, they tested for associations only. However, most involved statistical analyses for a wide range of potential confounding factors ([table 1](#)).

The specific bias of reverse causality between the healthcare-related events that might occur before death was discussed in 14 (63.6%) of the studies. Four cohort studies did not discuss reverse causality.^{14 24 27 31} However, all of the studies included some measure of health/disease status as a potential confounding factor and some included several detailed measures of these in their models.

Five of the studies had a design which meant there was no overlap between the time for continuity measurement and the period during which deaths were counted.^{13 20 22 28 30} Seven studies have complete^{14 16 21 23 28 29 31} and four partial overlap of these periods.^{17 19 24 25} Five studies included additional analyses which either eliminated the overlap²³ or introduced a lag time^{19 21 26 28} between continuity and mortality measurement periods. In each of these additional analyses, continuity was still found to be significantly associated with mortality. One long-term study²⁴ calculated survival from the date of the last continuity measurement and stratified by the length of time in the study. Five studies^{19 21 25 28 29} used their continuity score as a time-dependent variable in the model.

Risk of bias across studies

There is a risk of publication bias. It may be that reports showing no effect are less likely to be published. However, two showed no association. In two, mortality was not the primary outcome and in six, it was part of a composite outcome. For 13 studies, mortality was not the only outcome. In 10 studies, the association of two or more factors, including doctor continuity of care, with

Table 1 Studies investigating the link between continuity and mortality that meet the inclusion criteria, ordered by study design

| First author and year of publication | Country of origin | Patients | Study design | Number of patients if cohort study | Continuity measure | Continuity with measured for | Length of time continuity measured for | Confounding factors checked and/or adjusted for | Mortality measure | Quality score (mean of two reviewers) | Mortality primary outcome? | Length of time mortality counted |
|--------------------------------------|-------------------|---|--------------|------------------------------------|--------------------|---|--|---|------------------------------------|---------------------------------------|----------------------------|----------------------------------|
| | | | | | | | | | | | | |
| Bentler 2014 ¹³ | USA | 65 years+, community residing Medicare beneficiaries who completed NHHSUQ survey, not in managed care, not in MMC plan. | PC | 1219 | 1, 2, 3, 5, 8 | Physician | 1–2 years | A,B,C,D,E,F,G, H,I,J,K,M,N,O | All-cause time to death | 9 | Yes | Up to 5 years |
| Cerovečki 2013 ¹⁴ | Croatia | With opioid dependence, treated with methadone in family medicine setting. | PC | 287 | 7 | Family physician | 12 years | A,B,M,O,S | All cause | 8.5 | Yes | 12 years |
| Spatz 2014 ¹⁵ | USA | 18 years+, hospitalised with acute myocardial infarction. | PC | 2454 | 8 | Doctor | N/A | A,B,C,D,E,F,G, I,J,M,O,T | All cause | 9 | Yes | 12 months |
| van Walraven 2010 ¹⁶ | Canada | 18 years+, discharged into community from medical or surgical services of 11 Ontario hospitals. | PC | 3876 | 1 | Physician who saw patient before, during and/or after hospital stay | 6 months | A,B,H,L,N,O | All cause | 9 | Yes | 6 months |
| Blecker 2014 ¹⁷ | USA | 18 years+, hospitalised at least 2 days including at least one at weekend. | RC | 3391 | 1 | Discharge physician | 2 days | A,B,C,K,N,O,T | In hospital | 8 | No | Length of hospital stay |
| Brener 2016 ¹⁸ | Canada | 18 years+, discharged from hospital into community, family physician has history of hospital visits. | RC | 164 059 | 9 | Family physician | N/A | A,B,D,L,O,Q | All cause | 9 | Yes | 90 days postdischarge |
| Hoertel 2014 ¹⁹ | France | In CNAMTS insurance fund, saw a psychiatrist regularly. | RC | 14 515 | 2 | Psychiatrist | 3.5 years | A,B,D,K,N,O | All cause | 8.5 | Yes | 3 years |
| Justiniano 2017 ²⁰ | USA | 18 years+, underwent colorectal resection and readmitted within 30 days of DC. | RC | 20 016 | 0 | Surgeon | N/A | A,B,C,I,K,O, P,Q,T,U | All cause, colorectal cancer | 9 | Yes | 1 year |
| Leleu 2013 ²¹ | France | NHI reimbursement patients, >2 visits in 6 months. | RC | 325 742 | 1, 2 | Primary care physician/GP | 3 years | A,B,D,K | All cause | 9 | Yes | 3 years |
| Liao 2015 ²² | Taiwan | 31–99 years, with type 2 diabetes. | RC | 89 428 | 6 | Any physician | 1 year | A,B,H,K,N, O,PU | All cause | 8.5 | No | 4–9 years |
| Lustman 2015 ²³ | Israel | 40–75 years, with type 2 diabetes, remained in area, saw primary care provider >3x. | RC | 23 679 | 1 | Primary care physician/GP | 2 years | A,D,H,K,M, N,O | All cause, diabetes related causes | 8.5 | Yes | 2 years |
| Maarsingh 2016 ²⁴ | The Netherlands | 55–85 years, data available. | RC | 1712 | 3 | GP | 17 years | A,B,D,E,F,G, K,M,Q,O | All cause | 9 | Yes | 21 years |
| McAlister 2013 ²⁵ | Canada | 20 years+, DC from hospital with 1st time heart failure. | RC | 16 855 | 0, 1 | Physician who saw patient x2 in year before or 1x during admission | N/A | D,K,O,P,Q,R | All cause | 9 | Composite | 3 months/6 months |
| McAlister 2016 ²⁶ | Canada | 20 years+, DC from hospital with 1st time heart failure. | RC | 39 249 | 0, 1 | Any physician | N/A | A,B,Q,K,M, O,H | All cause | 8 | Composite | 30 days |

Continued

Table 1 Continued

| First author and year of publication | Country of origin | Patients | Study design | Number of patients if cohort study | Continuity measure | Continuity with measure | Length of time continuity measured for | Confounding factors checked and/or adjusted for | Mortality measure | Quality score (mean of two reviewers) | Mortality primary outcome? | Length of time mortality counted |
|--------------------------------------|-------------------|--|--------------|------------------------------------|--------------------|--|--|---|---|---------------------------------------|----------------------------|--|
| Pan 2017 ²⁷ | Taiwan | 35 years+, diagnosed with type 2 diabetes, in Taiwan NHI database. | RC | 396 838 | 2 | Any physician | 8 years | A, B, D, K, O, P, Q | All cause | 8.5 | Yes | Up to 8 years |
| Shin 2014 ²⁸ | South Korea | 20 years+, in Korean National Health Insurance, new diagnosis of hypertension, diabetes, hypercholesterolaemia or their complications. | RC | 47 433 | 2, 4, 5 | Physician | 2 years | A, B, D, F, K, N, Q, U | All cause | 9 | Yes | Up to 5 years |
| Sidhu 2014 ²⁹ | Canada | 'Adults' treated and released from 93 emergency departments with first-time diagnosis of heart failure. | RC | 12 285 | 0, 1 | Physician who saw patient x2 in year before or 1x during admission | 30 days | A, B, G, K, O, P, Q, N | All cause | 8.5 | Composite | 12 months but only give separate data for deaths for 30 days |
| Weir 2016 ³⁰ | USA | 20 years+, with incident diabetes and at least 2 years insurance. | RC | 285 231 | 1 | Physician who saw patient the most | 2 years | A, B, D, G, H, K, O | All cause | 8.5 | Composite | 1 year |
| Worrall 2011 ³¹ | Canada | 65 years+, with diabetes, 2+ fee-for-service family physician visits. | RC | 305 | 1 | Family physician | 3 years | A, B, N | All cause | 7 | Yes | 3 years |
| Baker 2016 ³² | England | Registered with 7858 general practices, with complete data, in England. | CS | N/A | 8 | GP | N/A | C, D, F, G, J, P | Premature-ratio observed to expected, age <75 | 8 | Yes | N/A |
| Honeyford 2013 ³³ | England | Registered with 229 general practices in the East Midlands between April 2006 and March 2009. | CS | N/A | 8 | GP | N/A | A, B, C, D, F, G, P, U | CHD under 75 and all age. | 8 | Yes | 3 years |
| Levene 2012 ³⁴ | England | 18 years+, registered with GP for at least 6/12 months of the year. | CS | N/A | 8 | GP | N/A | A, B, C, D, F, O, P | All cause, COPD, all cancer, CHD | 9 | Yes | 2 years |

Continuity measures: 1, usual provider of care index; 2, continuity of care index; 3, Herfindahl-Hirschmann Index; 4, Modified, Modified Continuity Index; 5, most frequent provider; 6, % consistency to physician; 7, loss of contact with family physician; 8, patient survey; 9, family physician visited patient in hospital; 0, follow-up by familiar doctor.

Confounding factors: A, age; B, sex; C, race; D, deprivation/social status/income; E, education; F, smoking; G, chronic conditions; H, prior hospitalisation; I, insurance; J, acute conditions; K, co-morbidity (including Charlston Index); L, LACE Index (risk of 30-day readmission or death after hospital discharge); M, marital/relationship status; N, number of healthcare visits/service intensity; O, other healthcare history; P, practice, hospital or doctor characteristics; Q, location; R, length of hospital stay; S, treatment plan; T, timing of admission; U, other.

CS, cross-sectional; CHD, Coronary heart disease; CNAMEITS, Caisse Nationale de l'Assurance Maladie des Travailleurs Salariés; COPD, Chronic Obstructive Pulmonary Disease; DC, discharged; GP, general practitioner; MMC, Medicare Managed Care; N/A, Not Applicable; NHHSUQ, National Health and Health Services Use Questionnaire; NHI, National Health Insurance; PC, prospective cohort; RC, retrospective cohort.

Table 2 Outcome measures of studies investigating the association of continuity of care with mortality

| First author and year of publication | Ratio (if available) | Other result | 95% CI | For measure | Continuity associated with mortality? | Results summary |
|--------------------------------------|----------------------|--------------|-----------------|--|---------------------------------------|---|
| Bentler 2014 ¹³ | 2.25† | | 1.33 to 3.81 | AHR above vs below mean patient-reported care site continuity. | Yes | Patient-reported duration continuity had significant, protective association with time to death. Seven claims-based continuity of care indicators and one patient-reported measure (site continuity) showed higher continuity associated with increased death hazard. |
| | 0.54* | | 0.37 to 0.8 | AHR, highest vs lowest tertile patient-reported duration continuity. | | |
| | 2.3† | | 1.56 to 3.38 | AHR, highest vs lowest tertile, UPC. | | |
| | 1.8† | | 1.12 to 2.88 | AHR, highest vs lowest tertile, inverse number of providers. | | |
| | 1.69† | | 1.13 to 2.52 | AHR, highest vs lowest tertile, MMCI. | | |
| | 1.7† | | 1.12 to 2.59 | AHR, highest vs lowest tertile, Ejlertsson's Index K. | | |
| | 2.33† | | 1.56 to 3.49 | AHR, highest vs lowest tertile, Bice-Boxerman CoC. | | |
| Cerovečki 2013 ¹⁴ | 12.6* | | 3.001 to 53.253 | OR, loss of CoC. | Yes | Loss of continuity of care one predictor of fatal outcome. |
| | 1.92* | | 1.19 to 3.12 | AHR, no usual source of care vs strong USOC relationship. | Yes | |
| | 1.03 | | 0.95 to 1.12 | AHR, increase of 0.1 in continuity score, preadmission physician. | No | |
| van Walraven 2010 ¹⁶ | 0.87 | | 0.74 to 1.02 | AHR, increase of 0.1 in continuity score, hospital physician. | | No significant association found for death risk with continuity with any doctor type studied. |
| | 0.97 | | 0.89 to 1.06 | AHR, increase of 0.1 in continuity score, postdischarge physician. | | |
| | 0.72 | | 0.29 to 1.8 | AOR, UPC 1 (complete continuity) vs 0, no continuity. | No | |
| Blecker 2014 ¹⁷ | 0.87* | | 0.82 to 0.93 | AOR, visited vs not, 90-day postdischarge. | Yes | Increased weekend UPC was significantly associated with decreased mortality in unadjusted analysis. No association after multivariate adjustment. |
| | 0.88* | | 0.81 to 0.86 | AOR, visited vs not, 30-day postdischarge. | Yes | |
| Brener 2016 ¹⁸ | 0.83* | | 0.83 to 0.83 | AHR, 0.1% increase in CoC index. | Yes | In unadjusted model, visited patients more likely to die at 90 days. In unadjusted model, visited patients less likely to die at 90 days. |
| | 0.53* | | 0.52 to 0.54 | AHR, perfect continuity vs imperfect continuity. | Yes | |
| Hoertel 2014 ¹⁹ | | | | | | 0.1 increase in CoC index associated with decreased likelihood of death. |

Continued

Table 2 Continued

| First author and year of publication | Ratio (if available) | Other result | 95% CI | For measure | Continuity associated with mortality? | Results summary |
|--------------------------------------|----------------------|---|--------------|---|---------------------------------------|---|
| Justiniano 2017 ²⁰ | 2.33 | | 2.10 to 2.60 | AHR, readmitted to original hospital but with different surgeon vs same hospital, same surgeon. | Yes | In comparison with patients readmitted to the same hospital and managed by the same surgeon, patients managed at the same hospital but by a different surgeon had > twofold risk of 1-year mortality. |
| Leleu 2013 ²¹ | 0.96* | | 0.95 to 0.96 | HR, 0.1 increase in CoC. | Yes | Increase in the CoC index associated with decrease in death risk. |
| Liao 2015 ²² | * | Significant trend (p<0.001, test for monotonic trend) | | Decreasing consistency in medical care-seeking behaviour with decreasing adjusted survival. | Yes | A significant monotonic trend was observed between decreasing consistency in medical care-seeking behaviour (from high consistency to low consistency) and decreasing multivariate-adjusted survival. |
| Lustman 2016 ²³ | 0.59* | | 0.5 to 0.7 | OR, high vs low UPC, measured at the same time. | Yes | Patients with a high UPC had lower risk of mortality. Not affected on adjusting for background characteristics. |
| | 0.7* | | 0.56 to 0.88 | OR, high vs low UPC, measured in successive years. | | |
| Maarsingh 2016 ²⁴ | 1.2* | | 1.01 to 1.42 | HR, lowest vs highest CoC. | Yes | In final model, participants in lowest CoC category showed greater mortality than those in maximum. |
| McAlister 2013 ²⁵ | 0.86 | | | HR, familiar vs unfamiliar (our calculation, CI not available). | Yes | After 6 months, death HR for familiar Dr 0.66 (95% CI 0.61 to 0.71) and 0.77 (0.68 to 0.88) with unfamiliar vs no follow-up. At 3 months, 1.6% of those who had a visit with a familiar Dr died, 3.3% who only saw an unfamiliar Dr, p<0.001. |
| McAlister 2016 ²⁶ | * | 3.1% vs 2.0%, p<0.0001 | | % mortality: follow-up by unfamiliar or familiar physician. | Yes | More died with follow-up with unfamiliar physician compared with those with at least one visit with familiar physician. |
| Pan 2017 ²⁷ | 0.47* | | 0.46 to 0.48 | AHR high (>50%) vs low (≤50%) CoC score. | Yes | Patients with diabetes with higher physician continuity had a lower risk of mortality. |
| Shin 2014 ²⁸ | 1.13* | | 1.05 to 1.21 | AHR, below vs above median most frequent provider. | Yes | Above median continuity associated with lower all-cause mortality using three different measures. |
| | 1.13* | | 1.05 to 1.21 | AHR, below vs above median MMCI. | | |
| | 1.12* | | 1.04 to 1.21 | AHR, below vs above median CoC. | | |
| Sidhu 2014 ²⁹ | * | 1.9% vs 1.4%, p<0.0001 | | % mortality: follow-up by unfamiliar or familiar physician. | Yes | More died with follow-up with unfamiliar physician compared with those with at least one visit with familiar physician. |
| Weir 2016 ³⁰ | 0.75* | | 0.61 to 0.94 | AOR, high vs low UPC. | Yes | High UPC associated with decreased mortality. |
| Worrall 2011 ³¹ | * | 9.0% vs 18.1%, (p=0.025, χ^2) | | % mortality: high vs low continuity group. | Yes | Proportion of people dying significantly lower in high-continuity group. |

Continued

Table 2 Continued

| First author and year of publication | Ratio (if available) | Other result | 95% CI | For measure | Continuity associated with mortality? | Results summary |
|--------------------------------------|----------------------|--------------|----------------|--|---------------------------------------|---|
| Baker 2016 ³² | | 21 deaths | -16 to 63 | Potential reduction in premature deaths in England in 1 year if there is a change of 1 percentile of patients expressing trust in their doctor. | No | Continuity not associated with mortality (except in less deprived practices in a separate subgroup analysis). |
| | | -49 deaths | -250 to 156 | Potential reduction in premature deaths in England in 1 year if there is a change of 1 percentile of patients able to get an appointment in advance. | | |
| Honeyford 2013 ³³ | 0.994* | | 0.989 to 1 | IRR, 1% change in survey response. | Yes | An increase in % of patients recalling being able to see their preferred GP was associated with decreased mortality. |
| Levene 2012 ³⁴ | 0.999 | | 0.997 to 1.01 | IRR, all-cause mortality. | Depends on mortality measure | No significant association with all-cause mortality. |
| | 0.997* | | 0.995 to 0.999 | IRR, all-cancer mortality. | | An increase in the % of patients recalling being better able to see their preferred doctor was associated with decreases in COPD mortality and in all-cancer mortality. |
| | 0.999 | | 0.995 to 1.07 | IRR, coronary heart disease mortality. | | |
| | 1.0002 | | 0.99 to 1.01 | IRR, stroke mortality. | | |
| | 0.993* | | 0.98 to 0.998 | IRR, COPD mortality. | | |

*Significant result showing higher levels of continuity associated with lower mortality.

†Significant result showing higher levels of continuity associated with higher mortality.

AHR, adjusted HR; AOR, adjusted GP, general practitioner; OR, CoC, Continuity of Care Index; IRR, incident rate ratio; MCI, modified continuity index; MMCI, Modified Modified Continuity Index; UPC, Usual Provider of Care Index; USOC, usual source of care.

outcomes was tested. Continuity and mortality as exposure and outcome, respectively, are reported in a range of studies, including where testing this association was not the primary aim.

DISCUSSION

Principal findings

In a substantial majority of studies (18, 81.8%) meeting the selection criteria, higher levels of continuity of care with doctors were associated with lower mortality rates. Two others, finding no significant association, had very short timescales for measurement of continuity, to the extent that the strength of any patient–doctor relationship was potentially questionable. Another study showing no significant association with all-cause mortality was cross-sectional, and the measurement methods related to questions on a national survey about seeing a particular general practitioner, again not necessarily indicative of a strong patient–doctor relationship.

One study⁸ found that for claims-based measures of continuity, increased mortality was associated with higher levels of continuity of care. However in the same study, higher levels of patient-reported continuity were associated with lower mortality rates. This emphasises the interpersonal relationship between patient and doctor as claims-based measures only give numbers of contacts and do not directly measure the quality of the relationship.

The effect sizes were generally small (table 2) but these were in the same range as some treatment effects, as very large, repeatable effects on mortality are rare.⁵³ In addition, for some studies included in this review, effect sizes were calculated using very small increments in the continuity measure.

Strengths and weaknesses of the evidence

All the studies found investigating the association of continuity of care with mortality were observational in nature, although the majority were high-quality cohort studies including three prospective cohort studies. The issue of reverse causality applies to all the evidence presented here. This could bias an association between continuity of care and mortality in either direction. As patient health worsens when approaching death, continuity of care may deteriorate for many reasons, for example, patients moving areas to accommodate increased health needs, the need to see more specialists or a loss of ability to obtain and attend appointments. Alternatively, deterioration of health could lead to a concerned doctor ensuring that the patient receives more continuity of care. For the cross-sectional studies, there is also a potential for confounding due to practice-level factors.

There have been randomised controlled trials into continuity of care but none on existing relationships or lasting longer than a year and none with mortality as an outcome.^{4–6} Observational studies which control rigorously for confounding factors and have a design aimed

at limiting the impact of reverse causality are the best evidence available.

Of the 16 cohort studies finding an association of higher continuity with lower mortality, most studies attempt to at least partially account or control for reverse causality in their study design or analysis. Most controlled for differences in health status and risk factors. Some carried out analyses measuring continuity and mortality in separate years, or with a lag. This method, particularly the lag between measurements, should help to minimise bias caused by rapid worsening prior to death. However, four cohort studies showing this association^{14 24 27 31} did not discuss this kind of reverse causality although one²⁴ nevertheless made several adjustments for health status and calculated survival from the date of the last continuity measurement. Measuring continuity and mortality over separate time periods is also one way of eliminating the potential bias caused by those who survived longer having more time to accrue continuity (time-dependent bias). Another way of reducing this is to model continuity as a time-dependent variable which was the case in five studies.^{19 21 25 28 29}

All studies included were rated as high quality, using the Newcastle-Ottawa Scale.

Several of the articles reported on studies using very large cohorts. The studies came from a number of different countries with different healthcare systems and cultures. Continuity of care in the studies included that received from specialist as well as generalist doctors, showing that the effect is not limited to one branch of medicine or health system.

As continuity research is an emerging field, no consensus on the best way to measure it has been reached. The measure used most was the UPC Index which does not take into account the total number, frequency or sequence of visits.⁵⁴

Doctors have been studied as a discrete category in numerous studies, and data systems usually allow them to be separately studied. The group studied included family doctors/general practitioners, physicians and psychiatrists so was already heterogeneous so expanding this to other professional groups would have complicated interpretation. As doctors are the most highly trained health professionals with the most influence over decisions, it is reasonable to assume that if interpersonal contact affects mortality, it is most likely to occur with doctors. Therefore, we eliminated articles, some with significant reductions in mortality, that measured continuity in relation to mixed profession teams or to other health professionals.^{35 43–49} This is the first systematic review investigating whether continuity of doctor care is associated with reduced mortality. We expect this to encourage studies with different selection criteria; for example, for continuity with other healthcare professionals.

Possible mechanisms and implications

This review, finding that increased receipt of continuity of care is associated with reduced mortality, comes after it

has been shown that continuity of care is associated with multiple benefits for patients.^{9–12} It therefore fits well with such earlier work. It is only recently that large databases and long-term cohort studies have made effective investigation into the links between continuity and mortality possible.

These known associations suggest possible mechanisms in that greater uptake of evidence-based preventative medicine such as immunisations as well as better concordance with treatments is likely to reduce mortality. Continuity of care is associated with patients perceiving that the doctor has become more responsive.⁵⁵ Patients then disclose more and medical management is more likely to be tailored to the needs of the patient as a person. The increased patient satisfaction may also be associated with an 'optimism' boost to health.⁵⁶ We have previously suggested that 'doctors tend to overestimate their effectiveness when consulting with patients they do not know, and underestimate their effectiveness when consulting with patients they know'.⁵⁷

The cumulative impact of these multiple gains may then be reflected in reduced mortality.

Historically, continuity of care has been considered a feature of the practice of medical generalists and featured in the job descriptions of the general practitioner.^{58 59} Recent studies included in this review found that continuity was associated with reduced mortality with specialist physicians,^{22 28} psychiatrists¹⁹ and surgeons²⁰ too.

Although this evidence is observational, with 18 of the 22 studies showing significant reductions in mortality with continuity of doctor care, the clear preponderance of evidence is in favour of the association. Three studies showed no significant association and one¹³ had mixed results but no study exclusively showed an association of higher continuity of care with higher mortality rates. Although there are difficulties in carrying out controlled trials on this subject, a few, with interventions to increase continuity of care, have been successful,^{4–6} and this could be attempted more widely. The presence of this association in nine countries, across three continents, and in very different populations and healthcare systems implies a basic human effect.⁶⁰ The policy implication as many studies noted is prioritising continuity of care.

For 200 years, medical advances have been mainly technical and impersonal which has reduced attention to the human side of medicine. This systematic review reveals that despite numerous technical advances, continuity of care is an important feature of medical practice, and potentially a matter of life and death.

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in article selection, data extraction and assessing article quality. KS-L carried out data analysis. All authors wrote and edited the manuscript. All authors approved the manuscript for publication.

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