Data and care integration for post-acute intensive care program of stroke patients: effectiveness assessment using a disease-matched comparator cohort

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Abstract

Purpose – To assess the effectiveness of an integrated care program for post-acute care of stroke patients, the return home program (RHP program), deployed in Barcelona (North-East Spain) between 2016 and 2017 in a context of health and social care information systems integration.

Design/methodology/approach – The RHP program was built around an electronic record that integrated health and social care information (with an agreement for coordinated access by all stakeholders) and an operational re-design of the care pathways, which started upon hospital admission instead of discharge. The health outcomes and resource use of the RHP program participants were compared with a population-based matched control group built from central healthcare records of routine care data.

Findings – The study included 92 stroke patients attended within the RHP program and the patients’ matched controls. Patients in the intervention group received domiciliary care service, home rehabilitation, and telecare significantly earlier than the matched controls. Within the first two years after the stroke episode, recipients of the RHP program were less frequently institutionalized in a long-term care facility (5 vs 15%). The use of primary care services, non-emergency transport, and telecare services were more frequent in the RHP group.

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Ethics statement: The Independent Ethics Committees of each of the participating hospitals approved the study protocol. Participants provided informed consent.

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Conflicts of interest: The authors declare no conflicts of interest regarding this work.
Originality/value – The authors’ analysis shows that an integrated care program can effectively promote and accelerate delivery of key domiciliary care services, reducing institutionalization of stroke patients in the mid-term. The integration of health and social care information allows not only a better coordination among professionals (thus avoiding redundant assessments) but also to monitor health and resource use outcomes of care delivery.

Keywords Information systems, Integrated care, Stroke, Domiciliary care, Home care

Paper type Research paper

Introduction
The continuity of care during transitions between different places or settings is the cornerstone of patient-centered integrated care. Of all these transitions, hospital discharge after a disabling health event has remarkable impact on patients’ life, particularly those with continuous complex care needs (Coleman, 2003). The difficulties associated with discharge of these patients can be overcome by appropriate comprehensive discharge plans that consider education on self-management, coordination of care providers, medication reconciliation, and assessment of financial barriers, among others (Coleman, 2003; Le Berre, 2017). Comprehensive discharge plans have shown to reduce post-discharge mortality, emergency department visits, and readmissions in patients with chronic complex diseases transiting from hospital to primary care (Le Berre, 2017).

Continuity of care is particularly important for the management of stroke, which affects over 80 million people worldwide, and it is leading cause of disability among adults (World Stroke Organization, 2016). Owing to their limited capacity for activities of daily living, stroke patients often require domiciliary care after hospital discharge, which is crucial for long-term results in these patients (Wolfe, 2000). Home care services, aimed to support people in their activities of daily living, are often prescribed and/or provided by social care services. In our country, as in many other areas, social care services are governed by political entities different from healthcare services, with different pathways and information systems. Hence, the continuity of care in this transition strongly depends on the adequate integration between social and health care services that identifies the patient needs from all spheres of care. Unsuccessful integration often results in a delay of domiciliary rehabilitation and care, which may worsen the health outcomes in stroke patients (Hu, 2010; Kwakkel, 2004; Langhorne, 2017).

Owing the separated governance of this healthcare and social services, information transfer of patients’ characteristics and needs is a mainstay for achieving continuity of care. In our area, the deployment of digital technologies for integrating social and healthcare data has significantly improved professional coordination, thus boosting integrated care pathways in the care of chronic patients (Cano, 2017; Baltaxe, 2019; Piera-Jiménez, 2020). Several of these programs were under the umbrella of the Nextcare European program, which aimed at fostering innovation of digitally-supported care services for chronic patients (Eurecat, 2022).

The healthcare burden of stroke is expected to increase with the population shift forecasted for the upcoming years. Although projections estimate a slight decrease of disability-adjusted life years despite the increasing incidence (Wafa, 2020), the raising number of people with rehabilitation needs is likely to impact health and social care systems in high-income countries. Furthermore, the aging population will increase the proportion of patients with continuous complex care needs, including domiciliary care services, contributing to the overload of healthcare services. This scenario stresses the need for implementing integrated care pathways that efficiently screen stroke patients and design adequate post-discharge care plans that cover both the social and health care needs of patients.

In Catalonia (North-East Spain), home care is provided by two different entities: the primary care team (ruled by the ministry of health) provides healthcare on demand aimed at managing healthcare needs, whereas social care services (ruled by the city council) provide
social care (named “domiciliary care” hereafter), which includes at-home rehabilitation and support with activities of daily living, if necessary. Typically, the hospital-based social worker requests home care services (and all other home needs) to both the primary care team and the social care team upon discharge. In 2016, a healthcare area in Catalonia started an integrated care program that included an integrated electronic record of health and social care information (with shared access by all stakeholders) and a new pathway in which home care resource allocation started early after admission. In this study, we assessed whether the implementation of this post-stroke intensive home care program based on the integration of social and health care for improving domiciliary care of stroke patients after hospital discharge would shorten the time to service provision and increase the time to key unwanted health outcomes, such death and institutionalization in a long-term care facility.

Methods
Study design and patients
This was a retrospective, matched control study to assess the effectiveness of an integrated care program for post-acute care of stroke patients (i.e. the return home program [RHP]) in Barcelona (North-East Spain). The study included all consecutive patients entering the program between February 15, 2016 and February 15, 2017. All patients with domiciliary care needs were consecutively screened for the program. Inclusion criteria were living alone or with another person with limited capacity to deliver adequate care, previous dependency, cognitive impairment, and stroke severity (National Institute of Health Stroke Scale, NIHSS >10). Patients leaving in a nursing home at the time of the stroke event were not considered for the analysis.

Following a similar approach as reported elsewhere (Yiu, 2012; Yamada, 2018), we built a comparator group using patients from the general population admitted to any of the tertiary hospitals of Barcelona because of stroke (Hospital Clínic, Hospital del Mar, Hospital de Sant Pau, and Hospital Vall d’Hebron) within the same period. Patients living in a long-term care facility, with missing data in any of the matching variables, and those who died during hospital stay or within the 6 months following the stroke episode were not considered for the control group. Case-control matching was performed at 1:4 ratio based on variables potentially affecting the utilization of healthcare resources, including sex, age (±5 years), the place of residence, and their healthcare risk, estimated using the morbidity adjusted groups (GMA, for Catalan Grups de Morbiditat Ajustada). Briefly, the GMA is a population-based case-mix tool that allows stratifying the general population into mutually exclusive health-risk groups based on all chronic conditions and recent acute diagnostic codes (Monterde, 2017). The tool has shown high predictive capacity of key clinical and resource utilization outcomes, including mortality, hospitalizations, and visits to primary care, polypharmacy, and overall expenditure (Vela, 2021; Monterde, 2020). The selection of control individuals was prioritized according to the matching of the following conditions: type of stroke (bleeding stroke, ischemic stroke, and others), annual income (grouped into four categories of pharmaceutical co-payment), admission to intermediate care after acute care discharge, place of residence, and least age difference.

Intervention and usual care
The RHP was developed to improve the post-stroke rehabilitation process and promoting the achievement of the highest rate of autonomy in the shortest possible time. The specific objectives of the RHP were (1) to identify stroke patients with social care needs during hospital stay, (2) to prescript and activate the appropriate domiciliary care according to the needs of the patient and his/her environment before hospital discharge, and (3) to coordinate synchronous interventions of rehabilitation, social care, and primary care. The RHP intervention includes
three consecutive phases: candidate screening, assessment and prescription of social needs, and activation of domiciliary care provided by the local social care service before patient discharge (Figure S1, Supplementary appendix). Candidate screening occurred within the 48 h following hospital admission; candidates were considered to have social needs if they met at least one the following indicators: living alone or with another person with limited capacity to deliver adequate care, previous dependency, cognitive impairment, and stroke severity (NIHSS >10). Once the patient was considered suitable for the RHP, the hospital-based social worker prescribes the social care resources needed for an 8-week period. The main social services prescribed at home were individual care supporting for activities of daily living, house cleaning, telemedicine, meals-on-wheels, and rehabilitation aids. Patients and their families were provided with the planned date for discharge; 72 h before discharge from acute hospital (and 5 days before discharge from intermediate care center) the prescribed services were notified to the city Social Care Services and activated.

Patients receiving usual care were assessed by the hospital-based social worker for dependency based on their clinical situation and social environment at discharge. In case the social worker considered that the patient required domiciliary care, two separate requests were made to the primary care team (managed by the autonomous government) and to the social care team (managed by the city council). The city social care team assessed the patients again and included them in a general waiting list and contacted them when possible to start a domiciliary care program.

At-home rehabilitation, aimed at relearn skills lost or damaged by the stroke incident and regain independence, was delivered equally in the two groups following the local guidelines for post-acute rehabilitation of stroke patients (Duarte, 2009).

Information system integration
Since year 2007, all centers in our area share the health-related information (i.e. clinical course, test results, digital imaging, etc.) through the Shared Electronic Health Record of Catalonia (“Història Clínica Compartida de Catalunya” [HC3]) with the main aim to improve the continuity of care (Marimon-Sunyol, 2010). Since 2015, an exchange of health and social care information was agreed among the public Catalan health system and the Barcelona City Council (Solans, 2018). This exchange includes information regarding the social context, grade and characteristics of dependency, home care resources, pharmacy, discharge planning, and a summary of the individual healthcare plan. The HC3 can be accessed by the hospital social worker and the physicians and nursing teams assigned to the patient. On the other hand, information regarding social care needs and resources is collected and handled by the Barcelona City Council. Social data can be accessed by the city council social worker. Furthermore, healthcare data are periodically transferred to the Ministry of Health of the autonomous community of Catalonia for healthcare planning and analytical purposes, including public health and resource utilization.

For the RHP project, we created an ad hoc registry of stroke patients entering the program that gathered relevant health and social information of the patient. The two entities governing the datasets (i.e. the Catalonia ministry of health and the Barcelona City Council) signed an agreement for data transfer, which defined the governance of the integrated registry, the access profile, and measures for ensuring confidentiality and adherence to the organic low 15/1999 on personal data protection. For patients receiving usual care, the social care team managed by the city council could not access health care data.

Study outcomes and data sources
The primary objective of the study was to assess the time to key events, including adverse endpoints (i.e. death and institutionalization in a long-term care facility) and service provision
endpoints (i.e. receiving domiciliary care, telecare, and at-home rehabilitation services). Other outcomes included health and social care resource utilization. Resource utilization included the monthly number or average of total hospital admissions, emergency hospital admissions, visits to the emergency department, visits to a specialist, drug units (packages) dispensed by the community pharmacist, non-emergency medical transport rides, domiciliary rehabilitation visits, and contacts with primary care services, total and grouped according to the following services: general practitioner, nurse, social worker, domiciliary care team, and remote consultations. Health care expenditure was estimated as described before (Vela, 2019), whereas social expenditure was provided by the Social Service Institute of Barcelona. Briefly, expenditure is attributed to each patient through a personal identification number. Expenses associated with primary care resources are estimated according to the type of healthcare resource (e.g., physician, nurse) and site where healthcare is delivered. Hospital procedures expenditure is estimated using diagnosis-related groups, whereas expenditure associated with inpatient care (e.g., hospital, skilled nursing facilities, and mental health centers) are estimated based on the length of stay. For all outcomes, patients were followed up to December 31, 2019 or a relevant outcome, such as death or institutionalization in a nursing home.

**Statistical analysis**

Quantitative variables were described using the mean and median, whereas categorical variables were described as the frequency and percentage over available data. Time-to-event analyses were performed using the Kaplan–Meier estimate and compared with the log-rank test. Categorical variables were compared using the chi-squared test and continuous variables with the T Student’s test, except expenditure, which was not normally distributed and was compared using the Kruskal–Wallis test. The between-group differences in resource utilization and expenditure were assessed using a ratio of monthly mean utilization rates or expenditure and computed using generalized linear mixed models (Poisson for resource utilization and lognormal for expenditure), considering the random effects for matched patients. The models were adjusted for age, sex, annual income, GMA status, type of stroke, admission to intermediate care center after acute care discharge and healthcare expenditure during the year preceding the stroke episode. The significance threshold for all analyses was set at a 5% two-sided alpha error. All analyses were conducted with R (R Core Team, 2021).

**Results**

**Study participants**

During the study period, 92 patients entered the RHP program after an ischemic ($n = 71$; 77.2%) or a hemorrhagic ($n = 18$; 19.6%) stroke. Table 1 summarizes the main demographic, clinical, and resource use characteristics of study patients and the matched population-based control group. The two groups were similar in all characteristics, except for type of stroke, which was more dominantly ischemic in the control group. The previous healthcare resource utilization, not included among the pairing criteria, was also similar in the two groups.

**Clinical and healthcare outcomes**

The survival analysis did not reveal significant differences between cases and controls regarding mortality after a stroke event (Figure 1a). On the other hand, patients in the control group were more likely to be institutionalized earlier than those in the intervention group (Figure 1b); two years after the stroke episode, the proportion of patients institutionalized in a long-term care facility was 14.2% ($n = 51$) and 4.8% ($n = 4$) in the control and intervention groups, respectively.
Patients in the intervention group were more likely to receive earlier key social and health home care services (Figure 2). Two years after the index stroke episode, domiciliary care was being provided to 85 (92.4%) of patients in the RHP group and 69 (19.1%) in the control group (Figure 2a). Likewise, telecare services, which were provided to nearly one-third of individuals before the stroke episode (27.2% and 29.6% in the RHP and control groups, respectively), amounted to 75.5% (n = 67) in the RHP group and 44.5% (n = 163) in the control group two years after the index episode (Figure 2b). At-home rehabilitation services were provided early in the two groups. However, a greater percentage benefited from this service in the RHP group; two years after the index stroke episode, the proportion of patients receiving at-home rehabilitation services amounted to 90 (98.4%) and 311 (84.8%) in the RHP and control group, respectively (Figure 2c).

Figure 3 summarizes the place of stay of stroke patients within the two years following the index episode. The proportion of patients at home in the RHP increased rapidly and remained higher than in the control group until the end of the follow-up, when the proportion of patients at home in the RHP and control group were 92.3 and 83.4%, respectively. Patients in the control group stayed more frequently in a long-term care facility throughout the follow-up period.

**Resource utilization**

Within the first month following the index stroke episode, the mean social and health care expenditure was remarkably higher in the RHP group (€3,152) than the control group (€2,364),
Figure 1.
Survival analysis of relevant health and healthcare outcomes after the index stroke episode: (a) all-cause mortality and (b) institutionalization in a long-term care facility
mostly due to the higher proportion of patients institutionalized in an intermediate care facility immediately after the stroke event. The monthly mean expenditure attributed to social services, intermediate care facility, outpatient visits, emergency visits, hospitalization, pharmacy, and primary care within the first year after the stroke episode is summarized in Figure S1 (Supplementary appendix). Between months 2 and 12 after the index episode, the mean monthly expenditure of the RHP and the control group were €784.58 and €657.15, respectively. The between-group differences in expenditure were more notable in the social care (€257.95 vs. €156.40) than the health care provision (€526.64 vs. €500.75). The mean monthly expenditure for the 1-to-12 months and 2-to-12 months' period according to the type of resource is provided in Table S1.

Figure 4 summarizes the difference in use of social and health care services between patients in the control and intervention group within the year following discharge (generalized mixed models); the monthly use of each resource within the two years following discharge is shown in Figure S2. The utilization of most services did not change significantly between groups. However, the overall expenditure was significantly higher in the intervention group, mostly due to the higher expenditure in social services and –less pervasive– the number of visits to primary care (Figure 4a). Of all primary care services, non-emergency transport, and telecare services were those which increased more in the intervention group (Figure 4b).

Discussion
Our retrospective analysis of an integrated care program for stroke patients discharged from the hospital showed that program recipients received key rehabilitation services, such as intensive post-acute home care and support for activities of daily living, earlier than stroke patients managed as usual. Of note, most stroke patients in the population-based matched control group eventually received domiciliary care services. Considering that, in the absence post-discharge supporting services (including adequate rehabilitation), most patients with
Figure 3.
Place of stay of patients in the control (a) and intervention (b) groups within the two years following discharge from the index episode.
stroke experience a progressive decline in functional mobility soon after discharge (Langhorne, 2017; Buvarp, 2020), this finding suggests that the later start of service provision among control patients was due to delay in planning, rather than the lack of need.

A remarkable finding of our analysis was that recipients of the RHP program were admitted in a long-term care facility less frequently and later. Thus, although stroke patients have higher risk for admission to long-term care facilities (Luengo-Fernandez, 2013; Ribera, 2022), this finding suggests that institutionalization is sometimes a consequence of inadequate or late domiciliary care provision. The dynamics of institutionalization in nursing homes have a remarkable impact on patients’ life, but also on the sustainability of the system. The implementation of a program for promoting domiciliary care is expected to increase the social expenditure in the short term, and this was the case in our experience. However, nursing home costs typically exceed domiciliary care costs (nearly four times in our area) and have been identified among long-term services with higher contribution to the economic burden of stroke (Rajsic, 2019). Therefore, in the mid- and long-term, early delivery of

**Note(s):** The central line represent the monthly rate, adjusted using a generalized linear model; values > 1 indicate higher average in the intervention group. The limits of the box indicate the 95% confidence interval.
domiciliary care in stroke patients would alleviate the financial burden of the whole system. This effect, however, would not manifest unless pooling budget policies, based on a joint health and social care funding, are implemented. Future cost-effectiveness analyses shall focus on the financial trade-off between increased social care services and savings associated with better and earlier recovery of skills lost because of the stroke episode.

One of the key elements of an integrated care program for early delivery of both social and health care services is the adequate integration between the two information systems, which in our country—as well as in most countries—are handled by different entities and often have non-overlapping access profiles. The integrated storage of social and health care information has been acknowledged as a cornerstone for a high-quality person-centered care (Øvretveit, 2017); still, the number of works reporting the integration of social and healthcare information are still limited. In our experience, the integration of these two information systems, in force by the time of designing the RHP program, was a mainstay for a successful deployment of the integrated care approach. Our intervention met some of the key features recommended for technologies facilitating integrated care, including the ease of adoption by professionals and tail of services to the specific needs of the user (Goodwin, 2018). Conversely, other features with potential benefits for care provision, such as the inclusion of patients (Yadav, 2019) or third sector organizations involved in long-term care (Piera-Jiménez, 2020) into the information hub, shall be implemented in future revisions of the program. Although the RHP program integrated only social and healthcare workers, the integration of the two information systems avoided redundant assessments by the health and social care teams, which in our area—like in most countries—belong to different government departments and entities, responsible to provide these services. In this regard, a systematic integration of health and social information systems would ease benchmarking analyses as well as health and social care planning.

Our study is strengthened by the inclusion of a population-based control group, which was feasible thanks to the central storage of healthcare data from nearly the entire population of Catalonia, as described elsewhere (Vela, 2021; Monterde, 2020). This design allowed us the quantitative assessment of the integrated care pathway, including hard endpoints, such as mortality or admission in a long-term care facility, which provide strong evidence on the effectiveness of the intervention. In this regard, our analysis is an example of how adequate information systems may enable establishing the so called “learning healthcare systems”, aimed at improving the quality of care by recursively adopting and assessing evidence-based solutions (Maddox, 2017). On the other hand, pairing of individuals in the RHP program and those in the control group was limited to the variables collected in central records. These records do not contain data related to the stroke episode such as stroke severity according to the National Institute of Health Stroke Scale, and relevant patient information such as the Barthel score for physical disability or the Pfeiffer scale for cognitive impairment. Of note, although stroke severity could not be included in the pairing, recipients of the RHP program were selected according to their health and social needs and are, therefore, expected to have poorer health status than those in the control group. Finally, the retrospective nature of the analysis precluded the administration of validated questionnaires for assessing the quality of care.

In summary, our analysis showed that an integrated care program of health and social care for stroke patients at discharge successfully promotes early domiciliary care delivery, associated with a less frequent and later institutionalization in a long-term care facility. The integration of health and social care information systems enables not only the appropriate coordination of the various stakeholders, but also the analysis of the outcomes of care delivery. The benefits of post-discharge integrated care services for stroke patients are likely to increase in the context of pooling budget approaches that consider and plan social and health care services as a whole.
References


Supplementary appendix

**Table S1.** Social and health care expenditure after discharge

<table>
<thead>
<tr>
<th>Period</th>
<th>Group</th>
<th>Outpatient resources</th>
<th>Inpatient resources</th>
<th>Pharmacy</th>
<th>Other</th>
<th>Total healthcare</th>
<th>Social care expenditure</th>
<th>Total expenditure</th>
</tr>
</thead>
<tbody>
<tr>
<td>Months</td>
<td>Control</td>
<td>117.48</td>
<td>398.54</td>
<td>110.46</td>
<td>29.06</td>
<td>655.54</td>
<td>145.63</td>
<td>801.17</td>
</tr>
<tr>
<td>1–2</td>
<td>Intervention</td>
<td>132.94</td>
<td>425.88</td>
<td>96.19</td>
<td>81.00</td>
<td>736.01</td>
<td>251.01</td>
<td>987.02</td>
</tr>
<tr>
<td>Months</td>
<td>Control</td>
<td>112.49</td>
<td>248.65</td>
<td>110.04</td>
<td>29.56</td>
<td>500.75</td>
<td>156.40</td>
<td>657.15</td>
</tr>
<tr>
<td>2–12</td>
<td>Intervention</td>
<td>133.10</td>
<td>211.90</td>
<td>97.87</td>
<td>83.77</td>
<td>526.64</td>
<td>237.95</td>
<td>764.58</td>
</tr>
</tbody>
</table>
Figure S2.
Monthly use of resources within the 24 months following discharge in the integrated care (intervention) group (a) and population-based control group (b).
Integrated care for stroke patients

Figure S2.
Figure S2.
Figure S2.
Figure S2.

Remote consultations

Visits to the primary care center
Figure S2.