Activity of daily living upon admission is an independent predictor of in-hospital mortality in older patients with community-acquired pneumonia

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Abstract

Background: Older patients hospitalized with community-acquired pneumonia (CAP) are at high risk for short-term mortality. Activity of daily living (ADL) is associated with clinical outcomes in older patients. We aimed to investigate the prognostic value of ADL upon admission on the in-hospital mortality in older patients with CAP.

Methods: We conducted a retrospective cohort study involving patients aged ≥65 years admitted to Beijing Chao-Yang hospital due to CAP between June 2012 and June 2020. ADL evaluation upon admission was performed by Barthel Index (BI). Data from all patients were extracted from the electronic medical records.

Results: Four thousand eight hundred eighty patients were included, 131 patients (2.7%) died during their hospitalization. Median BI in the Deceased group was 45 (20–65), Deceased group had lower BI scores than Survivors group (p < 0.001). Low BI (< 60) was more frequent in patients who died in the hospital than in patients discharged alive (69.5% vs. 13%, p < 0.001). In-hospital mortality was higher among patients with worse ADL upon admission (BI< 60) compared to those BI≥60 (12.6% vs. 0.9%). The worse ADL upon admission (BI< 60) was associated with an increase in the risk of death during CAP hospitalization, worse ADL upon admission (BI< 60) showed an odds ratio (OR) for in-hospital mortality of 7.53 (95%CI: 2.77–20.48; P < 0.01). This association remained significant after adjustment for age, comorbid conditions, respiratory failure, pathogens and laboratory findings (OR, 3.74; 95%CI, 2.37–5.91; P < 0.01). Receiver operating characteristic (ROC) curve revealed that BI upon admission is a predictor related to in-hospital mortality in elderly patients, the area under the ROC curve of BI in predicting in-hospital mortality was 0.81 (with 95% confidence interval: 0.78–0.85). The predictive value of ADL upon admission was better than age in our study population.

Conclusion: Activity of daily living upon admission is an independent predictor of in-hospital mortality in older patients with community-acquired pneumonia.

Keywords: Community-acquired pneumonia, In-hospital mortality, Activity of daily living, Barthel index, Older patients
Background
Community-acquired pneumonia (CAP) is the commonest cause of infectious death. The incidence and mortality of CAP are linked to increasing age [1, 2]. It is estimated that incidence of CAP patients ≥65 years old was 140 cases per 10,000 persons per year and 105 cases per 10,000 for hospitalized [3]. Mortality of CAP in older patients is more higher than non-elderly patients [3, 4]. Older patients hospitalized with CAP are at high risk for short-term mortality [5]. Thus, there is a clear need to focus on in-hospital mortality in older patients with CAP.

There is growing evidence that the functional status, is more important than age and comorbidity in predicting prognosis in the elderly [6, 7]. Activities of daily living (ADL) assessment is a convenient way to assess a person’s functional level. Even small changes in the ADL functional level are associated with clinically relevant outcomes. ADL functional status decline can lead to adverse clinical outcomes in infections [8–11], acute medical patients [6, 12–14], dementia [15], heart failure [16], hip fractures [17, 18] and geriatric trauma [19]. The ADL functional status may be one of the predictor factor for in-hospital mortality in older patients with CAP. However, there are few data on relationship between ADL and in-hospital mortality in older patients with CAP.

Thus, the purpose of this study is to investigate the prognostic value of ADL upon admission on the in-hospital mortality in older patients with CAP.

Methods
Study subjects
We conducted a retrospective study cohort up to hospital discharge. Between June 2012 and June 2020, 4880 patients diagnosed with CAP and age ≥65 years were admitted to Beijing Chao-Yang Hospital, Capital Medical University were included in this study. Patients with recent hospitalization or immunosuppressed treatment in the prior 90 days, solid organ or stem cell transplantation in the prior 90 days, cancer with neutropenia or undergoing chemotherapy, tracheotomy were excluded. Beijing Chao-Yang Hospital has the Beijing Institute of respiratory diseases, well-recognized at the national level and the hospital has 1900 beds. All patients underwent ADL evaluation by charge nurse within 2 h after admission. The main outcome was the in-hospital mortality. The patients were divided into Survivors (n = 4749) and Deceased (n = 131) groups depending on vital status at discharge.

The demographic and clinical information data from all patients were extracted from the electronic medical records. The following variables were collected: age, sex, smoking, co-morbidity, clinical symptoms, clinical conditions and laboratory findings on hospital admission.

Results
Characteristics of the study population
A total of 4880 patients diagnosed with CAP and aged ≥65 years were included in the study with a median age 72 years (range, 68–80 years) and over half (59.3%) were male. The characteristics of the 4880 patients in the study are shown in Table 1. One hundred thirty-one patients (2.7%) died during their hospitalization. 69.5% patients in Deceased group had a Barthel Index (BI) < 60. Median BI score in the Deceased group was 45 (20–65). Deceased group had lower BI than Survivors group (p < 0.001).
| Characteristic                        | Total patients | Vital status at discharge | Deceased n = 131 | P value* |
|--------------------------------------|----------------|---------------------------|------------------|----------|
|                                      | n = 4880       | Survivors n = 4749        |                  |          |
| Age, years                           |                |                           |                  |          |
|                                      | 72 (68–80)     | 71 (67–79)                | 75 (70–84)       | < 0.001  |
| Male, n (%)                          | 2895 (59.3)    | 2809 (59.1)               | 86 (65.6)        | 0.149    |
| Barthel index (BI)                   | 75 (65–85)     | 75 (65–85)                | 45 (20–65)       | < 0.001  |
| BI< 60, n (%)                        | 720 (14.8)     | 629 (13.2)                | 91 (69.5)        | < 0.001  |
| BI≥60, n (%)                         | 4160 (85.2)    | 4120 (86.8)               | 40 (30.5)        |          |
| Comorbid conditions, n (%)           |                |                           |                  |          |
| Smoking                              | 2336 (47.9)    | 2268 (47.8)               | 68 (51.9)        | 0.425    |
| COPD                                 | 676 (13.9)     | 656 (13.8)                | 20 (15.3)        | 0.609    |
| Lung cancer                          | 478 (9.8)      | 455 (9.6)                 | 23 (17.6)        | 0.004    |
| Diabetes                             | 1214 (24.9)    | 1172 (24.7)               | 42 (32.1)        | 0.037    |
| Chronic heart failure                | 922 (18.9)     | 893 (18.8)                | 29 (22.1)        | 0.365    |
| Hypertension                         | 2392 (49.0)    | 2321 (48.9)               | 71 (54.2)        | 0.250    |
| Cerebrovascular disease              | 330 (6.8)      | 316 (6.7)                 | 14 (10.7)        | 0.077    |
| Chronic renal failure                | 86 (1.8)       | 83 (1.7)                  | 3 (2.3)          | 0.520    |
| Chronic liver disease                | 146 (2.9)      | 145 (3.1)                 | 1 (0.8)          | 0.188    |
| Clinical symptoms, n (%)             |                |                           |                  |          |
| Fever                                | 1773 (36.3)    | 1715 (36.1)               | 58 (44.3)        | 0.065    |
| Cough and expectoration              | 1014 (20.8)    | 980 (20.6)                | 34 (25.9)        | 0.155    |
| Chest pain                           | 433 (8.9)      | 427 (8.9)                 | 6 (4.6)          | 0.086    |
| Dyspnea                              | 853 (17.5)     | 816 (17.2)                | 37 (28.2)        | 0.002    |
| Duration of symptoms                 | 6.90 ± 8.78    | 6.86 ± 9.03               | 6.93 ± 8.21      | 0.964    |
| Clinical data, n (%)                 |                |                           |                  |          |
| Respiratory failure                  | 210 (4.3)      | 147 (2.4)                 | 63 (48.1)        | < 0.001  |
| NIV/IMV                              | 167 (3.4)      | 115 (2.4)                 | 52 (39.7)        | < 0.001  |
| Impaired consciousness               | 40 (0.8)       | 28 (0.6)                  | 12 (9.2)         | < 0.001  |
| Respiratory rate ≥ 30/min            | 66 (1.4)       | 57 (1.2)                  | 9 (6.9)          | < 0.001  |
| Blood pressure SBP < 90 mmHg         | 26 (0.8)       | 23 (0.5)                  | 3 (2.3)          | 0.050    |
| DBP ≤ 60 mmHg                        | 594 (12.2)     | 570 (12.0)                | 24 (18.3)        | 0.265    |
| BUN≥7 mmol/L                         | 1531 (31.4)    | 1405 (29.6)               | 126 (96.2)       | < 0.001  |
| WBC < 4.0 × 10^9/L or > 10.0 × 10^9/L | 858 (17.6)    | 807 (17.0)                | 51 (38.9)        | < 0.001  |
| PLT < 10.0 × 10^9/L                  | 255 (5.2)      | 210 (4.4)                 | 45 (34.4)        | < 0.001  |
| Pathogens, n (%)                     |                |                           |                  | < 0.001  |
| Bacterial pneumonia                  | 4144 (84.9)    | 4050 (85.3)               | 94 (71.8)        |          |
| Viral pneumonia                      | 62 (1.3)       | 56 (1.2)                  | 6 (4.6)          |          |
| Fungal pneumonia                     | 146 (3.0)      | 129 (2.7)                 | 17 (12.9)        |          |
| Mycoplasma pneumonia/Chlamydial pneumonia | 528 (10.8) | 514 (10.8)                | 14 (10.7)        |          |

Data are presented as median (interquartile range), mean (standard deviation) or n (%)
COPD chronic obstructive pulmonary disease, IMV invasive mechanical ventilation, NIV non-invasive ventilation, SBP Systolic blood pressure, DBP Diastolic blood pressure, BUN blood urea nitrogen, WBC white blood cell, PLT platelet
*For comparisons between Survivors group and Deceased group
The ADL upon admission and in-hospital mortality in older patients with CAP

In-hospital mortality was higher among patients with worse ADL upon admission (BI<60) compared to those BI≥60 (12.6% vs. 0.9%).

For all clinical presentation presented in Table 1, we initially evaluated each variable that displayed statistical significance with \( p < 0.05 \) in difference between Deceased and Survivors group using univariate analysis (Table 2). Input variables for the multivariate model were selected from significant variables obtained from the univariate analysis. Dyspnea was not included in multivariate model to eliminate the influence of multicollinearity.

In logistic regression model (Table 2), the worse ADL upon admission (BI<60) was associated with an increase in the risk of death during CAP hospitalization, worse ADL upon admission (BI<60) showed an OR for in-hospital mortality of 7.53 (95%CI: 2.77–20.48; \( P < 0.01 \)). This association remained significant after adjustment for Age, Comorbid conditions, Respiratory failure, Pathogens, White blood cell and Blood urea nitrogen (OR, 3.74; 95%CI, 2.37–5.91; \( P < 0.01 \)).

**Discussion**

The main finding in this study was the activity of daily living upon admission is an independent predictor of in-hospital mortality in older patients with community-acquired pneumonia. We noted that worse ADL upon admission (BI<60) was associated with a nearly 7 times increase in the risk of death during hospitalization (OR, 7.53; 95%CI, 2.77–20.48; \( P < 0.01 \)). This association remained significant after adjustment for age, comorbid conditions, respiratory failure, pathogens, WBC and BUN (OR, 3.74; 95%CI, 2.37–5.91; \( P < 0.01 \)). ROC curve revealed that ADL upon admission is an important predictor of in-hospital mortality in older patients. The prognostic value of ADL function at admission was good, as shown by the ROC curve. Hitherto, association between ADL function upon admission and in-hospital mortality in older patients with pneumonia has not been studied, and our study had relatively adequacy sample size. Our study may represent useful information for planning of clinical strategies in older patients hospitalization for CAP.

Older patients hospitalized with CAP are at high risk for short-term mortality [5]. The in-hospital mortality observed in this study was 2.7%. Previous literature reported the in-hospital mortality of 1 to 5% [5, 23–26], with two multicenter studies reporting rates 2.2% [5, 23], our result is consistent with previous literature.

**Table 2** Logistic regression analysis for in-hospital mortality

| Variables                        | Univariate Analysis | Multivariable Analysis |
|----------------------------------|---------------------|------------------------|
|                                  | OR (95% CI)         | \( P \) value          | OR (95% CI)         | \( P \) value          |
| Age                              | 1.70 (1.08–2.66)    | 0.03                   | 1.58 (0.95–2.63)    | 0.09                   |
| Male                             | 1.01 (0.68–1.49)    | 0.38                   |                       |                       |
| BI< 60                           | 7.53 (2.77–20.48)   | \(< 0.01\)             | 3.74 (2.37–5.91)    | \(< 0.01\)             |
| Lung cancer                      | 3.05 (2.24–4.14)    | 0.01                   | 2.38 (1.35–4.21)    | 0.03                   |
| Diabetes                         | 1.54 (1.06–2.23)    | 0.02                   | 1.26 (0.99–1.59)    | 0.07                   |
| Dyspnea                          | 1.51 (1.19–1.92)    | 0.04                   |                       |                       |
| Respiratory failure              | 17.81 (12.04–26.36) | \(< 0.01\)             | 7.54 (3.72–15.27)   | \(< 0.01\)             |
| Impaired consciousness           | 1.47 (0.72–3.98)    | 0.16                   |                       |                       |
| Respiratory rate                 | 1.01 (0.99–1.03)    | 0.17                   |                       |                       |
| BUN                              | 5.38 (3.16–9.11)    | \(< 0.01\)             | 2.61 (1.51–4.52)    | 0.01                   |
| WBC                              | 2.36 (1.42–3.93)    | \(< 0.01\)             | 1.68 (1.30–2.17)    | 0.02                   |
| PLT                              | 1.36 (0.87–2.13)    | 0.11                   |                       |                       |
| Viral or Fungal pneumonia        | 5.58 (3.16–9.85)    | \(< 0.01\)             | 2.25 (1.25–4.05)    | 0.01                   |

\( OR \) odds ratio, \( CI \) confidence interval, BI Barthel Index, BUN blood urea nitrogen, WBC white blood cell, PLT platelet
Identifying predictor factors for pneumonia in older patients is crucial in clinical decision making [27–29]. The death of frail elderly with pneumonia is not frequently only due to pneumonia itself [5, 30]. Jason P found that early and aggressive management measures, implemented and valuation of prognosis within 24 h decrease mortality in severe CAP [31]. Different prognostic scales have been documented to assess in CAP, the most commonly be used are the PSI and CURB-65. However, both the PSI and the CURB-65, in contrast to our study data on ADL functional status was lacking. Increasing age is considered to be a factor affecting clinical outcomes in previous study [2, 32]. It is noteworthy that the predictive value of ADL upon admission was better than age in our study. Functional status in elderly patients are frequently individualized [33]. Increasing age does not always mean poor functional status, on the other hand, some older adults experienced a more significant decline in functional status than their peers. The functional status, is more important than age in predicting prognosis in the elderly.

There is growing evidence that poor ADL function is associated with increased mortality. Even little changes in the ADL function could lead to poor clinical outcomes. ADL functional status has been shown to be an independent predictor of mortality in heterogeneous populations [7–19]. In CAP, a worse ADL is directly related to increased short-term and long-term mortality [9]. It was reported that a BI level<80 was associated with 30-day mortality in pneumonia patients [8] and a low BI with increased mortality in institutionalized patients [7]. On the other hand, a high BI level has been reported was related to reduced 30-day and 18 months mortalities in elderly CAP patients [10]. It was found that BI was one of the risk factors for 6 month mortality in COPD patients [11]. While, a worse baseline BI was reported associated with greater mortality in elderly patients admitted to the emergency because of fever [14]. Our study show the same trend as all these previous studies, the worse ADL upon admission (BI< 60) was associated with an increase in the risk of death during CAP hospitalization. Activity of daily living upon admission is an independent predictor of in-hospital mortality in older patients with community-acquired pneumonia.

Assessment of ADL at admission could potentially be used in further management of CAP in older patients.
Barthel Index (BI) can effectively performed to evaluate ADL [34]. Barthel Index is a widely used functional assessment of ADL. BI is the official ADL tool of geriatric patients. All patients admitted to ward were evaluated in our hospital. The BI is a simple and rapid assessment tool with high reliability. It takes approximately 5 min to carry out, and it is easy to interpret in clinical practice. The Barthel Index (BI) is reliable, simple, and it can be used as a conventional method for the assessment of the ADL functional status upon admission in patients with CAP.

The study has some limitations. First, this was a single-center retrospective study in an urban area, the potential bias may be due to the single-center design. Secondly, the environment may have some influence on BI. Thirdly, it is unknown if measures such as functional exercise or nutrition supplementation could improve activities of daily living function to improve the prognosis, future research will be required.

Conclusions
Activity of daily living upon admission is an independent predictor of in-hospital mortality in older patients with community-acquired pneumonia.

Abbreviations
CAP: Community-acquired pneumonia; ADL: Activity of daily living; BI: Barthel index; IDSA/ATS: Infectious Diseases Society of America and the American Thoracic Society; OR: The odds ratios; CI: Confidence intervals; ROC: The receiver operating characteristic curve; AUC: The areas under the curves; COPD: Chronic obstructive pulmonary disease; SBP: Systolic blood pressure; DBP: Diastolic blood pressure; BUN: Blood urea nitrogen; WBC: White blood cell; PLT: Blood platelet; IMV: Invasive mechanical ventilation; NIV: Non-invasive ventilation; PSI: Pneumonia severity index

Supplementary Information
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Additional file 1: Supplementary Table 1. Diagnostic criteria for CAP.

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Authors’ contributions
YK conceptualized and designed the study, collected data, drafted the initial manuscript, and reviewed and revised the manuscript. XYF carried out the study. YK conceptualized and designed the study, collected data, drafted the initial manuscript, and reviewed and revised the manuscript. XYF carried out the study.

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Availability of data and materials
Raw data is available from the corresponding author on reasonable request.

Declarations

Ethics approval and consent to participate
The study protocol was approved by the Institutional Review Board for Human Studies of Beijing Chaoyang Hospital, Beijing, China. The review board exempted the acquisition of informed consent because this was a retrospective study. Patients’ data confidentiality was fully respected during data collection and the preparation of the manuscript.

Consent for publication
Not applicable – this study does not contain any patient personal details.

Competing interests
The authors declare they have no competing interests.

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References

1. GBD 2015 Mortality and Causes of Death Collaborators. Global, regional, and national life expectancy, all-cause mortality, and cause-specific mortality for 249 causes of death, 1980–2015: a systematic analysis for the Global Burden of Disease Study 2015. Lancet. 2016;388:1459–544.
2. Vila CA, Ochoa GO, Rodriguez BT, et al. EPIVAC study group. Epidemiology of community-acquired pneumonia in older adults: a population-based study. Respir Med. 2009;103:309–16.
3. Cillóniz C, Polverino E, Evog S, et al. Impact of age and comorbidity on cause and outcome in community-acquired pneumonia. Chest. 2013;144:999–1007.
4. Ochoa GO, Vila CA, De DC, et al. The burden of community-acquired pneumonia in the elderly: the Spanish EVAN-65 study. BMC Public Health. 2008;8:222.
5. Grant W, Wesley H, Mark C, et al. In-hospital deaths among adults with community-acquired pneumonia. Chest. 2016;154(3):628–35.
6. De Morton NA, Keating JL, Davidson M. Barthel index in the assessment of hospitalized older patients after admission for an acute medical condition. Arch Phys Med. 2008;89:641–7.
7. Nakazawa A, Nakamura K, Kitamura K, et al. Association between activities of daily living and mortality among institutionalized elderly adults in Japan. J Epidemiol. 2012;22(6):501–7.
8. Murcia J, Llorens P, Sánchez-Payá J, et al. Functional status determined by Barthel index predicts community-acquired pneumonia mortality in general population. J Inf Secur. 2010;61:4.
9. Shiao CC, Hsu HC, Chen IL, et al. Lowened witr Barthel index is associh higher risk of hospitalization-requiring pneumonia in long-term care facilities. Tohoku J Exp Med. 2015;236:281–8.
10. Torres OH, Munoz J, Rui D, et al. Outcome predictors of pneumonia in elderly patients: importance of functional assessment. J Am Geriatr Soc. 2004;52:1603–9.
11. Ranieri P, Bianchetti A, Margiotta A, et al. Predictors of 6-month mortality in elderly patients with mild chronic obstructive pulmonary disease discharged from a medical ward after acute non acidotic exacerbation. J Am Geriatr Soc. 2008;56:909–13.
12. Matsen LE, Jepsen DB, Ryg I, et al. Functional level at admission is a predictor of survival in older patients admitted to an acute geriatric unit. BMC Geriatr. 2012;12:32.
13. Yalçınli S, Ersel M, Karbek Akarca F, et al. Can Barthel index predict mortality in geriatric patients admitted to the emergency department with a high fever? Turk J Geriatr. 2015;18(4):266–72.
14. Bo M, Massaia M, Raspo S, et al. Predictive factors of in hospital mortality in older patients admitted to a medical intensive care unit. J Am Geriatr Soc. 2003;51:529–33.
15. Jo T, Yasunaga H, Sasabuchi Y, et al. Association between dementia and discharge status in patients hospitalized with pneumonia. BMC Pulm Med. 2017;17:128.
16. Pacho C, Domingo M, Nuñez R, et al. Predictive biomarkers for death and rehospitalization in comorbid frail elderly heart failure patients. BMC Geriatr. 2018;18(1):109.

17. Wong TM, Leung FKL, Lau TW, et al. Effectiveness of a day rehabilitation program in improving functional outcome and reducing mortality and readmission of elderly patients with fragility hip fractures. Geriatr Orthop Surg Rehabil. 2018;2151459318759355.

18. Nurul I, Mohd S, Mohamed S, et al. Activities of daily living and determinant factors among older adult subjects with lower body fracture after discharge from hospital: a prospective study. Int J Environ Res Public Health. 2018;15:1002.

19. Mangram AJ, Mitchell CD, Shifflette VK, et al. Geriatric trauma service: a one year experience. J Trauma Acute Care Surg. 2012;72:119–22.

20. Mahoney FI, Barthel DW. Functional evaluation: the Barthel index. State Med J. 1965;1461–5.

21. Respiratory Society of Chinese Medical Association. Guidelines for the diagnosis and treatment of community-acquired pneumonia in Chinese adults. Clin J Tuberc Respir Dis. 2016;39(4):253–79.

22. Mandell LA, Wunderink RG, Anzueto A, et al. Infectious Diseases Society of America/American Thoracic Society consensus guidelines on the management of community-acquired pneumonia in adults. Clin Infect Dis. 2007;44:527–72.

23. Adamuz J, Eulàlia M, Gorza M, et al. Care complexity individual factors associated with adverse events and in-hospital mortality. PLoS One. 2020;15(7):e0236370.

24. Flaatten H, Brattebø G, Alme B, et al. Adverse events and in-hospital mortality: an analysis of all deaths in a Norwegian health trust during 2011. BMC Health Serv Res. 2017;17:1–7.

25. Conway R, Byrne D, Riordan D, et al. Outcomes in acute medicine-evidence from extended observations on readmissions, hospital length of stay and mortality outcomes. Eur J Intern Med. 2019;66:69–74.

26. Uematsu H, Kunisawa S, Yamashita K, et al. The impact of patient profiles and procedures on hospitalization costs through length of stay in community-acquired pneumonia patients based on a Japanese administrative database. PLoS One. 2015;10:e0125284.

27. Manisha JM, Nathalie DR, Heather A, et al. Modifiable risk factors for pneumonia requiring hospitalization among community-dwelling older adults: the health, aging, and body composition study. J Am Geriatr Soc. 2013;61(7):1111–8.

28. Loeb M, Neupane B, Walter SD, et al. Environmental risk factors for community-acquired pneumonia hospitalization in older adults. J Am Geriatr Soc. 2009;57:1036–40.

29. Jackson ML, Nelson JC, Jackson LA, et al. Risk factors for community-acquired pneumonia in immunocompetent seniors. J Am Geriatr Soc. 2009;57:882–8.

30. Mortensen EM, Coley CM, Singer DE, et al. Causes of death for patients with community-acquired pneumonia: results from the pneumonia patient outcomes research team study. Arch Intern Med. 2002;162:1059–64.

31. Jason P, Nathan CD, Qi G, et al. Severe community-acquired pneumonia: timely management measures in the first 24 hours. Crit Care. 2016;20:237.

32. Prina E, Ranzani OT, Torres A, et al. Community-acquired pneumonia. Lancet. 2015;386:1097–108.

33. Na L, Streim JE. Psychosocial well-being associated with activity of daily living stages among community-dwelling older adults. Gerontol Geriatric Med. 2017;3:2333721417700011.

34. Milicic ME, Feng MC. Assessment of activities of daily living, self-care, and independence. Arch Clin Neuropsychol. 2016;31(6):506–16. https://doi.org/10.1093/arclin/acw049.

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