Mild Cognitive Impairment Predicts Institutionalization among Older Men: A Population-Based Cohort Study

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Abstract

Background: There is a lack of evidence on the contribution of mild cognitive impairment (MCI) to institutionalization in older adults. This study aimed to evaluate a range of risk factors including MCI of institutionalization in older men.

Methods: Men aged ≥70 years (n = 1705), participating in the Concord Health and Ageing in Men Project, Sydney, Australia were studied. Participants completed self-reported questionnaires and underwent comprehensive clinical assessments during 2005–2007. Institutionalization was defined as entry into a nursing home facility or hostel at any time over an average of 5 years of follow-up. Cox regression analysis was conducted to generate hazard ratios (HR) with 95% confidence intervals (CI).

Results: A total of 125 (7.3%) participants were institutionalized. Piecewise Cox proportional models were generated and divided at 3.4 years (1250 days) of follow-up due to violation of the proportional hazards assumption for the association between MCI and institutionalization ($\chi^2 = 6.44, p = 0.01$). Dementia, disability in Activities of Daily Living (ADL) and Instrumental Activities of Daily Living (IADL), poor grip strength, few social interactions, being a Non-English speaking immigrant and age were predictive of institutionalization during both time periods, whereas MCI (HR = 4.39, 95%CI 2.17–8.87) only predicted institutionalization in the period beyond 3.4 years of follow-up. Being married (HR = 0.42, 95%CI: 0.24–0.72) was protective only during the period after 3.4 years of follow-up.

Discussion: In this study, the strongest predictors of institutionalization were dementia, MCI, ADL and IADL disability. MCI was not a predictor of early institutionalization but became a significant predictor beyond 3.4 years of follow-up.

Introduction

Delaying older adults’ transition from living in the community to institutionalization is of major public health importance. It is also important for older individuals themselves, most of whom would prefer to remain living in the community [1]. A number of studies, conducted across a range of settings (e.g. population-based and dementia-based samples) have investigated risk factors for institutionalization in older adults [1,2]. Dementia, disability in Activities of Daily Living (ADL) and Instrumental Activities of Daily Living (IADL) are the most consistent risk factors for admission to a residential aged care facility (RACF) [1,2]. However, other studies have identified sociodemographic and socioeconomic factors as additional important predictors of institutionalization in older adults [3,4].

While studies that have investigated predictors of institutionalization have looked at severe cognitive impairment, the contribution of mild cognitive impairment (MCI) to institutionalization in older adults is not clear. One study has looked at the association of cognitive impairment not including dementia with adverse outcomes including institutionalization in older adults [5]. However, the diagnosis was based on clinical judgment rather than the use of specific diagnostic criteria.

More research is also needed to understand factors contributing to institutionalization in an ethnically diverse population. There is growing ethnic diversity in older populations in many western countries including Australia, Canada, and the USA [6,7,8]. Cultural differences in values and expectations of family support as well as the availability of culturally appropriate residential aged care services could all contribute to different rates of institutionalization for minority elders.

To our knowledge, no study has been conducted to investigate MCI as a risk factor of institutionalization in older adults. The objective of this study was to investigate a range of risk factors...
including demographics, socioeconomic status, health risk factors, health conditions including MCI, physical performance, medication use and service use as predictors of institutionalization in an ethnically diverse community-based cohort of older men, enrolled in the Concord Health and Ageing in Men Project (CHAMP). Investigating predictors of institutionalization in the CHAMP cohort represents a unique opportunity due to the availability of data including a range of clinical assessments, cognitive assessments, physical performance measures and use of community-based home care services.

Methods

Study Population

Participants were community-dwelling older men, participating in the CHAMP study, an ongoing cohort study in Sydney, Australia [9]. Eligible participants were aged ≥70 years at baseline and living in a specific study area (the Local Government Areas of Burwood, Canada Bay and Strathfield) near Concord Hospital. The only exclusion criterion was living in a RACF. The Electoral Roll was chosen as the sampling frame for the study. Registration on the Electoral Roll is compulsory and regularly updated, making it a suitable population-wide sampling frame. Invitation letters were sent to 3627 men and contact was made with 3005. Most of the 622 men who were not contacted did not have a listed telephone number. One hundred and ninety of the contacted men were not eligible for the study because they had moved out of the study area, moved into a nursing home, or had died. Of the 2815 eligible men contacted, 1511 (53.7%) participated in the study. An additional 194 (11.4%) men living in the study area heard about the study from friends or the local media and were recruited before receiving an invitation letter, giving a final sample of 1705 participants. All participants gave written informed consent. The study was approved by the Human Research Ethics Committee Concord RG Hospital.

Data Collection

Participants underwent baseline assessments that comprised self-completed study questionnaires and a clinical assessment that consisted of physical performance measures, neuropsychological testing and medication inventory. Following the initial baseline assessment, the men were contacted regularly at 4-monthly intervals to enable updating of data on institutionalization. Data collected during baseline assessments (2005–7), including self-reported questionnaire data and clinical information, were used in the current analysis along with longitudinal data on institutionalization.

Ascertainment of Predictor Variables

The main groups of predictor variables included demographic factors, socioeconomic status, health risk factors, health conditions, physical performance measures, medication use and service use. These predictors have been identified based on the clinical significance, and based on previous studies investigating risk factors for institutionalization [1,2].

Sociodemographic factors. Sociodemographic variables included age, marital status (married versus other) and living arrangements (live alone versus live with others). Social support was measured using the shortened Duke Social Support Index (DSSI) which measures both social support satisfaction and social interactions [10]. The first item in the DSSI was modified in the CHAMP which allowed the creation of two separate variables for the number of family and non-family supports. These variables were entered into models separately to the score for social interactions and subjective support. The men were also asked their country of birth which enabled grouping of the men into the categories of Australian-born, overseas-born from an English-speaking country (ESB), and overseas-born from a non-ESB. The CHAMP study area has a high proportion of immigrants and as a result, only 49.8% of men in the CHAMP study were born in Australia and 19.6% were born in Italy.

Socioeconomic status. Socioeconomic status was measured using four separate variables: age at leaving school, main lifetime occupation (managers and professionals versus other), source of income (government pension only versus other) and house ownership.

Health risk factors. Physical activity was assessed using the Physical Activity Scale for the Elderly (PASE) [11]. Participants were asked about whether they had ever consumed alcohol and whether they had consumed at least 12 alcoholic drinks in the past 12 months. This enabled categorization of current non-drinkers into lifelong abstainers and ex-drinkers. For those who consumed at least 12 drinks in the past year, the frequency and quantity of alcohol consumption was assessed, enabling categorization of drinkers as either safe drinkers (1–21 drinks per week) or harmful drinkers (>21 drinks per week) [12]. Smoking status (never smoker, ex-smoker, current smoker) was also assessed.

Health conditions. Data on medical conditions were obtained from the self-reported questionnaires in which participants reported if they had any of the following diseases: diabetes, thyroid dysfunction, osteoporosis, Paget’s disease, stroke, Parkinson’s disease, epilepsy, hypertension, coronary artery disease or myocardial infarction, angina, congestive heart failure, intermittent claudication, chronic obstructive lung disease, liver disease, chronic kidney (renal) disease or kidney (renal) failure, cancer (excluding non-melanoma skin cancers), or arthritis. The number of reported comorbidities was dichotomized at the upper quartile (≥4 versus >4).

Participants also self-reported the presence of shortness of breath, and a history of having fallen in the past 12 months. Data on self-rated health were obtained and dichotomized into excellent/good versus fair/poor/very poor. Corrected visual acuity was assessed using a Bailey-Lovie chart [13] and poor vision was defined as those with <6/19 visual acuity. The presence of incontinence was defined as leaking urine at least two or three times a week. Participants were asked about the presence of chronic pain (pain in the last six months that has lasted for ≥3 months and been experienced every day). Participants were also asked how much pain interfered in their normal activities in the past four weeks as part of the Short Form 12 [14]. Participants were considered to have chronic intrusive pain if they reported the presence of chronic pain and pain that interfered with normal activities moderately, quite a bit or extremely. Depressive symptoms were assessed with the 15-item Geriatric Depression Scale (≥5 indicative of depressive symptoms) [15]. Anxiety symptoms were measured using the Goldberg Anxiety Scale [16], with >5 considered as presence of anxiety.

Diagnosis of cognitive impairment. Participants were screened for cognitive impairment using the Mini Mental State Examination (MMSE) [17] and the Informant Questionnaire on Cognitive Decline (IQCODE) [18] during the baseline clinic assessment. In addition to the cognitive screen participants also completed other cognitive assessments including Addenbrooke’s Cognitive Examination [19], Trail Making Task B [20], Weigl-Colour Form Sorting test [21] and Logical Memory Recall test [22]. Participants with a MMSE less than or equal to 26 and/or IQCODE greater than 3.6 were invited to have detailed clinical assessments by the study geriatrician. This assessment included a
review of medical comorbidities and medications, a standardized neurological assessment, a more detailed informant interview [23] and the Rowland Universal Dementia Assessment Scale (RUDAS) [24]. At a weekly meeting two geriatricians, a neurologist and a neuropsychologist reviewed all medical, cognitive, informant and functional data and reached a final diagnosis of cognitive status for each participant. At the end of the screening and clinical assessments, participants were categorized as having dementia (n = 93), MCI (n = 120), unknown cognitive status (n = 164) or cognitively intact (n = 1328). Participants determined to be cognitively impaired but not demented were given the diagnosis of MCI, if they met the clinical criteria described by Petersen et al 2004 [25]. Although MCI was categorized according to the sub-types defined by Petersen et al, for the purposes of analyses and given small cell sizes, participants with all sub-types of MCI were grouped together. This is consistent with subjects fulfilling the general criteria for MCI [26]. Diagnosis and classification of dementia was based on the Diagnostic and Statistical Manual of Mental Disorders (4th edition) revised criteria and well recognized criteria for dementia subtypes [27,28,29]. All sub-types of dementia were grouped together for analyses given small cell sizes.

Physical function and performance. Functional status was measured with ADL and IADL scales. Disability in ADL was defined as needing help with ≥1 activities included in the modified Katz ADL scale [30]. Disability in IADL was defined as needing help with ≥1 activities included in the OARS IADL scale [31]. Physical performance was assessed by administering a standard performance battery that included the following tasks: (i) walking speed (m/s) over a 6-m course, adjusted for height; (ii) chair stands test-time to successfully complete five chair stands was assessed and time dichotomized at the slowest quartile; (iii) muscle (grip) strength (kg), and (iv) dynamic balance test. Muscle strength was measured using a Jamar dynamometer (Promedics, Blackburn, UK). The score was calculated as the grip strength (kg) of the dominant hand (best of two trials). Dynamic balance was assessed with a coordinated stability task [32]. Scores were dichotomized at the highest (worst) quartile. Participants, who did not complete the tests due to physical inability, were included in the worst quartile. A score of ≥1 was assigned for each of the four measures. Participants were categorized as having a coordinated stability task [32]. Scores were dichotomized at the highest (worst) quartile. Participants, who did not complete the tests due to physical inability, were included in the worst quartile for analyses given small cell sizes.

Medication assessment. Medication data were coded using the Iowa Drug Information Service (IDIS) drug code numbers. Polypharmacy was defined as the use of ≥5 regular prescription medicines [33]. Psychotropic medication use was defined as exposure to the following drug classes: anticonvulsants (IDIS code level 28120000), antidepressants and antipsychotics (IDIS code level 28160000) and anxiolytics (IDIS code level 28240000).

Service use variables. Participants were asked about their use of a number of community services during the past 12 months. These services included: spending at least one day in an aged care day centre, being visited by Home Care to help with personal or household duties, using services of the Community and Aged Care Packages (CACPs), or any service to deliver or prepare meals at home. Participants were categorized as using one or more of these services in the past year versus using none of these services. Participants were also asked whether they had spent at least one night in a hostel or nursing home in the past 12 months and this was entered into models as a separate variable.

Ascertainment of Outcome Variable

Institutionalization was defined as entry into a nursing home facility or hostel at any time during follow-up to 6.38 years. In Australia, there are two main forms of residential aged-care facilities: low-level care facilities (hostels) and high-level care facilities (nursing homes). Self-care retirement villages are not considered to be RACF and so moving into one of these facilities was not considered “institutionalization”. Data on institutionalization were ascertained through a regular phone contact with the participants or their nominated contact person at 4-monthly intervals. While our data does not enable us to discriminate between permanent and respite institutionalization, the majority of admission to aged-care facilities in Australia are permanent [34].

Statistical Analysis

Data are summarized as means (standard deviation) or numbers (proportions). Differences between institutionalized and non-institutionalized participants were compared using the two-sided t-test or χ²-test where appropriate. Initial univariate analyses of the association between the various study measures and institutionalization were conducted using Log-rank tests and examination of survival curves. Tests for linear trends were performed for continuous variables to determine the linearity of their relationship with institutionalization and, hence, whether to enter these variables into models as continuous or categorical variables. Testing for co-linearity between the variables was performed. There was no evidence of correlation between the variables. The appropriate parameterization of continuous variables as either categorical or continuous was also confirmed in the final model by using Akaïke’s Information Criterion (AIC). Univariate Cox regressions were conducted to determine unadjusted hazard ratios for admission to an aged care facility for the various study measures. Variables that had a p<0.25 in univariate analyses were included in the multivariate model with institutionalization as the outcome. Backward stepwise elimination was used to eliminate non-significant variables from the multivariate model. Backward stepwise elimination has an advantage over other methods (eg. forward) as it allows to examine a model with all independent variables as well as the joint predictive capability of all variables. Clinically significant interactions, and interactions identified in previous studies between dementia and urinary incontinence, dementia and falls, and arthritis and pain were examined by adding the interaction terms into the main effect models one at a time and including the significant interaction terms in the final model. None of the interaction terms remained significant in the model. In the final model, the proportional hazards assumption was assessed through use of a time-dependent covariate method, analysis of Schoenfeld residuals plots and graphical methods (eg. survival plots) for each variable. Upon the examination of the results of time-dependent covariate method, Schoenfeld residuals plots and survival curves, it was identified that the MCI covariate was violating the assumptions. To address this, the step function proportional hazards or piecewise Cox model was used to test the effect of MCI on institutionalization. Data were analyzed using SAS version 9.2 (SAS Institute Inc., Cary, North Carolina). The Kaplan-Meier survival curves were generated using SPSS software version 19.0 (SPSS Inc, Chicago, Illinois).

Results

Descriptive characteristics are provided in Table 1. Of 1705 men studied at baseline, a total of 125 (7.3%) were institutionalized during a mean follow-up of 4.94 (range: 0.08–6.38) years. The mean age, social support satisfaction and social interactions were significantly different between institutionalized men compared with non-institutionalized men. The proportion of men institutionalized increased up to the age of 84 years, and then slightly dropped (Figure 1). In relation to health conditions, there were
significant differences in all factors apart from the presence of anxiety symptoms between the two groups.

The multivariate Cox proportional model showed that age, marital status, social satisfaction, social interactions, country of birth, alcohol use, cognitive status, ADL disability, IADL disability, grip strength, and service use were significant predictors of institutionalization. However, the use of a time-dependent covariate and analysis of Schoenfeld residuals demonstrated that MCI violated the proportional hazards assumption ($\chi^2 = 6.44$, $p = 0.01$). Therefore, as the effect of MCI on institutionalization was not stable over entire follow-up time, and the proportional hazards assumption was not valid, the piecewise Cox proportional models were used to test the effect of MCI on institutionalization. The follow-up period was divided at 3.4 years (1250 days) based on examination of the survival curve for MCI and institutionalization (Figure 2).

Table 2 shows the results of the Cox proportional hazards models for the first 3.4 years of follow-up (Model 1) and beyond 3.4 years (Model 2) of follow-up. In the Cox regression model up to 3.4 years of follow-up, age, high social interactions, country of birth, dementia, ADL disability, IADL disability, grip strength and service use were significant predictors of institutionalization. Dementia (HR = 5.43, 95%CI: 3.00–9.81), ADL disability (HR = 3.22, 95%CI: 1.80–5.77), IADL disability (HR = 3.01, 95%CI: 1.32–6.36) and use of services (HR = 2.61, 95%CI: 1.46–4.66) were the most significant predictors of institutionalization. Interestingly, MCI was not associated with institutionalization during this time interval (HR = 0.72, 95%CI: 0.22–2.36). Participants were less likely to be institutionalized if they had high social interactions (HR = 0.48, 95%CI: 0.27–0.85) and were a NESB immigrant (HR = 0.31, 95%CI: 0.16–0.60).

In the Cox regression model for the period beyond 3.4 years of follow-up, age, marital status, social interactions, country of birth, cognitive status, ADL disability, IADL disability, and grip strength were statistically significant predictors of institutionalization. Dementia (HR = 6.05, 95%CI: 2.95–12.44), MCI (HR = 4.39, 95%CI: 2.17–8.87), ADL disability (HR = 2.72, 95%CI: 1.36–5.46) and IADL disability (HR = 2.71, 95%CI: 1.43–5.13) were the most significant predictor factors for institutionalization in this later time period. Married participants (HR = 0.42, 95%CI: 0.24–0.70), NESB immigrants (HR = 0.41, 95%CI: 0.22–0.77), and those with high social interactions (HR = 0.47, 95%CI: 0.26–0.87) were less likely to be institutionalized.

Discussion

In this prospective population-based study, we identified a number of predictors of institutionalization. The strongest predictors were dementia, MCI, ADL and IADL disability. Older adults with dementia had approximately six times the risk of institutionalization compared with those who did not have dementia. The predictive value of MCI changed with the length of follow-up. MCI was a significant predictor of institutionalization beyond 3.4 years of follow-up. In this period, participants with MCI had approximately four times the risk of institutionalization compared with those who were cognitively intact.

The rate of institutionalization of 7.9% in this study is slightly lower when compared to previous studies conducted in population-based settings in Australia, Europe and USA. An Australian study of community-dwelling adults aged ≥60 years, reported an 8.7% permanent nursing home placement over 14 years of follow-up [35]. In a study of adults aged ≥75 years living in Germany, 7.8% of participants were institutionalized during a mean follow-up of 7.6 years [36]. A USA study reported an institutionalization rate of 13.6% over 12 years of follow-up [37]. The difference in the institutionalization rate across studies may be due to the greater ethnic diversity in the CHAMP study. Moreover, older men are generally less likely to be institutionalized as their wives commonly act as their caregiver at home.

The findings of this study are consistent with meta-analyses that have highlighted dementia, ADL and IADL disability as the most important predictors of institutionalization in older adults [1,2,36]. However, our study is the first study to show that MCI is an important predictor of institutionalization. Individuals with MCI are at an increased risk of developing dementia [38,39]. Therefore, our finding that MCI only contributes to an increased risk of institutionalization after more than three years suggests that this increased risk is associated with a progression of the MCI to dementia with an associated increased risk of institutionalization. In the CHAMP population, of the 120 men diagnosed with MCI at baseline, 82 men were re-assessed at Year 2 follow-up. Of these, 12% (n = 10) had progressed to dementia over two years and three of these had been institutionalized. This progression rate of MCI to dementia is similar to other community-based studies [40]. However, other studies have reported higher conversion rates [41].

The prevalence of MCI has been found to be higher amongst older men than women in those living in nursing and veteran care homes [42]. Cognitive impairment, excluding dementia, has been shown to predict adverse outcomes including institutionalization and mortality in older adults [5]. Therefore, delaying the onset of dementia in individuals with MCI may reduce the risk of institutionalization, which is of major clinical and public health importance [43]. A recent study highlighted that not all predictors of institutionalization are robust with varying follow-up periods [44]. Identification of risk factors that predict institutionalization over short versus a longer period of time may inform future interventions to delay institutionalization [44].

We also found a strong relationship between country of birth, and risk of institutionalization. NESB immigrant men were about 70% less likely to be institutionalized compared with both Australian-born men. Different rates of institutionalization between ethnic groups has important implications for the planning of community and RACF services for older people, particularly as the proportion of older persons from an NESB in Australia is...
Table 1. Characteristics of the study population according to institutionalization status.

| Variable                      | Total population (n = 1705)* | Institutionalized (n = 124, 7.3%) | Not institutionalized (n = 1581, 92.7%) | P values |
|-------------------------------|-------------------------------|-----------------------------------|----------------------------------------|----------|
| **Socio-demographic factors** |                               |                                   |                                        |          |
| Age, mean (SD)                | 76.9 (5.5)                    | 81.4 (5.7%)                       | 76.6 (5.3%)                            | <0.001   |
| Currently married             | 1278 (74.9%)                  | 73 (58.9%)                        | 1205 (76.2%)                          | <0.001   |
| Live alone                    | 318 (18.1%)                   | 44 (35.5%)                        | 274 (17.5%)                           | <0.001   |
| Social support satisfaction, high (DSSS score ≥19) | 1294 (76.9%)                  | 69 (57.0%)                        | 1225 (78.5%)                          | <0.001   |
| Social interactions, high (DSSS score ≥9) | 1019 (61.2%)                  | 48 (40.3%)                        | 971 (62.8%)                           | <0.001   |
| Country of birth              |                               |                                   |                                        |          |
| Australia                     | 849 (49.8%)                   | 82 (66.1%)                        | 767 (48.5%)                           |          |
| ESB immigrant                 | 105 (6.2%)                    | 9 (9.9%)                          | 96 (11.2%)                            |          |
| Non-ESB immigrant             | 751 (44.1%)                   | 33 (28.7%)                        | 718 (48.4%)                           | 0.0002   |
| **Socio-economic factors**    |                               |                                   |                                        |          |
| Occupation, professional       | 505 (29.8%)                   | 21 (16.9%)                        | 484 (30.8%)                           | 0.0011   |
| Own house outright            | 1494 (88.9%)                  | 104 (86.0%)                       | 1390 (89.2%)                          | 0.27     |
| Years of education, ≥7 years  | 1429 (84.7%)                  | 115 (94.3%)                       | 1314 (83.9%)                          | 0.002    |
| Source of income, pension only | 773 (45.9%)                   | 66 (54.1%)                        | 707 (45.3%)                           | 0.06     |
| **Health risk factors**       |                               |                                   |                                        |          |
| Physical activity, normal/high (PASE score ≥80) | 1263 (74.9%)                  | 55 (45.5%)                        | 1208 (77.2%)                          | <0.001   |
| Alcohol consumption           |                               |                                   |                                        |          |
| Lifelong non-drinker          | 147 (8.8%)                    | 8 (6.5%)                          | 139 (9.0%)                            |          |
| Ex-drinker                    | 250 (14.9%)                   | 30 (24.4%)                        | 220 (14.2%)                           |          |
| Safe drinker (1–21 drinks per week) | 1151 (68.7%)                  | 77 (62.6%)                        | 1074 (69.2%)                          |          |
| Harmful drinker (>21 drinks per week) | 127 (7.6%)                   | 8 (6.5%)                          | 119 (7.7%)                            | 0.02     |
| Smoking status                |                               |                                   |                                        |          |
| Never smoker                  | 629 (37.3%)                   | 51 (41.8%)                        | 578 (37.0%)                           |          |
| Previous smoker               | 956 (56.7%)                   | 62 (50.8%)                        | 894 (57.2%)                           |          |
| Current smoker                | 101 (6.0%)                    | 9 (7.4%)                          | 92 (5.9%)                             | 0.38     |
| **Health conditions**         |                               |                                   |                                        |          |
| Comorbidities, ≥5             | 237 (14.0%)                   | 30 (24.6%)                        | 207 (13.2%)                           | 0.0005   |
| Urinary incontinence          | 232 (14.0%)                   | 27 (23.3%)                        | 205 (13.3%)                           | 0.003    |
| Visual acuity, low (<6/19)    | 74 (4.5%)                     | 15 (12.5%)                        | 59 (3.9%)                             | <0.001   |
| Chronic intrusive pain        | 223 (13.4%)                   | 24 (20.5%)                        | 199 (12.8%)                           | 0.02     |
| Self-rated health, good or excellent | 1176 (69.9%)             | 68 (57.1%)                        | 1108 (70.9%)                          | 0.002    |
| Shortness of breath           | 210 (13.3%)                   | 24 (19.4%)                        | 186 (11.8%)                           | 0.013    |
| Hearing loss                  | 1027 (61.1%)                  | 85 (70.3%)                        | 942 (60.4%)                           | 0.03     |
| History of falls              | 322 (19.1%)                   | 50 (41.7%)                        | 272 (17.4%)                           | <0.001   |
| Depressive symptoms           | 246 (14.6%)                   | 45 (37.5%)                        | 201 (12.9%)                           | <0.001   |
| Anxiety symptoms              | 123 (7.4%)                    | 10 (8.7%)                         | 113 (7.3%)                            | 0.57     |
| Cognitive status              |                               |                                   |                                        |          |
| Normal                        | 1492 (87.5%)                  | 73 (58.9%)                        | 1419 (89.8%)                          |          |
| Mild cognitive impairment     | 120 (7.0%)                    | 14 (11.3%)                        | 106 (6.7%)                            |          |
| Dementia                      | 93 (5.5%)                     | 37 (29.8%)                        | 56 (3.5%)                             | <0.001   |
| **Physical function and performance** |                       |                                   |                                        |          |
| ADL disability (needing help with ≥1 task) | 141 (8.3%)                  | 40 (32.3%)                        | 101 (6.4%)                            | <0.001   |
| IADL disability (needing help with ≥1 task) | 697 (41.6%)                   | 97 (80.8%)                        | 600 (38.6%)                           | <0.001   |
| Grip strength, poor (lowest quartile and unable) | 486 (28.7%)                  | 71 (58.2%)                        | 415 (26.4%)                           | <0.001   |
| Chair stands, slow (lowest quartile and unable) | 462 (27.7%)                  | 74 (63.3%)                        | 388 (25.0%)                           | <0.001   |
| Walking speed, slow (lowest quartile and unable) | 242 (14.5%)                  | 59 (49.6%)                        | 183 (11.8%)                           | <0.001   |
| Dynamic balance test, poor (lowest quartile and unable) | 478 (29.1%)                  | 70 (60.3%)                        | 408 (26.7%)                           | <0.001   |
Further research is required to confirm whether this difference in rates of admission is due to different cultural values about the role of family in supporting older persons to remain in the community or whether it is due to a relative lack of culturally and linguistically appropriate residential aged care services. Consistent with previous work [45,46], we also found that being married was associated with a reduced risk of institutionalization. Poor grip strength was also an important risk factor for institutionalization. One study has reported an association of weaker grip strength with increased risk of long-term nursing home stay in the unadjusted models only [47]. Interestingly, falls were not associated with an increased risk of institutionalization in our study, which is in contrast to previous studies [48,49]. Urinary incontinence was also not a significant predictor of institutionalization in this population, which is consistent with one study [50].

Table 1. Cont.

| Variable                        | Total population (n = 1705)* | Institutionalized (n = 124, 7.3%) | Not institutionalized (n = 1581, 92.7%) | P values |
|---------------------------------|-----------------------------|----------------------------------|----------------------------------------|----------|
| **Medication use**              |                             |                                  |                                        |          |
| Polypharmacy (≥5 medicines)     | 639 (37.7%)                 | 52 (42.3%)                       | 587 (37.3%)                            | 0.27     |
| Psychotropic medications       | 211 (12.4%)                 | 22 (17.9%)                       | 189 (12.0%)                            | 0.06     |
| **Service use**                 |                             |                                  |                                        |          |
| Stay in nursing home in past year | 29 (1.7%)                  | 9 (7.4%)                         | 20 (1.3%)                              | <0.001   |
| Use of services in past year   | 193 (11.3%)                 | 49 (39.5%)                       | 144 (9.1%)                             | <0.001   |

ADL = Activities of Daily Living [30]; DSSI = Duke Social Support Index [10]; IADL = Instrumental Activities of Daily Living [31]; PASE = Physical Activity Scale for the Elderly [11].

*Missing data not included in percentages.

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Figure 2. Kaplan-Meier survival curves of the time until institutionalization by cognitive status groups.

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but not with another [51] conducted in community-dwelling older people. It may be that factors such as the physical performance measures and urinary incontinence are not significant predictors of institutionalization in models that already include ADL and IADL disability as these composite measures of function are determined in part by physical function and continence status.

The major strengths of the CHAMP study include its representative sampling from the community, detailed assessment of cognitive status, comprehensive, objective and clinically validated physical performance measures, and availability of a range of important risk factors for institutionalization. However, there are some limitations to the present study. We were unable to investigate the association of caregiver characteristics with the onset of institutionalization. Some studies have shown that in addition to participant characteristics, caregiver characteristics are important determinants of nursing home placement for persons with dementia [4,52] while another study found that compared to participant characteristics, caregiver characteristics may not play an important role in predicting institutionalization [53]. This was a study of community-dwelling older men living in a defined geographical location, which may limit the study’s generalizability to men living in other areas. It should be noted that the response rate in the CHAMP study is similar to other comparable cohort studies of this type [9]. Also the findings of this study may not be applicable to older women. Moreover, the validity of self-report data in participants with cognitive impairment may be questionable.

In conclusion, in this population, the strongest predictors of institutionalization were dementia, MCI, ADL and IADL disability. Older adults with dementia had approximately a six times higher risk of institutionalization compared with those who did not have dementia. The contribution of MCI to institutionalization changed with time, with MCI being a significant predictor only beyond 3.4 years of follow-up. Participants with MCI had approximately a four times higher risk of institutionalization compared with those who were cognitively intact. Our findings suggest that in addition to other risk factors, MCI should also be considered when estimating the risks of long term institutionalization in older adults. Delaying the onset of dementia in individuals with MCI may reduce the risk of institutionalization in older adults, which is of major clinical and public health importance.

Author Contributions
Conceived and designed the experiments: DG FS RC LW FB VN DH DGL. Analyzed the data: DG FS. Contributed reagents/materials/analysis tools: DG FS RC LW FB VN DH DGL. Wrote the paper: DG FS.

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Table 2. Final multivariate models for predictors of institutionalization up to 3.4 years and beyond 3.4 years of follow-up.

| Predictors | Hazard Ratio (95% confidence interval) | P value | Hazard Ratio (95% confidence interval) | P value |
|-----------|---------------------------------------|---------|---------------------------------------|---------|
| Age group (80+ vs <80) | 1.90 (1.05–3.45) | 0.04 | 2.37 (1.37–4.08) | 0.002 |
| Marital status (currently married vs not married) | NA | NA | 0.42 (0.24–0.72) | 0.002 |
| Social interactions (high vs low) | 0.48 (0.27–0.85) | 0.01 | 0.47 (0.26–0.87) | 0.02 |
| ESB immigrant vs Australian born | 0.91 (0.35–2.37) | 0.85 | 0.92 (0.32–2.61) | 0.87 |
| NESB immigrant vs Australian born | 0.31 (0.16–0.60) | 0.0005 | 0.41 (0.22–0.77) | 0.005 |
| MCI vs normal | 0.72 (0.22–3.26) | 0.58 | 4.39 (2.17–8.87) | <0.001 |
| Dementia vs normal | 5.43 (3.00–9.81) | <0.001 | 6.05 (2.95–12.44) | <0.001 |
| ADL disability (yes vs no) | 3.22 (1.80–5.77) | <0.001 | 2.72 (1.36–5.46) | 0.005 |
| IADL disability (yes vs no) | 3.01 (1.32–6.86) | 0.009 | 2.71 (1.43–5.13) | 0.002 |
| Grip strength (low vs high) | 2.19 (1.22–3.93) | 0.008 | 1.95 (1.14–5.13) | 0.002 |
| Social interaction (high vs low) | 0.48 (0.27–0.85) | 0.01 | 0.47 (0.26–0.87) | 0.02 |
| Service use (yes vs no) | 2.61 (1.46–4.66) | 0.001 | NA | NA |

ADL = Activities of Daily Living [30]; ESB = English Speaking Background; IADL = Instrumental Activities of Daily Living [31]; MCI = Mild Cognitive Impairment; NESB = Non-English Speaking Background. NA, not applicable = not a significant predictor during the time period.

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