Research Report

Time Investment for Program Implementation to Manage Neuropsychiatric Symptoms: An Observational Longitudinal Study in In-Home and Residential Care Settings

Miharu Nakanishi^{a,*}, Junko Niimura^a, Canan Ziylan^b, Ton TJEM Bakker^{b,c}, Eva Granvik^d, Katarina Nägga^{e,f}, Yumi Shindo^g and Atsushi Nishida^a

^aResearch Center for Social Science & Medicine, Tokyo Metropolitan Institute of Medical Science, Setagaya-ku, Tokyo, Japan

^bResearch Centre Innovations in Care, Rotterdam University of Applied Sciences, Rotterdam, The Netherlands ^cStichting Wetenschap Balans, Rotterdam, The Netherlands

^dCenter of Excellence in Dementia, University Hospital, Malmö, Sweden

^eClinical Memory Research Unit, Department of Clinical Sciences Malmö, Lund University, Malmö, Sweden

[†]Department of Acute Internal Medicine and Geriatrics, Linköping University, Linköping, Sweden

^gNational Center for Geriatrics and Gerontology, Aichi, Japan

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Abstract.

Background: There are no studies on how the same psychosocial dementia care program is adapted to both in-home and residential care settings.

Objective: To evaluate the time investment required by professionals to implement a psychosocial dementia care program to manage neuropsychiatric symptoms.

Methods: A prospective observational study design was used. The program consisted of 1) a one-day training course, 2) three interdisciplinary discussion meetings in five months, and 3) a web-based tool for the continued assessment of neuropsychiatric symptoms. Care professionals implemented the intervention in in-home (19 in-home care management agencies and 14 multiple in-home service providers) and residential care settings (19 group homes and eight nursing homes) in Japan from October 2019 to February 2020. The level of neuropsychiatric symptoms for the participants was evaluated using the Neuropsychiatric Inventory (NPI: 0–144). The time investment was reported by participating professionals. A total of 125 persons with dementia were included at baseline.

Results: Neuropsychiatric symptoms were significantly decreased at the final follow-up in all types of providers (Cohen's $d_{rm} = 0.44-0.61$). The mean (SD) time required for the five-month implementation was 417.9 (219.8) minutes. There was a mean (SD) decrease of 8.6 (14.0) points in the total NPI score among the 103 persons with completed interventions.

*Correspondence to: Miharu Nakanishi, RN, PhD, Research Center for Social Science & Medicine, Tokyo Metropolitan Institute of Medical Science, 2-1-6 Kamikitazawa, Setagaya-ku, Tokyo

156-8506 Japan. Tel.: +81 3 6834 2292; E-mail: mnakanishi-tky@umin.ac.jp.

The time investment was significantly lower in in-home care management agencies than in group homes, and lower in follow-ups than at baseline assessment.

Conclusion: The program implementation may incur a substantial time investment regardless of setting. An additional benefit scheme to reward the time investment would be helpful to encourage implementation until the follow-ups.

Keywords: Behavioral symptoms, dementia, group homes, home care agencies, nursing homes, psychosocial support systems

INTRODUCTION

As the number of people with dementia increases globally [1], dementia has become a public health priority [2, 3]. Dementia has significant social and economic implications in terms of direct medical and social care costs, and the costs of informal care [4-6]. Neuropsychiatric symptoms contribute to the overall costs of dementia care, both in the community [7, 8] and in care homes [9]. Neuropsychiatric symptoms are highly prevalent in persons with dementia [10, 11], resulting in negative health consequences, including lower quality of life [12], increased psychological suffering [13], higher psychotropic drug use [14], shorter time to institutionalization [5], and increased mortality [15]. Furthermore, family caregivers suffer from a higher burden and elevated risk of being sick [13]. As neuropsychiatric symptoms are considered expressions of distress [16-18], psychosocial interventions are globally recommended as first-line treatments to target the underlying causes [2, 19, 20].

Psychosocial interventions of neuropsychiatric symptoms involve the provision of care tailored to the individual's needs, considering their life context [21]. The underlying causes can lie in different life areas, including psychiatric aspects. Therefore, dementia case management models should enable interdisciplinary team approaches, integrate complex networks of care professionals, and respond appropriately to the unmet needs of persons with dementia [2, 20, 22]. The Swedish BPSD-registry was launched in November 2010 to improve the care of persons with dementia by focusing on psychosocial interventions for neuropsychiatric symptoms [15]. The BPSD-registry is based on an interdisciplinary team approach, and was primarily established for use in nursing homes. Adapted forms of the program have been developed in Denmark [23], the Netherlands [24], and Japan [25].

The Japanese adaptation of the BPSD-registry, called DEMBASE[®], has been developed with a focus on care managers of in-home service users

with dementia. The results of a cluster-randomized controlled trial indicated a reduction in neuropsychiatric symptoms upon application of the adapted program [25], with a medium observed effect size (Cohen's $d_{ppc} = 0.52$). Based on the results, the program was implemented for routine dementia care in the Tokyo metropolitan area in April 2018. However, several in-home care managers raised barriers to the interdisciplinary approach, as it is not usually incorporated into their daily practice with in-home service providers [26]. The program is expected to incur substantial time investment, particularly from interdisciplinary discussion meetings to evaluate neuropsychiatric symptoms, identify unmet needs, and establish an action plan. The time investment required across various care settings warrants further investigation to design a rewarding system that can facilitate program implementation. Most psychosocial interventions require minimal to moderate financial investment by family caregivers in in-home settings as well as care professionals in residential care settings [27]. However, there is scarce evidence for care professionals' management of the neuropsychiatric symptoms of dementia in homebased practices. Furthermore, there are no studies on how the same psychosocial dementia care program is adapted to both in-home and residential care settings. Uncovering differences could have global implications for the dissemination of psychosocial interventions for neuropsychiatric symptoms in people with dementia across various care settings.

This study evaluates the time investment required to implement a psychosocial dementia care program for neuropsychiatric symptoms in both in-home and residential care settings. Our research questions were as follows:

(1) Are neuropsychiatric symptoms reduced via the program implementation in both residential and in-home care settings?

(2) Is care professionals' time investment higher in in-home than residential care settings?

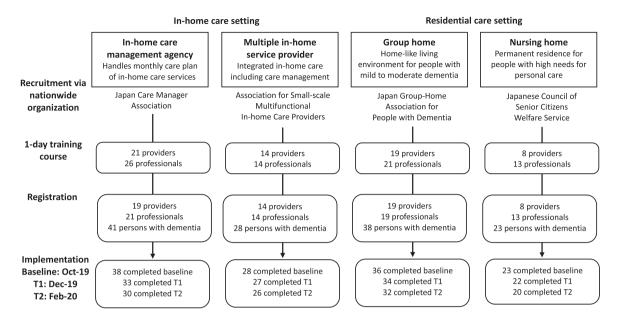


Fig. 1. Participant recruitment, registration, and implementation of the psychosocial dementia care program. Care professionals working as long-term service providers were invited to participate in the program by four nationwide voluntary organizations. Each organization selected candidate providers and asked for participation. The directors of providers asked their employees (care professionals) to participate. In-home care management agencies have care managers who handle monthly care plans for in-home care users and work independently from in-home care service providers such as day care centers and home help. Multiple in-home service providers offer home-based care services to older adults in the community. Group homes offer small-scale, homelike accommodation for residents with mild to moderate dementia. Nursing homes offer permanent residence for older adults with and without dementia who are stable but require regular nursing care. Each provider organization was asked to recruit 20 professionals and 40 persons with dementia.

MATERIALS AND METHODS

Study design

A prospective, observational study design was adopted.

Participants

Care professionals working as long-term service providers were invited to participate in the program by four nationwide voluntary organizations in joint commission with the Ministry of Health, Labour and Welfare (Fig. 1). Each organization selected candidate providers and asked for participation within three regions (Chiba, Kanagawa, and Saitama) around the Tokyo metropolitan area. The three regions were selected to enable employees to participate in the training course held in Tokyo and return home the same day. The directors of providers asked their employees (care professionals) to participate.

Multiple in-home service providers, group homes, and nursing homes were sampled for comparison with in-home care management agencies. In-home care management agencies have care managers who handle monthly care plans for in-home care users and work independently from in-home care service providers such as day care centers and home help. Multiple in-home service providers offer homebased care services to older adults in the community. Group homes offer small-scale, homelike accommodation for residents with mild to moderate dementia. Nursing homes offer a permanent residence for older adults with and without dementia who are stable but require regular nursing care. Nursing home residents have greater personal care needs but lower healthcare needs than group home residents [28]. In-home care service users are those in-home care management agencies or multiple in-home service providers. Residential care settings in this study comprised group homes and nursing homes. We hypothesized that in-home care management agencies would invest more time in program implementation than the other three types of providers, because they provide care management to users independently from service providers, thus increasing the difficulty of holding interdisciplinary discussion meetings [26].

The sample size was calculated for the analysis of variance in time investment for repeated measures between factors, using G*Power 3.1.9.7 [29, 30]. Assuming an alpha level of 0.05 and 95% power, 132 participants were required to observe an effect size of 0.3 (Cohen's f, medium) in time investment. To allow for a 16% dropout rate based on a previous study in Tokyo [25, 31], the sample size was set at 157 persons with dementia. Therefore, each provider organization was asked to recruit 20 professionals and 40 persons with dementia, to collate a total of 160 persons with dementia at baseline. It was intended that each type of provider has an equal number of participants.

Intervention

The program comprised 1) a one-day training course, 2) interdisciplinary discussion meetings, 3) a web-based tool for ongoing behavioral assessments, and 4) a three-hour debriefing meeting.

Training course

The training course guided 1) the process of interdisciplinary meetings for the evaluation of neuropsychiatric symptoms, specification of unmet needs using a 23-item checklist, and establishment of an action plan using an interdisciplinary approach; 2) implementation of the action plan; and 3) use of a web-based tool. The training course was based on consideration of neuropsychiatric symptoms as communicating unmet needs [16-18]. Psychosocial interventions with goal-setting, such as providing pleasant activities, outdoor activities, and removal of environmental triggers were recommended to address these unmet needs based on global evidence [22]. The web-based tool was explained to each participating professional during the training course, held at the end of September 2019.

Interdisciplinary discussion meeting

Once the training course was completed, the professionals held an interdisciplinary discussion meeting with other care professionals to evaluate the neuropsychiatric symptoms of each participant with dementia, to specify their unmet needs, and to establish an interdisciplinary action plan to meet those needs. The medications prescribed to each participant with dementia were also assessed. These components were included to promote a plan-dostudy-act (PDSA) cycle, developed according to the team-based dementia case management model. The PDSA cycle is widely used as a quality improvement method in healthcare settings [32]. If the level of neuropsychiatric symptoms assessed was not reduced during follow-up evaluations, care professionals reviewed unmet needs and revised the action plan during the discussion meeting. Unmet needs, contents of the action plan, and type of caregivers involved in the meeting were categorized and recorded in the web-based tool (Supplementary Table 1). The categories of unmet needs were developed in reference to the findings on associations between neuropsychiatric symptoms and basic physical needs [33] and environmental sources of discomfort [34, 35].

Web-based tool

The web-based tool provided a visualization of longitudinal changes in neuropsychiatric symptoms measured by the Neuropsychiatric Inventory to inform interdisciplinary decision-making. The professionals input the information collected during the discussion meeting using the web-based tool. The individual characteristics of persons with dementia were recorded at registration, including birth year and month, sex, and type of dementia.

Debriefing meeting

Participating professionals attended a three-hour debriefing meeting in mid-November 2019, six weeks after the training. They were divided into groups of four to six members to share their experiences with the program.

Data collection

A questionnaire completed on the day of training evaluated the attitudes towards persons with dementia, self-reported competence in dementia care, and care professionals' personal characteristics. The care professionals responded to the questionnaire before the training course.

Care professionals assessed the neuropsychiatric symptoms and prescribed medication for each person with dementia. They input the information on the web-based tool at baseline (October 2019), after two months (T1 = December 2019), and after four months (T2 = February 2020). Care professionals also reported, via paper questionnaire, the time investment required to implement the program per person at each time of assessment.

Ethical considerations

The study protocol was approved by the Ethics Board of the Tokyo Metropolitan Institute of Medical Science (number 19–26). The research was conducted in accordance with the principles of the Declaration of Helsinki (version November 2013) and is in agreement with the law regarding medicalscientific research in humans. Written informed consent was obtained from each person with dementia and/or by proxy (typically, next of kin) as appropriate. This trial is registered at the UMIN Clinical Trials Registry (UMIN000037254).

Measurements

Assessment of persons with dementia

Information was made available via the web-based tool. We collected information on neuropsychiatric symptoms and prescribed medications. The Neuropsychiatric Inventory - Nursing Home version (NPI-NH) was used to assess the incidence, frequency, and severity of neuropsychiatric symptoms. The original NPI-NH comprised 12 items to rate the frequency and severity of neuropsychiatric symptoms in persons with dementia [36-39]. Scores for each item range from 0 to 12, with higher scores indicating more severe symptoms. Frequency and severity scores are multiplied to determine a total score ranging from 0 to 144. The Japanese version of the NPI-NH has good validity and reliability [40]. The drug name and daily dosage were recorded for each prescribed medication. In this study, the presence of prescribed medication for the nervous system was used for analysis based on the Anatomical Therapeutic Chemical classification. Percentages of the prescriptions were calculated for analgesics (N02), antipsychotics (N05A), anxiolytics (N05B), hypnotics and sedatives (N05C), antidepressants (N06A), and anti-dementia drugs (N06D).

Time investment

Care professionals completed a questionnaire at baseline, T1, and T2 to report the time required to implement the program per person with dementia. The program implementation comprised 1) preparation, 2) transportation, 3) meeting, and 4) management of the web-based tool (Supplementary Table 2). The care professionals also recorded the number of other professionals who had participated in the discussion meeting, and if any professionals had been invited but declined to participate. If so, the reason for declining was also reported (Supplementary Table 3).

Assessment of care professionals

Care professionals completed a questionnaire on the day of the training course, which evaluated their self-reported competence in dementia care and their personal characteristics. Competence in dementia care was measured using the Japanese version of the Sense of Competence in Dementia Care Staff (SCIDS) scale [41], comprising 17 items on a fourpoint Likert scale ranging from '1 = not at all' to '4 = very much'. The total score ranges from 17 to 68, with higher scores indicating a higher sense of confidence. The Japanese version of the SCIDS has been validated [42]. Care professionals' personal characteristics comprised age, sex, primary work qualification, tenure in care for older adults, and educational attainment. Primary work qualification was classified into 1) care manager/social worker, or 2) nurses or other direct care workers in the multivariate analysis.

Statistical analysis

Differences in personal and professional characteristics at baseline were assessed between four provider types (in-home care management agency, multiple in-home service, group home, and nursing home). One-way analyses of variance were used for continuous variables, and chi-square tests were used for categorical variables.

Changes in neuropsychiatric symptoms per type of provider were examined using multilevel linear regression analysis, including persons with dementia and care professionals as random effects, and time of assessment as fixed effects. Panel-data format was adopted in which the same person with dementia could appear two or three times (data at baseline, T1, and T2) to utilize information on cessation cases during follow-ups. The within-subject effect size was calculated using Cohen's d_{rm} for the NPI-NH total score. The effect size was considered low if d values varied by approximately 0.20, medium if approximately 0.50, and large if greater than 0.80 [43, 44].

Multilevel linear regression analysis was also executed to examine the associations between time investment and type of provider. The model included other independent variables from professional characteristics (primary work qualification, tenure in care for older adults, and competence in dementia care), characteristics of persons with dementia (age, sex,

Baseline, Oct 2019 N (%)	In-hon	ne care	Reside	ntial care	Test statistic	p	
or mean (SD)	Care management $(N=38)$	Multiple in-home $(N=28)$	Group home $(N=36)$	Nursing home $(N=23)$			
Age, y	85.1 (5.6)	85.1 (6.0)	84.0 (7.3)	86.7 (7.0)	F(3) = 0.81	0.489	
Sex, male	14 (36.8)	5 (17.9)	6 (16.7)	5 (21.7)	$\chi^2(3) = 5.14$	0.162	
Alzheimer's disease	28 (73.7)	16 (57.1)	26 (72.2)	16 (69.6)	$\chi^2(3) = 2.40$	0.494	
Prescribed medication							
Anti-dementia drugs (N06D)	17 (44.7)	15 (53.6)	22 (61.1)	11 (47.8)	$\chi^2(3) = 2.19$	0.534	
Antipsychotics (N05A)	5 (13.2)	8 (28.6)	6 (16.7)	3 (13.0)	$\chi^2(3) = 3.19$	0.363	
Analgesics (N02)	1 (2.6)	5 (17.9)	4 (11.1)	2 (8.7)	$\chi^2(3) = 4.44$	0.217	
Hypnotics and sedatives (N05C)	2 (5.3)	2 (7.1)	5 (13.9)	1 (4.3)	$\chi^2(3) = 2.53$	0.470	
Antidepressants (N06A)	1 (2.6)	2 (7.1)	5 (13.9)	1 (4.3)	$\chi^2(3) = 3.88$	0.275	
Anxiolytics (N05B)	3 (7.9)	0 (0.0)	4 (11.1)	0 (0.0)	$\chi^2(3) = 5.47$	0.140	
Neuropsychiatric symptoms (0–144)	18.7 (16.1) ^a	25.2 (17.7)	28.8 (18.2)	32.9 (19.2) ^a	F(3) = 3.64	0.015	

 Table 1

 Baseline characteristics of persons with dementia

SD, standard deviation. Level of neuropsychiatric symptoms was assessed using the Japanese version of the Neuropsychiatric Inventory-Nursing Home version (NPI-NH). ^aSignificant difference between the same alphabet letter at p < 0.05, Bonferroni correction.

type of dementia, use of anti-dementia drugs and antipsychotics), and level of neuropsychiatric symptoms. The interclass correlation coefficient (ICC) for care professional was calculated to evaluate the contribution of the difference of professionals to the variance in time investment. A sensitivity analysis was employed using the multilevel model where the dependent variable comprised each category of time investment rather than total minutes. Time investment for transportation was not included as the average number of minutes was small (4.2–5.8).

Statistical significance was considered at an overall $\alpha = 0.05$. Adjustments for multiple testing included the Bonferroni correction with a significance threshold at 0.0125 for neuropsychiatric symptoms per type of provider. All statistical analyses were conducted using STATA version 16.1 (StataCorp, Texas).

RESULTS

Characteristics of persons with dementia

Registration

There were 130 persons with dementia registered in the web-based tool. Of these, 125 received a complete baseline assessment; 5 did not receive baseline assessments due to consent withdrawal (n = 2), hospital admission (n = 1), acute deterioration of physical health (n = 1), and the care professional's change in place of work (n = 1). Those who completed the baseline assessment had a mean age of 85.1 (SD = 6.5) years and included 30 men (24.0%). Eighty-six persons had a diagnosis of Alzheimer's disease (68.8%); 65 had been prescribed anti-dementia drugs (52.0%); and 22 had been prescribed antipsychotics (17.6%). These personal characteristics at baseline did not differ by type of provider except for the level of neuropsychiatric symptoms. The average total NPI-NH score varied between 18.7 in in-home care management agencies and 32.9 in nursing home residents (Table 1).

Follow-ups

Of the 125 persons with dementia at baseline, 116 received a complete T1 assessment, and 108 received a complete T2 assessment (Fig. 1). Seventeen persons did not receive complete follow-ups due to an unavailable schedule for the discussion meeting (n = 6), hospital admission (n = 5), change in care professional's place of work (n = 3), acute deterioration in physical health (n = 2), or death (n = 1) during the period. The 17 cessation cases did not differ from the remaining 108 cases in baseline characteristics or type of provider.

Characteristics of care professionals

Registration

A total of 74 care professionals from 62 providers participated in a 1-day training course for the program in Tokyo. Although the sample size was lower than expected, we concluded the recruitment by mid-September due to Typhoon Faxai. Of the 74 participating professionals, 7 declined to implement the program as they had been affected by Typhoon Hagibis and its aftermath in early October 2019. The remaining 67 professionals from 60 providers registered 130 persons with dementia in the program at baseline (Fig. 1). These 67 professionals completed the three-hour debriefing meeting. On average,

Baseline, Sep 2019 N (%)	In-home	e care	Resider	tial care	Test statistic	р	
or mean (SD)	Care management (N=21)	Multiple in-home (N=14)	Group home (N=19)	Nursing home (N=13)			
Age, y	51.2 (9.0) ^a	44.1 (6.3)	46.9 (9.1) ^b	37.5 (12.3) ^{a,b}	F(3) = 6.08	0.001	
Sex, male	5 (23.8)	5 (35.7)	13 (68.4)	6 (46.2)	$\chi^2(3) = 8.51$	0.037	
Primary qualification for work					$\chi^2(12) = 49.23$	< 0.001	
Professional who handles care							
plan/care coordination							
Care manager	21 (100.0)	5 (35.7)	2 (10.5)	2 (15.4)			
Social worker	0 (0.0)	0 (0.0)	1 (5.3)	0 (0.0)			
Professional who provides care							
Nurse	0 (0.0)	1 (7.1)	0 (0.0)	0 (0.0)			
Direct care worker other than nurse	0 (0.0)	8 (57.1)	16 (84.2)	10 (76.9)			
OT / PT	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)			
Other, unspecified	0 (0.0)	0 (0.0)	0 (0.0)	1 (7.7)			
Educational attainment					$\chi^2(9) = 8.76$	0.460	
Junior high school	1 (4.8)	0 (0.0)	0 (0.0)	0 (0.0)			
High school	4 (19.0)	3 (21.4)	6 (31.6)	4 (30.8)			
College, vocational school	9 (42.9)	4 (28.6)	4 (21.1)	7 (53.8)			
University	7 (33.3)	7 (50.0)	9 (47.4)	2 (15.4)			
Tenure in care for older adults, month	231.1 (74.1) ^{a,b,c}	151.4 (81.6) ^a	134.7 (81.5) ^b	117.5 (74.5) ^c	F(3) = 7.75	< 0.001	
Competence in dementia care (17–68)	40.0 (7.6)	40.4 (6.1)	40.8 (6.0)	43.2 (7.6)	F(3) = 0.63	0.598	

Table 2 Characteristics of care professionals

SD, standard deviation. Competence in dementia care was measured by the Japanese version of the Sense of Competence in Dementia Care Staff scale. ^{a,b,c} Significant difference between the same alphabet letter at p < 0.05, Bonferroni correction.

1.1 professionals per provider and 1.9 persons with dementia per professional were registered.

Baseline characteristics

The 67 care professionals had a mean age of 45.9 (SD = 10.3) years and included 29 men (43.3%). There were 34 direct care workers (other than nurses) (50.7%) and 30 care managers (44.8%). The mean number of months of tenure in care for older adults was 165.1 (SD=89.1). Twenty-five professionals had graduated from university (37.3%) and 24 from vocational school or college (35.8%). The mean sense of competence in dementia care measured by SCIDS was 40.9 (SD=6.8). Care professionals from in-home care management agencies were older, included fewer men, had a longer tenure in care for older adults, and included no professionals other than care managers. Most care professionals from group homes and nursing homes were direct care workers. There were no significant differences in measured competence in dementia care between types of providers or professionals (Table 2).

Intervention

At baseline, the most frequently specified unmet needs were another person bothering the person (n =67, 53.6%), sleepiness or tiredness (n = 58, 46.4%), and pain (n = 55, 44.0%). The action plan was mostly linked to pleasant activity/recreation (n = 38, 30.4%). The most frequent caregivers involved in the interdisciplinary meeting were direct care workers who were not nurses (n = 123, 98.4%) and care managers (n = 80, 64.0%) (Supplementary Table 1). On average, the baseline assessment was performed 19.8 (SD = 10.6) days after the training course. The mean number of days from baseline to T2 was 97.1 (SD = 21.9).

The average total NPI-NH score was 19.4 (SD = 16.4) at T1, and 16.0 (SD = 15.2) at T2. The total score decreased significantly from baseline to T1 and T2 in each type of provider, except for nursing homes between baseline and T1. The observed effect size at T2 was medium (0.44–0.61) at each type of provider (Table 3).

Time investment of care professionals

At baseline, the 119 persons with dementia had complete information on the time investment of care professionals for the program implementation. There were 111 persons with complete T1 information, and 108 with complete T2 information. The average total minutes for the five-month interventions with completed follow-ups was 417.9 (SD = 219.8) with a decrease of 8.6 points (SD = 14.0) in the NPI total

	In-home care					Residential care				
	Care management		Multiple in-home		Group home		Nursing home			
Time of assessment	Ν	Mean (SD)	Ν	Mean (SD)	Ν	Mean (SD)	Ν	Mean (SD)		
Baseline	38	18.7 (16.1)	28	25.2 (17.7)	36	28.8 (18.2)	23	32.9 (19.2)		
T1	33	13.3 (11.2)	27	17.8 (14.4)	34	19.3 (13.4)	22	30.9 (23.3)		
Coefficient (95%CI)	-5.41*		-6.13*		-9.94*		-1.98			
	(-9.09, -1.73)		(-10.82, -1.44)		(-13.78, -6.09)		(-10.67, 6.70)			
T2	30	11.5 (12.1)	26	16.7 (17.7)	32	17.7 (14.8)	20	19.3 (16.3)		
Coefficient (95%CI)	-7.40*		-7.64*		-10.45^{*}		-12.08^{*}			
	(-11.20, -3.59)		(-12.40, -2.89)		(-	-14.38, -6.52)	(-21.04, -3.12)			
Effect size, Cohen's d _{rm}	0.55		0.44		0.61		0.61			
ICC in the null model										
Person with dementia	0.304		0.350		0.308		0.197			
Care professional	0.304		0.350		0.308		0.197			
Random effect in the full model										
Residual	59.320		77.709		65.855		218.613			
Person with dementia		60.914	108.487		87.739		81.133			
Care professional		60.914	108.487		87.739		81.133			

 Table 3

 Average total Neuropsychiatric Inventory scores by type of provider at baseline, T1, and T2

Coefficients and 95% confidence intervals, and interclass correlation coefficient were estimated using multilevel linear regression analysis including person with dementia and care professional as random effects. Level of neuropsychiatric symptoms was assessed using the Japanese version of the Neuropsychiatric Inventory-Nursing Home version (NPI-NH). *p < 0.0125, Bonferroni significance threshold.

Table 4 Time investment of professionals for program implementation by type of provider

	Total N	Mean (SD)		In-home care			Residential care			
			Care management		Multiple in-home		Group home		Nursing home	
			N	Mean (SD)	N	Mean (SD)	N	Mean (SD)	N	Mean (SD)
Minutes										
Baseline	119	193.7 (103.3)	38	184.5 (71.1)	25	172.4 (118.0)	33	219.6 (126.6)	23	194.8 (92.4)
T1	111	126.5 (77.6)	33	133.2 (85.5)	27	101.9 (60.8)	30	151.8 (87.6)	21	111.3 (58.2)
T2	108	101.2 (58.7)	30	102.9 (53.4)	26	64.7 (21.4)	32	127.7 (77.0)	20	123.5 (41.9)
Total	103	417.9 (219.8)	30	420.4 (190.4)	24	311.0 (153.6)	29	508.2 (281.6)	20	411.4 (172.4)
Total cost, yen	103	28836.6	30	26421.0	24	25122.3	29	31824.9	20	32584.0
		(14912.1)		(15231.3)		(13269.6)		(15006.9)		(15479.1)
Change in NPI	103	8.6 (14.0)	30	8.2 (11.1)	24	6.6 (15.5)	29	9.3 (13.3)	20	10.3 (17.6)

Estimated total cost accounted for time investment, national average wage, and number of participants in the discussion meeting. Level of neuropsychiatric symptoms was assessed using the Japanese version of the Neuropsychiatric Inventory-Nursing Home version (NPI-NH).

score (Table 4). Two-fifths of the total time investment was spent on the discussion meeting, one-third on preparation for the meeting, and one-fifth on using the web-based tool to input evaluation of the NPI, unmet needs, and action plan determined in the meeting (Supplementary Table 2).

The average number of professionals who participated in the baseline meeting was 3.1 (SD = 1.9). Forty-eight persons (40.3%) experienced other professionals declining to participate in the meeting at baseline. The most frequent reason for declining to participate was time constraints. The percentage of professionals declining to participate did not differ by type of provider ($\chi^2(6) = 12.08$, p = 0.060) (Supplementary Table 3). The multilevel linear regression analysis showed a significantly lower time investment at T1 and T2 than at baseline. The time investment among in-home care management agencies was significantly lower than that in group homes. There were significant associations of lower time investment with both a shorter tenure in care for older adults and greater competence in dementia care. The ICC for professionals was 0.280 (Table 5). Sensitivity analyses using each category instead of total time investment displayed a significantly higher time investment for preparation and use of the web-based tool among group homes than in-home care management agencies. Time investment for preparation was also significantly higher among nursing homes than care management agencies.

 Table 5

 Factors relating to time investment for the program implementation

	1.8.1.
Coefficient (95% confidence interval)	Total minutes
Type of provider, reference = in-home	
care management	
Multiple in-home service provider	7.97 (-32.39, 48.33)
Group home	59.83* (16.80, 102.85)
Nursing home	36.20 (-9.68, 82.08)
Time of evaluation,	
reference = baseline	
T1	-66.92* (-80.43, -53.41)
T2	-91.72* (-105.34, -77.61)
Characteristics of care professionals	
Primary qualification, care	14.61 (-21.32, 50.54)
manager/social worker	
Tenure in care for older adults, month	0.23* (0.06, 0.39)
Competence in dementia care	-2.29* (-4.09., -0.49)
(17–68)	
Characteristics of persons with	
dementia	
Age at baseline, year	0.70 (-1.17, 2.57)
Sex, male	21.59 (-7.52, 50.70)
Type of dementia, Alzheimer's	15.79 (-10.75, 42.33)
disease	
Prescribed medication	
N05A: antipsychotics	-6.33 (-35.74, 23.09)
N06D: anti-dementia drugs	1.17 (-21.86, 24.21)
Level of neuropsychiatric symptoms	0.18 (-0.35, 0.72)
(0–144)	,
Random effect	
Residual	2512.072
Person with dementia	1599.603
Care professional	1599.603

Multilevel linear regression analysis including person with dementia and care professionals as random effects. Interclass correlation coefficient inn the full model was 0.280 for care professional. Competence in dementia care was measured by the Japanese version of the Sense of Competence in Dementia Care Staff scale. Level of neuropsychiatric symptoms was assessed using the Japanese version of the Neuropsychiatric Inventory-Nursing Home version (NPI-NH). *p < 0.05.

There were significant associations between a lower time investment for preparation or use of the webbased tool and shorter tenure in care for older adults. Time investment for preparation was significantly lower among care professionals whose primary qualifications were nurses or other direct care workers, and professionals with greater competence in dementia care (Supplementary Table 4).

DISCUSSION

The level of neuropsychiatric symptoms declined significantly in group home and nursing home residents as well as in in-home service users. The effect size of the reduction in neuropsychiatric symptoms was medium (0.44–0.61) for each type of provider.

The time investment for program implementation among in-home care management agencies was significantly lower than for group homes and smaller at follow-ups than at baseline. To the best of our knowledge, this is the first study to have uncovered the time investment of a psychosocial dementia care program for neuropsychiatric symptoms both in inhome and residential care settings. Our findings have implications for exploring dissemination strategies for psychosocial dementia care interventions. Further, professionals with shorter tenure in care for older adults and those with greater competence in dementia care took significantly less time for implementation than professionals with longer tenure and lower competence.

Contrary to our expectations, the time investment was not higher among in-home care management agencies than in other types of providers. Furthermore, the time investment among in-home care management agencies was significantly lower than that in group homes. Although group home residents did not show a significant difference in their level of neuropsychiatric symptoms from in-home service users, persons with dementia living in group homes tended to have more severe neuropsychiatric symptoms than in-home service users [45]. Thus, care professionals may have needed more time for discussions to identify unmet needs and devise an action plan. It is globally recommended to start advance care planning to share the person's values, goals, and preferences regarding the rest of life at the time of dementia diagnosis [19]; to date, however, there is no legislation or regulations regarding advance care planning in Japan [46]. The small-scale homelike environment in group homes could have added another disadvantage to program implementation. Although this environment has positive aspects such as increased decision authority among nursing staff [47], it has barriers in terms of integrated care for nursing staff working alone for a large part of the day [48]. Whereas the lack of regular multidisciplinary meetings is perceived as a facilitator to psychosocial management of neuropsychiatric symptoms [49], it could also escalate the time required for care professionals to become acquainted with the DEMBASE® program. It would be beneficial to explore the incorporation of advance care planning in the program implementation to make information more available to specify unmet needs and form an action plan.

In previous research in Tokyo, where the majority of participating professionals were in-home care managers, the professionals presumed that their challenges in holding interdisciplinary discussion meetings were specific to in-home care settings [26]. In this study, however, more than half of group home and nursing home residents received declines from professionals other than participating employees to attend the discussion meeting. Time constraint was the most frequently cited reason for the decline across all types of providers, which is consistent with international findings regarding the implementation of psychosocial interventions for persons with dementia [50, 51]. Therefore, the largest time investment at the first implementation is a unique barrier to program implementation regardless of setting. Since the time investment gradually decreased from baseline to follow-ups, care professionals who managed the implementation and other professionals who participated in the discussion meeting acquired knowledge regarding the individual needs of the person with dementia, resulting in a faster implementation process. The facilitators to program implementation included visualized feedback regarding mitigated neuropsychiatric symptoms via a web-based tool that motivated care professionals only at follow-up assessments [26]. This delayed-return feature of the program may increase the risk of cessation before follow-ups when the time investment for baseline assessment is felt not to be worth the expense. In this study, the most frequent reason for cessation was time constraints for the discussion meeting. An additional benefit scheme under the public long-term care insurance program would be helpful to reward the time investment and encourage program implementation until follow-ups.

The time investment was significantly lower among care professionals with greater professional competence in dementia care. Their enhanced competence may enable care professionals to more effectively explain the program and encourage other professionals to participate in discussion meetings. Our results imply that effective program implementation may require a certain level of competence among care professionals who manage the implementation for each provider. A shorter professional tenure in the care for older adults was also significantly related to lower time investment in all categories except for the discussion meeting. More experienced professionals may have required more time to adapt to new dementia care practice. Alternatively, less experienced professionals were aware of fewer signs of unmet needs and acted sooner to determine the action plan. However, even after adjustment for competence in dementia care and tenure in care for older adults.

28% of the variance (ICC = 0.280) in time investment was accounted for by individual care professionals, implying that time investment may depend on which professional manages the implementation. Further refinement of the program components should be explored to help care professionals proceed with implementation with minimum time investment. It should be cautioned that discussion in the care team is an essential aspect and that working too fast may increase the risk of overlooking the essence of a successful action plan. Instruction on how to structure the discussion meeting would be helpful to shorten the time required for professionals.

The total time investment for program implementation was 418 minutes on average, per person, besides an 8.6-point decrease in NPI total score during the five-month period, with three interdisciplinary discussion meetings. Using the national mean wage for long-term care providers in 2018 [52], the total cost for the five-month implementation was estimated at 3353 yen for a one-point decrease in NPI score. Since the incremental cost of a one-point increase in NPI score was estimated as 30 USD per month [7] or 247-409 USD per year [53], which would be driven by caregiving time [8, 54], the program may have yielded a fair return on the time investment. It should be noted that reduction in neuropsychiatric symptoms could impact the costs of psychotropic drug use, physical/psychological examination, consultation, and treatment for the management of neuropsychiatric symptoms [55, 56]. Future research should evaluate the change in these costs by a reduction in neuropsychiatric symptoms, as well as the time investment for program implementation.

Strengths and limitations

The main strength of our study lies in including both in-home and residential care settings in the implementation of a common psychosocial dementia care program. A significant reduction in neuropsychiatric symptoms was observed in both in-home and residential care settings, which incurred a substantial time investment. However, our study has some limitations. The number of participants per type of provider did not fulfil the expected sample size due to the effect of natural disasters during the recruitment process. Care professionals were also unable to access training courses following the government's advice to avoid large gatherings during the COVID-19 pandemic, caused by the severe acute respiratory syndrome coronavirus 2 (SARS- CoV-2). As the pandemic and consequent distancing might increase neuropsychiatric symptoms of people with dementia [57, 58], the psychosocial dementia care program delivered remotely through technology should be further explored. Currently, an e-learning training course of the program has been developed in the Netherlands and Japan to enable care professionals encouraged to avoid travel due to COVID-19 to participate in the program remotely.

The time investment was based on professionals' reports, which may have resulted in response bias toward overestimation. Because participant recruitment was conducted by nationwide provider organizations, participating professionals probably had higher competence (mean 40.9) than care professionals in Tokyo (mean 37.0) [31]. This sampling bias may have resulted in lower time investment for the program implementation and underestimation of the association between time investment and professional competence. The present study did not assess the quality of life of persons with dementia, which should be evaluated to measure the cost-effectiveness of the program in the future.

Conclusions

The program implementation incurred a substantial time investment in both in-home and residential care settings. The total time investment may have a fair return when the level of reduced neuropsychiatric symptoms is considered. However, there is a risk of cessation before follow-ups when the time investment for the baseline assessment is felt not to be worth the expense. An additional benefit scheme to reward the time investment would be helpful to encourage the program implementation until follow-ups.

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CONFLICT OF INTEREST

The authors declare that there is no conflict of interests.

SUPPLEMENTARY MATERIAL

The supplementary material is available in the electronic version of this article: https://dx.doi.org/ 10.3233/ADR-200235.

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