Table 2 Continued

Variable	Category	Male caregiver n %	n = 399	Female caregiver n %	n = 1193	<i>P</i> -value
Money management	Independent	107	26.8	310	26.0	NS
	Mostly independent	75	18.8	194	16.3	
	Fairly dependent	58	14.5	171	14.3	
	Completely dependent	158	39.6	518	43.4	
Medication	Independent	149	37.3	389	32.6	NS
	Mostly independent	47	11.8	148	12.4	
	Fairly dependent	65	16.3	208	17.4	
	Completely dependent	137	34.3	447	37.5	
Telephone	Independent	138	34.6	409	34.3	NS
	Mostly independent	70	17.5	180	15.1	
	Fairly dependent	50	12.5	156	13.1	
	Completely dependent	141	35.3	448	37.6	
Shopping	Independent	30	7.5	84	7.0	NS
	Mostly independent	56	14.0	162	13.6	
	Fairly dependent	94	23.6	241	20.2	
	Completely dependent	219	54.9	706	59.2	
Transportation use	Independent	28	7.0	84	7.0	NS
•	Mostly independent	48	12.0	163	13.7	
	Fairly dependent	99	24.8	237	19.9	
	Completely dependent	224	56.1	709	59.4	

The χ²-test was conducted between male and female caregiver groups. ADL, activity of daily living; IADL, instrumental activity of daily living; NS, not significant.

The ADL of the dependents in the male and female caregiver groups were matched, except for auditory capacity. Dependents in the female caregiver group were more dependent in IADL. A good explanation for this is that the male dependents were generally unskilled<sup>22</sup> and more female caregivers cared for a male dependent than male caregivers.

As for the characteristics of the caregivers, consistent with previous studies,<sup>3-5</sup> male caregivers were more likely to be older than their female counterparts. Also, as mentioned earlier, there was a kinship difference between the male and female caregiver groups. According to previous studies,<sup>17,23</sup> the differences in age and kinship should be taken into account in analyzing our results.

In items of care services, consistent with previous studies, the male caregiver group was more likely to use home help.<sup>3,19</sup> It is generally believed that men are less experienced with housework<sup>3,7</sup> and our results probably reflect this situation. In addition, female caregivers were more likely to use a day care/service in our study. Sugiura *et al.* and Colline *et al.* previously explained that women tended to prefer respite care.<sup>3,6,22</sup> Our results seem to support their suggestions. Contrary to what might be expected, more frequent use of home-visit nursing care by male caregivers was observed in this study. Male caregivers

were less likely than female caregivers to help an older person with different types of illnesses who may need more nursing care. However, to our knowledge, few studies have so far dealt with this issue. Additional studies are needed to obtain a more accurate appraisal of the gender differences in the use of inhome care.

Inconsistent with earlier studies, 3.24 which found that female caregivers showed a higher depression rate than male caregivers, we detected no significant difference in GDS-15 in this study. In addition, the nurses' subjective assessment in this study showed that male caregivers used less formal or informal care, and that male caregivers were in worse health. It is possible that this had a negative effect on the male caregivers, resulting in a higher depressive mood, because there is a strong relation between caregiver burden and depression. 20.21.25 However, the GDS-15 was developed to assess the depressive mood of the elderdy and not that of a younger population. We should think of this result only as a suggestion.

#### Gender differences in caregiver burden

This study focused on differences in caregiver burden according to gender. Our results, regardless of adjusting, did not reveal any difference between male and female caregivers with respect to caregiver burden.

Table 3 Gender differences in main caregiver characteristics

Variable	Categories	Male caregiver N %, mean ± SD (range)	n = 399	Female caregiver N %, mean ± SD (range)	n = 1193	<i>P</i> -value
Age (years)		68.3 ± 12.7 (31-91)		62.5 ± 12.1 (31-93)		<0.001
Kinship	Spouse	234	58.6	417	35.0	< 0.001
	Child	143	35.8	419	35.1	
	Daughter/son-in-law	5	1.3	311	26.1	
	Sibling	8	2.0	27	2.3	
	Other	9	2.3	18	1.5	
	Unknown	0	0.0	1	0.1	
Types of care service use	Day care/service	166	41.6	586	49.1	0.009
· ·	Home-visit rehabilitation	31	7.8	90	7.5	NS
	Home-visit bathing	46	11.5	165	13.8	NS
	Short stay	37	9.3	142	11.9	NS
	Home help	211	52.9	451	37.8	< 0.001
	Family physician home-visit	256	64.2	707	59.3	NS
	Home-visit nursing care	232	58.1	609	51.0	0.014
	Housing adjustments	93	23.3	297	24.9	NS
	Care implements rental	237	59.4	770	64.5	NS
Depressive mood GDS-15		5.5 ± 4.0 (0–15)		5.1 ± 3.9 (0–15)		NS
Nurse's assessment						
Use of care service by	Sufficient	157	39.3	531	44.5	NS
caregiver	Average	189	47.4	545	45.7	
	Insufficient	53	13.3	115	9.6	
	Unknown	0	0.0	0	0.0	
Caregiving by family	Sufficient	167	41.9	656	55.0	<0.001
	Average	183	45.9	446	37.4	
	Insufficient	47	11.8	85	7.1	
	Unknown	2	0.5	0	0.0	
Caregiver's health	Excellent	141	35.3	522	43.8	< 0.001
-	Normal	193	48.4	529	44.3	
	Below standard	63	15.8	135	11.3	
	Unknown	2	0.5	0	0.0	

The  $\chi^2$ -test for categorical variables or the unpaired t-test for continuous variables was conducted between male and female caregiver groups. GDS, geriatric depression scale; NS, not significant.

This finding supports the result of Aoki et al.'s study<sup>5</sup> and differs from various other studies which suggested that female caregivers showed more caregiver burden than their male counterparts.3,6-11 Our results suggest that further studies are needed to prove the gender difference in caregiver burden, at least in Japan. In Japan, the public long-term care insurance system which was implemented in 2000 provides a care-management system by professional care managers.<sup>27,28</sup> Care management facilitates the selection of appropriate care services for elderly people among available care services provided in the community based on a care need assessment.28 A care manager needs to monitor a dependent's physical and mental condition to assess the latest care need as occasion demands.27,28 Therefore, the system provides for a high level of care and helps caregivers cope with

stress, giving them relief from caregiver burden.<sup>29</sup> It is possible that the care management system lessened the female caregiver burden and narrowed the gender gap in caregiver burden. Moreover, female caregivers reportedly tend to seek informal support from family and neighbors.<sup>3,5</sup> We did not investigate the use of informal care, except family care, and therefore we were unable to determine the extent to which caregivers were given informal support by care providers except family.

#### Study limitation

The current study has several limitations. Although the NLS-FE is a large-scale observational study, it does not include the complete spectrum of elderly patients in the Nagoya area. In addition, the selection of subjects was somewhat biased because the par-

Table 4 Gender differences in caregiver burden and depressive mood

(0-81) n = 399	000	aregiver n %, mean ± SD			Odds ratio		Odds ratio adjusted		Odds ratio adjusted for age and other	
26.0 ± 18.5(0–81)	880 II I	(range)	n = 1193	۵	n = 1193 P unadjusted	95% CI	for age	95% CI	variablest	95% CI
26.0 ± 18.5(0–81) 's assessment Severe 113 28.3			Í							
113 283		27.3 ± 17.6(0-84)		SN	1.004	0.997-1.011 1.005	1.005	0.998-1.012	1 004	0.005_1.012
113 283							) } !			310.
0::	113 28.3	274	23.0	SR						
	53.9	715	59.9							
Light 69 17.3		196	16.4							

cerebrovascular disease person in charge of medication, auditory capacity, food preparation, home maintenance, laundry, caregiver's age, kinship, use of daycare/service, use of home help, use of home-visit nursing care, caregiving ₽ confidence interval; NS not significan peripheral vascular disease, logistic caregiver quality of family relationship, cognitive heart failure, ulcer disease, female ਠੰ and conducted ş was continuous and the ģ t-test unpaired 휼 ŏ categorical variables and caregiver's health status. ₫ The x2-test family,

ticipants were groups of users of home nursing stations using home visiting nurses or care planning services.

Another limitation is that we requested that each station perform its own evaluation due to a shortage of staff and the large quantity of settings. This may have biased the assessors' evaluation and limited the validity of the results, including the nurses' subjective assessment.

Finally, this study is an analysis of data from a large-scale study. Therefore, our database does not always capture the full extent of the dependents' and caregivers' characteristics needed to obtain a precise analysis. A lack of data concerning caregiving period, caregiving hours per day or details of required care weakened the impact of our findings.<sup>23,30</sup>

#### CONCLUSION

We conducted a subanalysis of a large scale observational study in Japan. Our results indicated that there were no differences in caregiver burden between male and female caregivers. Further studies are needed to confirm whether or not gender differences do in fact exist.

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#### Conflict of interest declaration

The authors declare that they have no competing interests.

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#### ORIGINAL ARTICLE

# **Predicting Recovery of Upper-Body Dressing Ability After Stroke**

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ABSTRACT. Suzuki M, Omori M, Hatakeyama M, Yamada S, Matsushita K, Iijima S. Predicting recovery of upper-body dressing ability after stroke. Arch Phys Med Rehabil 2006;87: 1496-502.

Objective: To identify predictors of the recovery of independent dressing ability after stroke.

Design: Prospective cohort study.

Setting: Rehabilitation unit at a university hospital.

Participants: Sixty-three consecutive stroke patients were enrolled in the study. Twelve patients were not able to complete the study because they were discharged or transferred to another hospital before study completion.

Intervention: Fifty-one patients underwent and completed 15 days of dressing training based on the time-delay method, which included the 10 component actions of upper-body dressing and 4 cues given by therapists.

Main Outcome Measures: The dressing item of the FIM instrument, Brunnstrom motor recovery stages, presence or absence of deep and tactile sensation, Rey-Osterrieth complex figure test, Kohs block design test, body image test, Weintraub cancellation task, and presence or absence of the visual extinction phenomenon and the motor impersistence phenomenon.

Results: The FIM upper-body dressing item score and the cancellation task score at the start of training were significantly better in patients who achieved independence in dressing within 15 training days than in patients who did not (P < .05). The motor impersistence phenomenon was found less frequently among patients who achieved independence in upperbody dressing than among patients who did not (P < .05). However, logistic regression analysis showed that only the FIM score for upper-body dressing on the first day of training was a significant independent predictor of dressing ability at the end of training (odds ratio, 4.33; 95% confidence interval, 1.51-12.37). The receiver operating characteristic curve indicated that a cutoff score of 3 would provide the best balance between sensitivity and specificity for the FIM upper-body dressing item. The positive predictive value of this cutoff score was .90, and the negative predictive value was .70.

Conclusions: Our findings indicate that the FIM upper-body dressing score on the first day of dressing training is an indepen-

dent predictor of recovery of upper-body dressing ability after stroke.

**Key Words:** Activities of daily living; Prognosis; Rehabilitation; Stroke.

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NDEPENDENCE IN DRESSING enables a person to maintain a sense of dignity, self-respect, and achievement. Therapists working with stroke patients spend a large proportion of the day teaching patients how to put on and take off items of clothing. Despite such instruction, however, many patients are still unable to dress themselves independently for several weeks after hospital admission. The Dressing is more difficult than undressing, and upper-body dressing requires more advanced recognition than lower-body dressing. The

vanced recognition than lower-body dressing. 1.2 Many studies 1.2.7-13 have shown a relation between difficulty in dressing and cognitive and physical impairments. Despite more than 50 years of research, it is still difficult to predict the extent or duration of loss of dressing ability.7-15 There is no clear understanding of the effect of early neurologic impairments and early dressing disorder on a stroke patient's ultimate recovery from dressing disorder. 1.2.7-15 In a single-blind randomized controlled trial, Kwakkel et al14 investigated the effects of different intensities of arm and leg rehabilitation on the functional recovery of activities of daily living (ADLs) including dressing. They found no differences in ADL scores between the arm-training and control groups, and they suggested that stroke patients compensate for the loss of function in the paretic arm by using the nonparetic arm during ADLs. They noted that impairment was not always associated with dressing disorder in their patients. Jongbloed<sup>15</sup> performed a critical review of 33 studies and concluded that the admission ADL score is a strong predictor of discharge ADL status, but its relation to improvement in cognitive and physical impairment is unclear.

Which is a stronger predictor of recovery of dressing ability, early neurologic impairments or early dressing disorder? Because it is still difficult to identify predictors of dressing ability, success in training is largely due to the extent of a therapist's experience rather than any scientific data. If we could identify predictors of recovery of dressing ability and predict the extent or duration of dressing ability loss, training in upper-body dressing skills would become more evidence based.

We conducted a prospective cohort study to identify predictors of the recovery of independent dressing ability after stroke.

#### **METHODS**

#### Outcome Measures

The study protocol is shown in figure 1. Each patient's upper-body dressing ability was assessed according to the FIM instrument dressing item<sup>16</sup> on the first day of dressing training.

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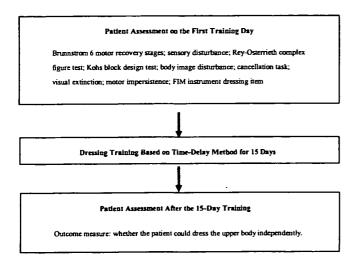


Fig 1. Study protocol.

The ability to dress the upper body is precisely defined by the FIM, and scores range from 1 to 7, with 1 indicating complete dependence during an activity and 7 indicating complete independence.

Many studies 1,2,7-12 have shown a relation between difficulty in dressing and cognitive and physical function impairments including motor palsy, sensory disturbance, constructional disorder, body image disturbance, visual inattention, unilateral spatial neglect, and motor impersistence. Therefore, the presence or absence of these impairments was also determined. The severity of motor palsy was assessed according to the 6 motor recovery stages of Brunnstrom<sup>17</sup> representing muscle conditions ranging from the initial flaccidity of palsy to normal coordination. Sensory disturbance was evaluated according to the presence or absence of deep and tactile sensation. Constructional disorder was assessed by the Rey-Osterrieth complex figure test<sup>18</sup> and Kohs block design test<sup>19</sup>; these are 36-point and 131-point scales, respectively. Body image disturbance was assessed by an unpublished test of each patient's ability to discriminate the head, nose, right shoulder, left shoulder, abdomen, and back of the neck (6-point scale). Visual inattention was evaluated by a cancellation task involving a sheet of paper containing 360 randomly arranged shapes, 60 of which were target stimuli. 20 Unilateral spatial neglect was measured by the presence or absence of the visual extinction phenomenon.<sup>21</sup> Motor impersistence was assessed by each patient's ability to sustain tongue protrusion and eye closure simultaneously for 20 seconds.<sup>22</sup> The reliability and validity of the 4 tests (FIM, Brunnstrom stages, Rey-Osterrieth complex figure test, Kohs block design test) have been established. 21,23-32 We assessed 2 tests for their test-retest reliability in 15 stroke patients with an interval of 5 days between measurements. The intraclass correlation coefficients were .78 for the target cancellation task (P<.01) and 1.00 for the body image test (P < .01).

#### **Participants**

Sample size calculation was based on a desired 95% statistical power to detect a 1-point difference in the FIM dressing item score, with a 2-sided  $\alpha$  of 5%. The average value and standard deviation (SD) of FIM dressing item scores in 20 stroke patients were assessed to determine the standard effect size. The average FIM dressing item score was  $2.45\pm1.50$  points, and the standard effect size was .66. A sample size of 53

was derived by insertion of 1-power (.05),  $\alpha$  (.05), and standard effect size (.66) values in the Hulley matrix.<sup>33</sup> We therefore planned to recruit about 50 stroke patients.

Between May 1, 2001, and May 1, 2004, 63 consecutive stroke patients from the Department of Rehabilitation Medicine, St. Marianna University School of Medicine Hospital, were enrolled in the study. Stroke was diagnosed according to the World Health Organization definition.<sup>34</sup> Eligibility criteria included hemiplegia, dependence on spoken cues or physical assistance to accomplish upper-body dressing, ability to sit up with a backrest for more than 30 minutes, lucid consciousness, a period of less than 3 months since the stroke event, absence of severe cardiopulmonary or respiratory insufficiency, and a desire to participate in the study. Baseline characteristics of patients who satisfied the inclusion criteria are presented in table 1. The mean age of participants was 69.4±10.6 years. There were 25 women and 38 men, 45 patients with cerebral infarction and 18 with cerebral hemorrhage, and 27 patients with right hemiplegia and 36 with left hemiplegia. The average time since the stroke event was 23.0±17.2 days.

Twelve patients (6 with right hemiplegia, 6 with left hemiplegia) withdrew from the study because they were discharged or transferred to another hospital before study completion.

The study was approved by the St. Marianna University School of Medicine Institutional Committee on Human Research. Informed consent was obtained from each patient before his/her participation in the study.

#### Intervention

ADLs such as dressing are considered behavioral chains of component actions.<sup>35</sup> Such chains have been learned and performed since childhood. A patient with hemiplegia cannot dress by means of the behavioral chains of component actions used by a healthy person and thus has to learn new behavioral chains of component actions to achieve independence in dressing. It is necessary to control the cue stimulations and rewards

Table 1: Baseline Characteristics of Patients Who Satisfied the Inclusion Criteria

Characteristics	Values
Age (y)	69.4±10.9
Sex (n)	
Male	38
Female	25
Diagnosis (n)	
Infarction	45
Hemorrhage	18
Time poststroke at assessment (d)	23.0±7.2
Paralysis side (n)	
Right	27
Left	36
Sensory disturbance (n)	
Tactile sense	21
Deep sense	17
Visual extinction phenomenon (n)	28
Motor impersistence (n)	23
FIM dressing item	2.0 (2.0-3.0)
Brunnstrom motor recovery stage	3.0 (2.0-4.0)
Kohs block design test score	0.0 (0.0-17.0)
Rey-Osterrieth complex figure test score	4.0 (0.0-21.4)
Target cancellation task	21.0 (0.0-49.0)
Disturbance of body image	6.0 (4.0–6.0)

NOTE. Values are mean ± SD, n, or median (interquartile range [IQR]).

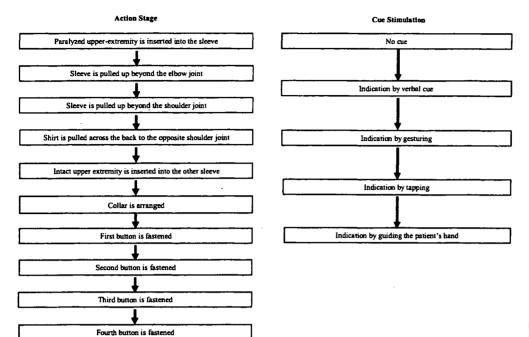


Fig 2. Upper-body dressing training by the time-delay method.

presented in training to support the organization of new behavioral chains. 35,36 For the purpose of the study, upper-body dressing was viewed in 10 separate stages: (1) the paralyzed upper extremity is inserted into the sleeve, (2) the sleeve is pulled up beyond the elbow joint, (3) the sleeve is pulled up beyond the shoulder joint, (4) the shirt is pulled across the back to the opposite shoulder joint, (5) the intact upper extremity is inserted into the other sleeve, (6) the collar is arranged, (7) the first button is fastened, (8) the second button is fastened, (9) the third button is fastened, and (10) the fourth button is fastened (fig 2). These 10 component actions describe the entire process of upper-body dressing. The list was developed after observation of the behavioral chains of upper-body dressing used by 33 stroke patients with hemiplegia. The 22 stroke patients (66.7%) who achieved the greatest independence in upper-body dressing used these 10 component actions as a behavioral chain. Thus, these actions were selected for use in our current study.

The study patients underwent 15 days of training based on the time-delay method,<sup>37</sup> which is a recognized and effective training method.<sup>38</sup> In the time-delay method, cues are given after a set interval of time has elapsed, in this case 10 seconds. The starting position for dressing training was the patient grasping the shirt collar. Dressing training began with the verbal instruction, "Please put on the shirt." If the patient responded with inadequate component actions or if the patient did nothing for 10 seconds, the therapist offered cues at 4 levels in the following order: (1) verbal cue, (2) gesturing, (3) tapping, and (4) physical assistance. Verbal cues were instructions such as, "Can you pass your right hand into the sleeve?" or "Can you pull the sleeve up to your elbow?" Gesturing consisted of the therapist mimicking the component action of upper-body dressing. Tapping consisted of the therapist tapping the patient's clothes and body. Physical assistance consisted of the therapist taking the patient's hand and guiding it in the appropriate direction. When the patient performed each component action, the therapist praised him/ her. After 15 days of training, each patient was assessed for his/her ability to dress the upper half of the body independently.

#### Statistical Analysis

Patients were classified into 2 groups: those who could perform the upper-body dressing tasks independently after the 15 days of training and those who required assistance. Cognitive and physical function and upper-body dressing ability on the first day of dressing training were compared between the 2 groups. Differences in categoric variables were analyzed by the chi-square test or Fisher exact test. The Mann-Whitney U test was used to analyze ordinal variables. Logistic regression analysis was used to identify the best independent predictors of independent upper-body dressing ability after 15 days of training. All statistical procedures were performed with SPSS software<sup>a</sup> with a significance level set at P equal to .05. A receiver operating characteristic (ROC) curve was used to assess the clinical utility of the independent predictors.<sup>39</sup> Constructing the ROC curve involved setting several cutoff points for significant variables and calculating sensitivity, specificity, positive predictive value, and negative predictive value at each point.

#### **RESULTS**

Statistics related to subjects' performances of each task are presented in table 2. Upper-body dressing ability and visual attention at the start of training were significantly better in patients who achieved independence in dressing within 15 training days than in patients who did not. The FIM upper-body dressing item score for independent patients was higher than that for dependent patients (median score, 3 points; interquartile range [IQR], 2-3 points vs median score, 2 points; IQR, 1-2 points; P < .001). The target cancellation task score for independent patients was higher than that for dependent patients (median score, 40 points; IQR, 10.5-57.5 points vs median score, 2 points; IQR, 0-21 points; P=.004). In addition, motor impersistence was found less frequently among patients who achieved independence in upper-body dressing than among patients who did not. There were 4 (14.3%) independent patients and 12 (54.2%) dependent patients (P=.014) with motor impersistence. Independence in dressing was not

Table 2: Predictors of Upper-Body Dressing Ability After Stroke

Characteristics	Independent (n=28)	Dependent (n=23)	P*	Odds Ratio (95% C
Age* (y)	69.8±10.0	70.3±10.0	.837	NS
Sex (% male)	53.6	65.2	.580	NS
Diagnosis (% cerebral infarction)	71.4	69.6	.758	NS
Paralysis side (% right hemiplegia)	46.4	34.8	.259	NS
Tactile sense disturbance (% positive cases)	25.0	34.8	.543	NS
Deep sense disturbance (% positive cases)	17.9	26.1	.732	NS
Visual extinction phenomenon (% positive cases)	28.6	60.9	.079	NS
Motor impersistence (% positive cases)	14.3	54.2	.014	NS
FIM dressing item	3.0 (2.0-3.0)	2.0 (1.0-2.0)	<.001	4.33 (1.51-12.37
Brunnstrom motor recovery stage	3.0 (3.0-4.3)	3.0 (2.0-3.5)	.163	NS
Kohs block design test score	7.0 (0.0–18.8)	0.0 (0.0-1.0)	.063	NS
Rey-Osterrieth complex figure test score	13.5 (1.3–24.5)	2.5 (0.0-6.5)	.088	NS
Target cancellation task	40.0 (10.5-57.5)	2.0 (0.0-21.0)	.004	NS
Disturbance of body image	6.0 (4.0-6.0)	6.0 (4.0-6.0)	.668	NS

NOTE. Values are mean ± SD, median (IQR), or as otherwise indicated. Odds ratios show logistic regression analysis. Abbreviations: CI, confidence interval; NS, not significant.

significantly associated with the severity of motor palsy or body image disturbance. Logistic regression analysis of the 14 variables showed only the FIM upper-body dressing score to be a significant predictor of the recovery of dressing ability (odds ratio, 4.33; 95% confidence interval, 1.51–12.37).

FIM upper-body dressing scores were plotted as an ROC curve (fig 3). Sensitivity, specificity, and predictive value at several cutoff points are presented in table 3. The curve indicated that a cutoff score of 3 being "moderate" would provide the best balance between sensitivity and specificity for the FIM upper-body dressing item (sensitivity, .68; 1 – specificity, .09). The positive predictive value of this cutoff score was .90, and the negative predictive value was .70. Characteristics of patients who withdrew from the study were similar to those of patients who completed the study (table 4).

#### DISCUSSION

Our results indicate that the FIM upper-body dressing score on the first day of dressing training is an independent predictor

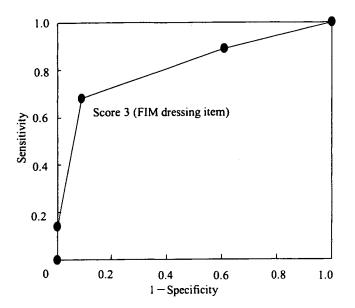


Fig 3. ROC curve for the FIM upper-body dressing item.

of recovery of upper-body dressing ability after stroke. Williams<sup>8</sup> correlated the constructional abilities of 136 hemiplegic patients with their abilities to relearn upper-extremity dressing skills. Patients with normal constructional ability were more likely to be independent in upper-extremity dressing or had a greater capacity to achieve this skill than patients with a constructional disorder. In an evaluation of 60 stroke patients by the Nottingham Stroke Dressing Assessment and other physical and cognitive assessments, Walker and Lincoln<sup>2</sup> found that difficulty in lower-body dressing was associated with physical impairment and that difficulty in upper-body dressing was associated with visual inattention and sensory disturbance. Chen et al<sup>11</sup> investigated the relation between patterns of visuospatial inattention and performance of ADLs by means of the Klein-Bell ADL Scale and the Random Chinese World Cancellation Test in 64 patients with a right brain lesion. They found that hemi-inattention was highly related to poor ADL performance and that independence in dressing appeared to be more adversely affected by hemi-inattention than was independence in bathing and hygiene, eating, or use of the telephone. Hier et al<sup>12</sup> evaluated 41 patients with unilateral right hemisphere stroke for hemiparesis, hemianopia, constructional apraxia, spatial neglect, dressing disorder, and motor impersistence. Dressing disorder was associated with severe constructional apraxia, spatial neglect, motor impersistence, and hemianopia.

In marked contrast to the findings of earlier studies, 1.2.7-12 the final regression model in our study showed no significant relation between the recovery of upper-body dressing ability and any underlying cognitive or physical impairment assessed in this study. As noted above, patients with hemiplegia are unable to use the same behavioral chains that are used by healthy people to accomplish dressing tasks. However, if a new behavioral chain is learned, patients can achieve a degree of

Table 3: Sensitivity, Specificity, and Predictive Values at 3
Cutoff Points

Cutoff Point	Sensitivity	Specificity	Positive Predictive Value	Negative Predictive Value
2	0.893	0.391	0.641	0.750
3	0.679	0.913	0.905	0.700
4	0.143	1.000	1.000	0.489

<sup>\*</sup> $\chi^2$  test or Fisher exact test (categoric variables), Mann-Whitney U test (ordinal variables).

Table 4: Characteristics of Subjects Who Completed the Study and Those Who Withdrew

Characteristics	Patients Who Completed the Study	Patients Who Withdrew From the Study	P
Age (y)	70.0±9.9	66.6±15.4	.716
Sex (% male)	60.8	58.3	.383
Diagnosis (% cerebral infarction)	70.6	75.0	.563
Paralysis side (% right hemiplegia)	41.2	50.0	.290
Tactile sense disturbance (% positive cases)	29.4	50.0	.218
Deep sense disturbance (% positive cases)	21.6	50.0	.281
Visual extinction phenomenon (% positive cases)	43.1	50.0	.021
Motor impersistence (% positive cases)	31.4	0.7	.002
Period from crisis to measurement (d)	23.0±16.5	22.8±21.8	.620
FIM dressing item	2.0 (2.0-3.0)	2.0 (2.0-3.5)	.446
Brunnstrom motor recovery stage	3.0 (2.5-4.0)	2.0 (2.0-3.5)	.104
Kohs block design test score	0.0 (0.0–17.0)	0.0 (0.0–17.5)	.932
Rey-Osterrieth complex figure test score	4.0 (0.5–22.5)	0.0 (0.0–6.0)	.092
Target cancellation task	21.0 (0.0-50.5)	15.0 (0.5–23.0)	.453
Disturbance of body image	6.0 (4.0-6.0)	6.0 (2.5-6.0)	.617

NOTE. Values are mean ± SD, median (IQR), or as otherwise indicated.

independence in dressing. The reason why early dressing disorder is a stronger predictor of the recovery of dressing ability than cognitive or physical impairment is related to the degree of change to the original behavioral chain.

According to univariate analysis, visual attention at the start of training was significantly better in patients who achieved independence in dressing within 15 training days than in patients who did not. Also, motor impersistence was found less frequently among patients who achieved independence in upper-body dressing than among patients who did not. These findings corroborate those reported by Walker and Lincoln<sup>2</sup> and Hier et al. 12 There is still no clear understanding of the effect of underlying neurologic impairments on a stroke patient's ability to relearn to dress. Prior studies<sup>1,2,7-12</sup> have failed to answer this question for several reasons. In some cases, multivariate analysis was not performed, and in others, variables were not tested for independent prediction. In addition, no prospective cohort study to identify the predictors of recovery of independent dressing ability has been conducted. In the present study, the target cancellation task score and motor impersistence on the first day of dressing training were independent predictors of recovery of upper-body dressing ability after stroke. However, the recovery of independent dressing ability was more strongly related to the FIM upper-body dressing score than to the target cancellation task score and motor impersistence.

Our analysis indicated that the FIM upper-body dressing score can serve as a valuable predictor of the ability to dress the upper body independently after stroke. Ninety percent of patients with a FIM upper-body dressing score of 3 or more on the first training day recovered the ability to dress the upper body independently within 15 training days. However, 70% of patients with a score of 2 or less could not perform this task independently after 15 days of training. Such patients require other solutions, such as different types of training or changes in materials or types of clothes. In addition, therapists should be consulted about the appropriate method for assisting these patients. Therapists can predict the recovery of independent dressing ability after stroke scientifically by an initial assessment with the FIM dressing item. Our findings will contribute to an increasingly evidence-based approach to upper-body dressing training for stroke patients.

The FIM score was investigated in relation to the burden of care; a 1-point change in the total FIM score was equivalent to

an average of 2 to 5 minutes of help from another person per day. 25-27 Rogers et al<sup>38</sup> examined the effectiveness of a behavioral rehabilitation intervention based on the time-delay method for improving the performance of morning care routines by nursing home residents with dementia. In their study,<sup>38</sup> physical assistance were provided for significantly smaller proportions of a morning care session during the behavioral rehabilitation intervention. However, the intervention took considerably more time than was needed for the usual care. Our results indicated that the time spent in nonassisted dressing increased for patients who could perform upper-body dressing independently after the 15 days of training. However, a therapist may spend more time with a patient who requires only partial assistance after the 15 days of training than with patients who require full assistance. Therefore, therapists should devise a method in which assistance and promotion of independence are balanced.

#### **Study Limitations**

Because this was a prospective cohort study, we did not randomize patients into groups before the training. Patients were classified into 2 groups after training: those who could perform the upper-body dressing tasks independently after the 15 days of training and those who required assistance. We also did not evaluate cognitive or physical function of patients after training. Therefore, the effect of dressing training based on the time-delay method is not clear in this study. If patients were allocated before or re-evaluated after the training, the effect of any natural recovery could be excluded. Further research in a randomized controlled trial is needed to verify the effect of dressing training based on the time-delay method.

The FIM upper-body dressing item used in this study is part of a standardized ADL test. However, this item does not distinguish the component actions of upper-body dressing, and it does not account for cues given by the therapist during evaluation of upper-body dressing ability. The difficulty of upper-body dressing varies according to the dressing components and is also affected by cues given during evaluation. Thus, the FIM upper-body dressing item cannot be used to evaluate details of upper-body dressing or the level of assistance required. Therefore, further research is needed to develop an upper-body dressing assessment scale that accounts for the individual components of dressing activities

and controls for cue stimulation during any evaluation of dressing skills.

The number of participants in our study was determined on the basis of Hulley's matrix for sample-size estimation.<sup>33</sup> However, a larger number of participants will be needed in further studies to remove the influence of natural differences between people in recovery from cognitive and physical impairments. With the addition of a detailed examination classifying participants by types of lesion and by attributes and the inclusion of a large number of patients, the results of a study like ours would be more generalizable.

#### **CONCLUSIONS**

We conducted a prospective cohort study to investigate the influence of early neurologic impairments or early dressing disorder on the recovery of independent dressing ability after stroke. Our findings indicate that early dressing disorder (as measured by the FIM upper-body dressing score) on the first day of dressing training is an independent predictor of the upper-body dressing ability after stroke. We expect our findings will contribute to a more evidence-based method of training in upper-body dressing skills.

The most popular behavioral chain of component actions in dressing training based on the time-delay method was used in this study. However, the behavioral chain of component actions will vary according to the seriousness of impairments in cognitive and physical function. There was no significant relation between the recovery of upper-body dressing ability and any underlying cognitive or physical impairment assessed in this study, but this finding may be related to the fact that only 1 behavioral chain was targeted. Therefore, it is necessary to investigate the relation between cognitive and physical impairments and several different behavioral chains of component actions.

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#### Supplier

a. SPSS Inc, 233 S Wacker Dr, 11th Fl, Chicago, IL 60606.

〈原 著〉

高齢重度認知症患者および高齢進行癌患者の在宅終末期ケアに関する研究 ~在宅終末期ケアを推進する診療所群における前向き研究から~

平川 仁尚" 益田雄一郎" 葛谷 雅文" 井口 昭久" 旭 多貴子" 植村 和正"

要 約 目的:高齢化に伴い、高齢認知症患者と高齢癌患者が増加しているが、両者の終末期ケアの相違に関する実証研究はほとんどない、本研究は、在宅における認知症高齢者と高齢進行癌患者の終末期ケアの実態を明らかにすることを目的として、日本ホスピス在宅ケア研究会の協力を得て実施した「高齢者の在宅終末期ケアに関する前向き研究」のデータの二次解析を行った。方法:この研究は、日本ホスピス在宅ケア研究会に所属する医師 16 名が担当した患者のうち、2002 年の 10 月から 2004 年の 9 月までの間に、最終的に自宅で看取られた 65 歳以上の高齢患者 240 名を対象としたものである。調査内容は、患者の特徴(性別、年齢、障害老人の日常生活自立度(JABC)、痴呆性老人の日常生活自立度、死因など)、死亡前 48 時間以内に観察された症状と実施された終末期ケア、などであった。これらのデータの収集は、患者の死後、カルテや家族の話などを参考に、患者の担当医師が質問紙に回答する形式で行われた。解析にあたって、重度認知症患者群 (痴呆性高齢者の日常生活自立度 III 以上)と進行癌患者群を比較・検討した。結果:進行癌患者群と比較して、重度認知症患者群において、疼痛、悪心・嘔吐、せん妄が少なく、発熱、咳嗽が多かった。また、痰の吸引や抗生剤の投与の実施頻度が高く、輸液量は多かった。麻薬系鎮痛剤の投与について、重度認知症患者群では実施されていなかったが、進行癌患者群では約半数に実施されていた。結論:今回の結果から、重度認知症患者と進行癌患者の終末期の特徴の違いに配慮する必要があることが示唆された。

Key words:麻薬,症状,終末期ケア,疼痛,痴呆

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#### 緒 言

わが国は世界でも類を見ない速さで高齢社会を迎えている。そして、人は老いて死ぬことが避けられない以上、高齢社会の到来は、高齢者の死の増加を意味する。とくに、認知症や癌は高齢者によくみられる疾患であり、その多くが進行性であることから、認知症患者や癌患者の死亡が増加することが予想される。

終末期ケアの目標は、苦痛な症状を管理し、患者の QOLを最大限に高めることである。そのためには、終 末期によくみられる症状や実施可能な終末期ケアに関す る充分な情報に基づいて議論を行い、患者と家族の希望 に沿った終末期のケアの計画を立てる必要がある<sup>1)</sup>. 高 齢認知症患者と高齢癌患者の終末期の特徴には違いがあ

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ることが指摘されているが<sup>2)~0</sup>, 先行研究は少なく, 両者の違いに関する実証データの蓄積が必要である.

一方、近年、高齢者の在宅終末期ケアが、注目されている<sup>5</sup>、わが国では、他の先進国と同様<sup>6</sup>、病院で死亡する患者が多いが<sup>5</sup>、在宅で最期を迎えたいと希望する高齢者が多いことが示唆されている<sup>517</sup>、最期を迎える場所により、終末期ケアが違う可能性があり<sup>8</sup>、在宅終末期ケアに関するデータの蓄積も必要である。

本研究の目的は、「高齢者の在宅終末期ケアにおける前向き研究」のデータを用いて、高齢重度認知症患者と高齢進行癌患者の終末期の特徴を比較検討し、その違いを明らかにすることである。

#### 対象と方法

「高齢者の在宅終末期ケアに関する前向き研究」は、 日本ホスピス在宅ケア研究会の協力を得て実施した。こ の前向き研究は、日本ホスピス在宅ケア研究会に所属す る医師 16 名が担当した患者のうち、2002 年の 10 月か ら 2004 年の 9 月までの期間に、最終的に自宅で看取ら

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	T dole			
項目	内訳	認知症(n=36) n (%)	癌 (n=116) n (%)	р
性別	女	24 (66.67)	51 (43.59)	0.017
年齢(平均 ±SD,歳)		87.61 ± 0.837	74.52 ± 1.238	0.000
障害老人の日常生活自立度	J	0	2 (1.71)	0.004
	A	1 (2.78)	11 (9.40)	
	В	4 (11.11)	25 (21.37)	J
	С	31 (86.11)	59 (50.43)	1
	不明	0	19 (16.24)	
死因 (原発巣)	胃	_	22 (18.80)	-
	肺	<del>-</del>	30 (25.64)	_
	肝	_	19 (16.24)	_
	大腸		9 (7.69)	_
	膵	-	3 (2.56)	_
	腎		3 (2.56)	_
	血液	_	0	_
	脳	_	1 (0.85)	_
	その他		17 (14.53)	
	不明	_	12 (10.26)	
死因(非癌疾患)	呼吸器	14 (38.89)	_	_
	循環器	5 (13.89)	_	_
	脳血管	2 (5.56)	_	
	腎	2 (5.56)	_	_
	肝	0	_	
	消化器	0	<del>-</del>	_
	その他	12 (33.33)	_	_
	不明	1 (2.78)	<del>-</del>	_

Table I 患者の特徴

れた 65 歳以上の高齢患者 240 名を対象とした. 調査内容は, 患者の特徴(性別, 年齢, 寝たきり度(障害老人の日常生活自立度, JABC), 痴呆性老人の日常生活自立度, 疾病および死因など), 死亡前 48 時間以内に観察された症状と実施された終末期ケア, などである. 今回の調査で使用した症状とケアの項目は次の通りである.

#### 症状

呼吸困難, 疼痛, 自制内疼痛, 昏睡, せん妄, 不安, めまい, 吐き気・嘔吐, 食欲不振, 下痢, 便秘, 発熱, 尿便失禁, 吐血, 喀血, 下血, その他の出血, 咳, 痰, など

#### 終末期ケアおよび検査

心臓マッサージ, 気管内挿管, 人工呼吸器, 酸素吸入, エアウェイ留置, 痰の吸引, 高カロリー輸液, 末梢点滴, 抗生剤, 昇圧剤, 輸血(成分輸血を含む), 麻薬, 尿導 カテーテル留置, 心理的ケア, 宗教的癒し, など

これらのデータの収集は、患者の死後、カルテや家族 の話などを参考に、患者の担当医師が質問紙に回答する 形式で行われた。

解析にあたって、statview5.0 日本語版を使用した.

連続量には t 検定を, 離散量には  $\chi2$  乗検定を用いた. p< 0.05 を統計学的に有意差があるものとし、高齢重度認知症患者群 (以下、認知症患者群) と高齢進行癌患者群 (以下、癌患者群)を比較・検討した。尚、本調査において、重度認知症を痴呆性高齢者の日常生活自立度 III 以上と定義し、認知症患者群に分類した。また、質問紙の死因欄に癌が明記されている場合にのみ癌患者群に分類した。すなわち、癌患者であっても、癌以外の疾患で死亡した可能性がある患者は解析から除外した。

#### 成 續

解析対象者は、認知症患者群 36 人、癌患者群 116 人であった、対象者の特徴 (表1) について、平均死亡年齢は、認知症患者群で有意に高かった (認知症患者群 87.61±0.84 歳 vs 癌患者群 74.52±1.24 歳). また、性別は、認知症患者群で女性が多かった。障害老人の日常生活自立度について、認知症患者群で自立度が低い傾向がみられた。死因について、認知症患者群では、呼吸器疾患が約 40% と最も多く、循環器疾患が約 15% と続いた、癌患者群の癌の部位では、肺が約 25% と最も多く、胃

症状	認知症(n=36) n (%)	癌(n=116) n(%)	р
呼吸困難	14(38.89)	56 (47.86)	0.324
我慢できない疼痛	0	25(21.37)	0.002
自制内疼痛	2(5.56)	57 (48.72)	0.000
昏睡	12 (33.33)	50(42.74)	0.297
せん妄	3(8.33)	26 (22.22)	0.060
不安	2(5.56)	14(11.97)	0.266
眩暈	1 (2.78)	2(1.71)	0.691
悪心嘔吐	3(8.33)	33(28.21)	0.013
食欲不振	18 (50.00)	70 (59.83)	0.272
下痢	2(5.56)	7 (5.98)	0.915
便秘	2(5.56)	9 (7.69)	0.656
発熱	18 (50.00)	29(24.79)	0.005
失禁	3 (8.33)	17(14.53)	0.327
吐血	1 (2.78)	4(3.42)	0.844
喀血	0	1 (0.85)	0.576
下血 .	4(11.11)	7(5.98)	0.304
出血(吐血・下血・喀血以外)	1(2.78)	9(7.69)	0.292
咳嗽	15(41.67)	15(12.82)	0.000

Table 2 死亡前 48 時間以内に観察された症状

Table 3 死亡前 48 時間以内に実施された終末期ケア

15(41.67)

5(13.89)

ケア	認知症(n=36) n (%)	癌 (n=116) n (%)	р
心臓マッサージ	2(5.56)	1 (0.85)	0.077
挿管	0	0	–
人工呼吸器	0	0	-
酸素吸入	8(22.22)	45 (38.46)	0.068
エアウェイ	0	3(2.56)	0.330
吸痰	16 (44.44)	29(24.79)	0.026
髙カロリー輸液	2(5.56)	14(11.97)	0.266
抗生剤	12 (33.33)	11 (9.40)	0.001
昇圧剤	0	0	-
輸血	0	0	-
末梢点滴	11 (30.56)	38 (32.48)	0.805
輸液量 (平均 ±SD, ml)			
24 ~ 48 時間前	$880 \pm 110.353$	502.22 ± 55.920	0.003
0~24時間前	$800 \pm 108.711$	467.14 ± 54.079	0.004
麻薬	0	60(51.28)	0.000
尿道カテーテル	6(16.67)	23(19.66)	0.673
心理的ケア	0	3(2.56)	0.330
宗教的癒し	0	1 (0.85)	0.576
その他	3 (8.33)	9(7.69)	0.911

#### が約20%と続いた.

死亡前 48 時間以内に観察された症状を表に示す (表 2). 我慢できない疼痛および自制内疼痛は、癌患者群と 比較して、認知症患者群で有意に少なかった、悪心・嘔 吐とせん妄は、認知症患者群で少なかったが、せん妄に

喀痰・痰詰り

その他

関しては統計学的に有意差が認められなかった.一方, 発熱と咳嗽は、認知症患者群で有意に多かった.

35 (29.91)

29 (24.79)

0.200

0.169

死亡前48時間以内に実施されたケアを表に示す(表3). 心臓マッサージ, 挿管, 人工呼吸器は, 両群ともほとんど実施されていなかった. 痰の吸引や抗生剤の投与

は、癌患者群と比較して、認知症患者群で広く実施されていた。末梢点滴の実施率には両群で有意差がみられなかったが、輸液量は認知症患者群で有意に多かった。また、麻薬の投与は、認知症患者群では実施されていなかったが、癌患者群では約半数に実施されていた。

#### 考 察

平成15年度の厚生労働省「人口動態統計」(http://www.mhlw.go.jp/toukei/saikin/hw/jinkou/geppo/nengai03/index.html)によると、加齢とともに死因全体に占める肺炎の割合が増加し、90歳以上の高齢者の死因では心疾患が一位(19.9%)、二位に肺炎(17.1%)、悪性新生物は三位(12.2%)となる。本研究における認知症患者の死因では呼吸器疾患が全体の約40%を占め、一般の高齢者と比べて死因に占める割合が高い、加齢以外の要素として、重度認知症患者は嚥下性肺炎で死亡することが多いこと<sup>214</sup>が影響していると考えられる。

前述のように、高齢重度認知症患者と高齢進行癌患者において、終末期にみられる症状やケアに違いが見られることが指摘されている<sup>2/4/5/</sup>が、今回の結果は、これを裏付けるものであった。すなわち、疼痛、せん妄、悪心・嘔吐、発熱、咳で、その頻度に両群間に違いがみられた。

疼痛について、認知症患者群ではほとんど観察されなかったのに対して、癌患者群では多くの患者で観察された、癌は疼痛を伴うことが多い疾患である一方、認知症患者は疼痛に対して寛容であるという意見があり<sup>9)</sup>、こうした背景が結果に反映されたと考えられる。また、今回の結果では認知症患者群は癌患者群と比較して高齢であったが、一般的に高齢者は若年者と比べて疼痛の訴えが少ないといわれており<sup>277101</sup>、この年齢の違いが両群にみられた理由と考えられる。さらに、認知症患者群は、痛みを感じてもコミュニケーション能力の低下などにより疼痛を訴えることができなかったとも考えられる<sup>9111</sup>、

認知症はせん妄の危険因子であるが<sup>12</sup>、癌患者群と比べて、認知症患者群のせん妄の出現率は低かった。我々の知り得る範囲では、癌患者と認知症患者のせん妄の出現率を比較した調査はないため、新たな知見である可能性がある。麻薬の使用や疼痛もせん妄の危険因子であるため<sup>12</sup>、これらの頻度が癌患者群で高かったことも今回の結果に影響を与えている可能性がある。

今回の結果では、悪心・嘔吐は癌患者群で広く観察された。悪心・嘔吐は、終末期癌患者、とくに消化器癌の 患者において高頻度に見られる症状である<sup>13</sup>、癌患者群 において約3分の1が胃癌と大腸癌である一方で認知症 患者群において消化器疾患で死亡した患者がいなかった こと、癌患者群で麻薬の使用や疼痛が多かったこと、などが理由として考えられる<sup>13</sup>.

咳と発熱は、癌患者群と比較して、認知症患者群で広く観察された。咳の原因として、肺炎や肺癌など肺疾患が多いい。今回の結果では、認知症患者群の約40%、癌患者群の約25%、が肺疾患で死亡していたため、この差が咳の出現率の差の説明になり得る。しかし、転移性肺癌は、今回の調査結果には含まれていないことを考慮する必要がある。また、高齢者では、発熱は見られないことがしばしばあるが150、今回の結果では、認知症患者群で、抗生物質を多く投与され、高齢であったにもかかわらず、発熱が多くみられた。ステロイドや非ステロイド性解熱鎮痛剤など解熱効果のある薬剤の使用頻度は調査していないが、高齢の重度認知症患者においても発熱の管理が必要であることが示唆される。

死亡前48時間以内に、両群において、心臓マッサージや人工呼吸器の使用など苦痛を伴う延命治療の実施はほとんどみられなかった。場所により終末期ケアが異なると言われているが。在宅は、他の場所と比較して3、終末期患者への延命治療が差し控えられる傾向があるかもしれない。

認知症患者群では、痰の吸引が広く行われていた、痰の出現頻度に有意差がみられていなかったことから、癌患者群では、吸引を必要としない程度の少ない痰であったと推察できる.

癌患者群と比較して、認知症患者群で抗生剤は広く使用されていた。発熱が癌患者群と比較して高頻度で観察されたことがその主要因と考えられるが、終末期の重度認知症高齢者への抗生剤の使用には、副作用の出現など問題点を指摘する意見があり3060、議論を深める必要がある

わが国において、終末期癌患者に対する適切な輸液量に関しては、医師の間で一致した見解は得られていないが「"、終末期癌患者に対する過剰な輸液を控えることで、気道の分泌物を減らし、苦痛を軽減することができるといわれている「①18」、今回の結果では、癌患者において約500m1/日であり、認知症患者と比べて輸液量が少なかった。本研究に参加した医師は、終末期ケアに関心が高いと考えられ、苦痛を軽減するため、輸液を控えた可能性が示唆される。一方、重度認知症患者の終末期においても、輸液量を控えることで、苦痛を緩和できる可能性がある「9」、今後、重度認知症高齢者の終末期の輸液のあり方も議論されるべきであろう。

麻薬の使用は、認知症患者では報告されず、癌患者群で多く報告された、疼痛が癌患者群で高頻度にみられ、

認知症患者では疼痛がほとんど見られなかったことから、妥当な結果であるといえる. しかし、認知症患者群で呼吸困難を訴える患者が少なからず存在した. 麻薬は呼吸困難を緩和する作用があり<sup>2</sup>、認知症患者群において麻薬の使用が適切であったかどうかは、さらなる詳細な調査が必要である.

本研究の限界について以下に述べる.

- 1. 疼痛以外は、症状の程度や頻度を調査していない.
- 2. 今回の対象者は、在宅で終末期ケアを受け、最期に在宅で看取られた患者であった。そのため、症状が重度であった患者は、看取りの直前に病院に転送された可能性がある。
- 3. 今回の調査では対象施設は、在宅終末期ケアに関心を持つ集団であったことが結果に影響を与えた可能性は否定できない。
- 4. 両群で性別の割合に違いがみられたが、性別は、 症状に影響を与える可能性がある<sup>20)</sup>.

以上より、今回の結果をわが国の在宅終末期ケアの実態として一般化することは難しい、今後、これらの課題をふまえ、追加調査が必要であると考える。

#### まとめ

今回、高齢者の在宅終末期ケアに関する前向き調査のデータの二次解析を行い、重度認知症高齢者と高齢癌患者の終末期にみられる症状と行われたケアの相違を明らかにした。死亡前48時間以内に、認知症患者において、疼痛、悪心・嘔吐、せん妄が少なく、発熱、咳嗽が多かった。また、行われたケアについて、痰の吸引や抗生剤の投与について認知症患者群で頻度が高く、輸液量については認知症患者群で有意に多かった。また、麻薬系鎮痛剤の投与について認知症患者群では全員に行われていなかったが、癌患者群では約半数で使用されていた。

今回の結果から、重度認知症患者と進行癌患者の終末期の特徴の違いに配慮する必要があることが示唆されたが、本研究の限界を踏まえ、追加調査が必要である.

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## Home end-of-life care for advanced dementia vs advanced cancer elderly patients: Dying elderly at home project

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#### **Abstract**

AIM: The aim of this study was to assess the frequency of symptoms and end-of-life care received in advanced dementia and advanced cancer elderly patients dying at home during the last two days of their lives and to evaluate the differences observed between the two groups.

METHODS: We used data from the Dying Elderly at Home (DEATH) project, which was a prospective study of home elderly patients dying with end-stage illness. Consecutive deceased subjects aged 65 or older who were seen at 16 study clinics belonging to the Japanese Society of Hospice and Home-care with diagnoses of all illnesses including advanced dementia and advanced cancer and died at home from October 2002 to September 2004 were included in the study. We evaluated 36 deceased subjects with advanced dementia and 116 with advanced cancer. We collected the following information:sociodemographics, ADLs, cognitive impairment, observed symptoms and end-of-life care provided during the last 48 hours of life.

RESULTS: Deceased subjects with advanced dementia were less likely to show symptoms of pain, acute confusion, or nausea/vomiting and more likely to display fever or cough than advanced cancer patients. Also, those with advanced dementia were more likely to receive intravenous drip injection or narcotic analgesia and more likely to be given sputum suction, or antibiotics.

**CONCLUSION**: We observed that the dying process and end-of-life care for advanced dementia elderly patients was different from that for advanced cancer elderly patients.

**Key words**: Opioid, Symptom, End-of-life care, Pain, Dementia (Jpn J Geriat 2006; 43: 355–360)

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### 簡便な操作で痛みの強さを記憶する痛み計の臨床試行

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本研究の目的は,患者が簡便な操作で随時,主観的な痛みの強さを記録できる「痛み計」を開発し,臨床における有用性 を確認することである。「痛み計」(23cm x 6cm x 2cm,160g) は,0-10 Numeric Rating Scale (NRS)を採用した 11 個の 押しボタンを有し,患者が疼痛に相当する数値のボタンを押すとその数値と日時を記憶する.「痛み計」をパソコンに接続 すると,痛みの強さをグラフとして印刷できる.研究方法は事例検討である.大学病院に入院中でがん性疼痛のある患者 1 例に 14 日間,痛み計の使用を依頼した.印刷したグラフは患者と医療スタッフに渡した.その結果,以下の点が示唆され た、1.0-10NRS を用いたことにより、患者は痛みを円滑に表現できた、2. 操作を簡便にしたことにより、患者は随時、入 力できた、3.1 日の痛みの推移をグラフで出力したことは、疼痛アセスメントに有効であった。

Keywords: 疼痛測定, 数値的評価スケール, がん性疼痛, 測定機器, 端末機

#### 緒雲

米国の Joint Commission on Accreditation of Healthcare Organizations (JCAHO) は, 2001 年に疼痛を第5のパイタルサインと 位置づけて全ての患者に疼痛のモニタリングが必要であるこ とを明言した[1]. 痛みを積極的にアセスメントし, 緩和ケア を促進するためにその意義は高い. 疼痛の強さは熱や脈と異 なり、適切な客観的測定法は未だなく [2], 患者が主観的に認 知する程度が指標となる.しかし,がん患者は痛みの訴えを 躊躇する傾向 [3] があり、医療者が患者の主観的な疼痛をモニ タリングすることは難しい. そのため患者による症状の記録 が重要であり、近年では紙の記録から電子記録に変わり、携 帯性やパソコンとの互換性に優れた Personal Digital Assistance (PDA) の開発について研究されている [4]. 痛みの記憶は思い 出した時の感覚に影響される [5] ため, PDA と紙の記録を比 較した研究では, 実際は多くの患者が振り返って記入してい た紙の記録ではなく、入力時間が記録される PDA の有用性が 報告されている[6]. しかし、PDAの操作は必ずしも患者にとっ て容易ではなく [7], 入力忘れが多いことも指摘されているた め[8]、患者が簡便な操作で随時、主観的な痛みの強さを記録 できる「痛み計」の着想に至った.本研究の目的は、「痛み計」 を実際に事例に試みて有用性を確認することである.

#### 方法

#### 1. 痛み計の着想(図1)

痛み計は,縦6cm,横23cm,厚さ2cm,重量160gの電子機 器であり、主観的な痛みの強さを記録するための道具である. 痛みの強さの評価は、0(全く痛みが無い)から10(最悪の痛み)

の Numeric Rating Scale (NRS: 数値的評価スケール) を採用し た.操作は,患者が簡便に入力できることを重視し,電源を入 れ、痛みの強さに該当する数値のボタンを押すという2つの 手順とした、押した数値は表示窓に表示され、15秒後に電源 は自動的に切れる.入力された痛みのデータは,日時とともに 痛み計内に記録され、専用のソフトをインストールしたパソ コンに取り込むことによって出力することができる. 本研究 では、痛みの強さの推移が視覚的に分かるよう、1日単位の折 れ線グラフで印刷した。

#### 2 車例への試み

大学病院に入院中の患者に対し,試作した痛み計を2週間 使用するように依頼して協力を得た. 患者の負荷を軽減する ため入力時間や回数は任意とし、さらに入力回数は痛み計の 簡便性を評価する指標とした.グラフは1日に1回出力し,患 者と医療スタッフの双方に渡した.なお,本研究は名古屋大学 医学部倫理委員会保健学部会の承認および対象施設の承諾を 得て実施し,対象者に研究の主旨と倫理的配慮について説明 して依頼し,文書で承諾を得た.



図 1. 痛み計.

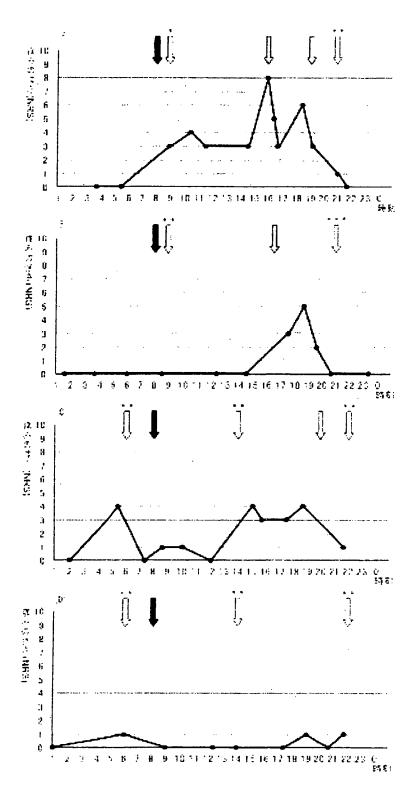


図 2. 痛み計使用による痛みの強さの推移 .(A) 痛み計使用 2 日目, (B) 痛み計使用 6 日目, (C) 痛み計使用 10 日目 , (D) 痛み計使用 13 日目 .

白色矢印:塩酸オキシコドン徐放剤

\*5mg, \*\*10mg, \*\*\*15mg

灰色矢印:塩酸モルヒネ内服薬 5mg 黒色矢印: 非ステロイド性消炎鎮痛薬 10mg

NRS: Numeric Rating Scale

#### 結果

#### 1. 事例紹介

A氏は58歳の女性で,肺がんの術後に気管支およびリンパ 節転移に対して、化学療法と放射線療法を施行中である. 左胸 部と背部の疼痛に対して,塩酸オキシコドン徐放剤(以下,オ キシコドン)と非ステロイド性消炎鎮痛薬を内服している.

#### 2. 痛み計使用の実際 (図2)

痛み計導入日(オキシコドン開始9日目)までのA氏は「痛 かったけど、もうすぐ鎮痛剤の時間だから我慢してた.」と 語り、痛みを我慢することがあった、痛み計導入当初は、レス キューを複数回使用する状況であり (図 2-A), まずオキシコド ンを増量した(図2-B).しかし,21時の定期内服前に疼痛が強 くなる傾向があり、A氏本人の「薬が切れるせいか、夕方にな ると痛みが出てくる」という自覚を考慮して, 痛み計9日目 にオキシコドンをさらに増量し、12時間ごとの内服を8時間 ごとの内服に変更したところ (図 2-C), 痛み計 13 日目には終 日 0-1 のレベルで過ごすことができるようになった (図 2-D). 痛み計への入力操作はスムーズで,1日平均10.6(範囲:4-14) 回であった.

A氏は、痛み計3日目に「昨日はつい我慢してしまい、本当 に辛かった、だからか、レスキューを飲んでもあんまり効か なくて,また痛くなった.」とグラフを見て語り,痛み計7日 目には「夜中は、オキシコンチンが効いてるけど、トイレに行 きたくて目が覚めるついでに記録してあるの.だから,0や1 ばっかりでしょ.このことも先生にも伝えなきゃね.」と説明 した、また、痛み計を2週間使用した後に「疼痛が強くなった ときに,その場で記録しておくことは難しいけど大切.値はす ぐ忘れてしまうので、この器械があったのは本当に良かった. ぜひ改良して皆さんに使ってあげてほしい.」と感想を述べ, 改良が必要な点として,小型化,起動時間の短縮などをあげ た.

#### 考察

A氏は0-10NRSを用いた疼痛の評価に混乱する様子はなく, 痛みを数値化することができた、すべての患者が痛みの強さ を数値化できるとは限らないが、痛み計に採用したことは妥 当であったと判断できる.

A氏の入力回数が、1日平均 10.6回であったことは、これま でに開発された疼痛を記録する電子機器では,1日3回と規定 した入力回数のコンプライアンスが 91-94%[6, 8], 本研究と同 様に入力回数を任意とした報告 [5] では平均 6.08 回であった ことから、痛み計の操作の簡便性を支持する結果であった。

データをグラフとして出力したことにより,A氏と医療ス タッフの双方にとって痛みの強さの推移が分かりやすくなっ た.A氏による痛み計3日目の振り返りは,グラフとして表示 した痛みの強さを見たことで、改めて痛みと鎮痛薬のタイミ ングの理解につながったことを示した. 医療スタッフは, グラ フに表示された痛みの傾向と A氏の自覚をもとに、オキシコ ドンは1日2回の投与で有効とされる薬剤であるが、増量お よび1日3回の投与に変更した.実際の疼痛緩和は,オキシコ ンチンの増量,投与間隔の短縮,化学療法および放射線療法の 影響といった様々な要因が考えられるが、医療スタッフに痛 みの推移を視覚的に提示したことは, 疼痛アセスメントの一 助となったと言える.

A氏は痛み計によって,痛みの強さを随時, 記録することの 重要性や痛みの記録がサポートされたと感じ, そのデータを 主体的に医療スタッフとの会話に活用していた.また,改善が 必要な点も指摘しており、今後はさらに多くの患者で痛み計 の効果を検証すること,および患者の意見とコストを考慮し た改善を重ねることが課題である。

#### 結論

疼痛のある患者自身が、簡便に痛みの強さを入力できる痛 み計を開発し、1事例に試みた結果、入院患者が痛み計を使用 することは,以下の点でペインコントロールに役立ち,有用な 道具としての可能性を示唆した.1. 痛み計に用いた0-10NRS によって、患者は痛みを円滑に表現できた、2. 痛み計の操作は 簡便でシンプルな機能にしたことにより,患者は随時.入力で きた.3.1日の痛みの推移をグラフ化して視覚的に分かり易く 出力することは、疼痛アセスメントに有効であった.

ただし、今回は1事例であるため、より多くの患者の協力を 得て検討を重ねる必要がある.

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#### 铭爈

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